

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/97ISR Number: 3025593-0Report Type:Periodic Company Report #A0052454  
 Age:38 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG/ TWICE PER DAY/ ORAL		Breast Discharge	Health	Wellbutrin	PS		ORAL
		Breast Tenderness	Professional				
		Galactorrhoea	Company Representative	Cetirizine Hydrochloride Beta Carotene Fluticasone Propionate Aspirin Trazodone Clonazepam Amlodipine Medroxyprogesterone Midrin Metoclopramide Enalapril Maleate Estratest Darvocet-N Oxpentifylline Calcium Carbonate Parafon Forte Hydroxychloroquine S04	C C C C C C C C C C C C C C C C C C C		

Date:11/03/97ISR Number: 3025599-1Report Type:Periodic Company Report #A0052699  
 Age:48 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG/ TWICE PER DAY/ ORAL		Amnesia	Health	Wellbutrin	PS		ORAL
			Professional				
			Other	Dexfenfluramine			

15 MG / TWICE

Capsule

SS

ORAL

PER DAY /

ORAL

Date:11/03/97ISR Number: 3025602-9Report Type:Periodic Company Report #A0052754

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oedema Peripheral	Health	Wellbutrin	PS		ORAL

ORAL

Professional  
Company  
Representative

Date:11/03/97ISR Number: 3025606-6Report Type:Periodic Company Report #A0052828

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Influenza Like Illness	Health	Wellbutrin	PS		ORAL

150 MG / ORAL

Professional  
Company  
Representative

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Freedom Of Information (FOI) Report

Date:11/03/97ISR Number: 3025611-XReport Type:Periodic Company Report #A0052857  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea	Health	Wellbutrin	PS		ORAL
150 MG/ORAL			Professional				

Date:11/03/97ISR Number: 3025615-7Report Type:Periodic Company Report #A0052904  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Skin Discolouration	Health	Wellbutrin	PS		ORAL
ORAL			Professional				

Date:11/03/97ISR Number: 3025619-4Report Type:Periodic Company Report #A0052934  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema Multiforme	Health	Wellbutrin	PS		ORAL
150MG / TWICE			Professional				
PER DAY/ ORAL							

Date:11/03/97ISR Number: 3025622-4Report Type:Periodic Company Report #A0052935  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema Multiforme	Health	Wellbutrin	PS		ORAL
150MG / TWICE			Professional				
PER DAY /							
ORAL							

Date:11/03/97ISR Number: 3025625-XReport Type:Periodic Company Report #A0052936  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema Multiforme	Health	Wellbutrin	PS		ORAL
150MG / PER			Professional				
DAY / ORAL							

Date:11/03/97ISR Number: 3025631-5Report Type:Periodic Company Report #A0052941  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State	Health	Wellbutrin	PS		ORAL
ORAL		Tremor	Professional Company Representative				

Date:11/03/97ISR Number: 3025635-2Report Type:Periodic Company Report #A0052982  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Injury	Consumer	Wellbutrin	PS		ORAL
150MG / TWICE		Therapeutic Response					
PER DAY /		Decreased					
ORAL				Calcium Salt Multivitamin	C C		



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Freedom Of Information (FOI) Report

Date:11/03/97ISR Number: 3025637-6Report Type:Periodic Company Report #A0052988  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG / TWICE	Dry Mouth	Consumer	Wellbutrin	PS		ORAL
	PER DAY /	Feeling Abnormal					
	ORAL	Lethargy					
		Paraesthesia					

Date:11/03/97ISR Number: 3025641-8Report Type:Periodic Company Report #A0053021  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG / PER	Blood Pressure Increased	Health	Wellbutrin	PS		ORAL
	DAY/ ORAL	Weight Decreased	Professional				
				Codimal Dm	C		
				Loratadine	C		

Date:11/03/97ISR Number: 3025644-3Report Type:Periodic Company Report #A0053026  
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	ORAL	Hyperventilation	Health	Wellbutrin	PS		ORAL
		Snoring	Professional				

Date:11/03/97ISR Number: 3025650-9Report Type:Periodic Company Report #A0053069  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	ORAL	Dermatitis	Health	Wellbutrin	PS		ORAL

Date:11/03/97ISR Number: 3025654-6Report Type:Periodic Company Report #A0053087  
 Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Wellbutrin	PS		ORAL
ORAL			Professional	Wellbutrin	SS		ORAL
150 MG TWICE							
PER DAY ORAL							

Date:11/03/97ISR Number: 3025656-XReport Type:Periodic Company Report #A0053150  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Wellbutrin	PS		ORAL
150 MG/ PER		Hyperhidrosis					
DAY / ORAL		Insomnia		Alprazolam	C		

Date:11/03/97ISR Number: 3025660-1Report Type:Periodic Company Report #A0053152  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Jittery	Health	Wellbutrin	PS		ORAL
150MG / PER		Insomnia	Professional				
DAY / ORAL		Nausea					

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Freedom Of Information (FOI) Report

Date:11/03/97ISR Number: 3025663-7Report Type:Periodic Company Report #A0053182  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus	Health	Wellbutrin	PS		ORAL
ORAL			Professional Company Representative				

Date:11/03/97ISR Number: 3025668-6Report Type:Periodic Company Report #A0053203  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tachycardia	Health	Wellbutrin	PS		ORAL
150MG / TWICE			Professional				
PER DAY /			Company				
ORAL			Representative				

Date:11/03/97ISR Number: 3025671-6Report Type:Periodic Company Report #A0053307  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Clonic Convulsion	Health	Wellbutrin	PS		ORAL
150MG / THREE			Professional				
TIMES PER DAY			Company				
/ ORAL			Representative	Paroxetine Hydrochloride	C		
				Semisodium Valproate	C		

Date:11/03/97ISR Number: 3025674-1Report Type:Periodic Company Report #A0053396  
 Age: Gender:Unknown I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Dermatitis	Health Professional Company Representative	Wellbutrin	PS		ORAL

Date:11/03/97ISR Number: 3025677-7Report Type:Periodic Company Report #A0053400  
 Age:38 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG / TWICE PER DAY / ORAL		Urticaria	Consumer	Wellbutrin	PS		ORAL

Date:11/03/97ISR Number: 3025680-7Report Type:Periodic Company Report #A0053474  
 Age:83 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG / TWICE Other PER DAY / ORAL		Tremor	Health Professional Company Representative	Wellbutrin	PS		ORAL

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Freedom Of Information (FOI) Report

Date:11/03/97ISR Number: 3025683-2Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #A0053475

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Health	Wellbutrin	PS		ORAL
200 MG /			Professional				
TWICE PER DAY							
/ ORAL				Thyroxine Sodium	C		
				Oral Contraceptive	C		

Date:11/03/97ISR Number: 3025686-8Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #A0053520

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Health	Wellbutrin	PS		ORAL
ORAL		Coordination Abnormal	Professional				
		Malaise	Company Representative				

Date:11/03/97ISR Number: 3025690-XReport Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0053528

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Deafness	Health	Wellbutrin	PS		ORAL
ORAL		Tinnitus	Professional				

Date:11/03/97ISR Number: 3025694-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0053589

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flatulence	Consumer	Wellbutrin	PS		ORAL
150MG / PER							

DAY / ORAL

Date:11/03/97ISR Number: 3025696-0Report Type:Periodic Company Report #A0053644  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyskinesia	Health	Wellbutrin	PS		ORAL
150MG / TWICE			Professional				
PER DAY /							
ORAL							

Date:11/03/97ISR Number: 3025706-0Report Type:Periodic Company Report #A0053701  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Health	Wellbutrin	PS		ORAL
150MG / TWICE		Balance Disorder	Professional				
PER DAY /		Dermatitis					
ORAL		Dyspnoea		Triamcinolone			
		Headache		Acetonide	C		
		Nausea		Paracetamol	C		
		Pharyngeal Oedema		Lopoeramide			
				Hydrochloride	C		

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Date:11/03/97ISR Number: 3025709-6Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #A0053777

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
75 MG / THREE		Tooth Discolouration	Consumer	Wellbutrin	PS		ORAL
TIMES PER DAY							
/ ORAL							

Estrogen C  
Atenolol C

Date:11/03/97ISR Number: 3025712-6Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0053779

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
ORAL	18 MON	Alopecia	Consumer	Wellbutrin	PS		ORAL

Medroxyprogesterone  
Ace. C  
Nizatidine C

Date:11/03/97ISR Number: 3025714-XReport Type:Periodic  
Age:25 YR Gender:Female I/FU:I

Company Report #A0053780

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG / THREE		Grand Mal Convulsion	Health	Wellbutrin	PS		ORAL
TIMES PER DAY			Professional				
/ ORAL			Company				
			Representative				

Date:11/03/97ISR Number: 3025717-5Report Type:Periodic  
Age:71 YR Gender:Male I/FU:I

Company Report #A0053781

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Health	Wellbutrin	PS		ORAL
150MG / PER		Gastrointestinal	Professional				
DAY / ORAL		Obstruction		Salbutamol	Sulphate	C	
		Nausea		Famotidine		C	

Date:11/03/97ISR Number: 3025719-9Report Type:Periodic Company Report #A0053803  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Periorbital Oedema	Health	Wellbutrin	PS		ORAL
ORAL			Professional Company Representative				

Date:11/03/97ISR Number: 3025721-7Report Type:Periodic Company Report #A0053804  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral	Health	Wellbutrin	PS		ORAL
ORAL			Professional Company Representative				



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Date:11/03/97ISR Number: 3025723-0Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0053873

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Generalised	Consumer	Wellbutrin	PS		ORAL
TWICE PER DAY/ ORAL;							
THREE TIMES PER DAY ORAL							

Date:11/03/97ISR Number: 3025725-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0054003

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Galactorrhoea	Health	Wellbutrin	PS		ORAL
ORAL							
			Professional	Oxazepam	C		

Date:11/03/97ISR Number: 3025727-8Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0054026

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Health	Wellbutrin	PS		ORAL
ORAL							
			Professional Company Representative				

Date:11/03/97ISR Number: 3025730-8Report Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #A0054027

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Clonic Convulsion	Health	Wellbutrin	PS		ORAL
Hospitalization - 150MG / PER							

Initial or Prolonged      Coordination Abnormal      Professional  
DAY / ORAL

Fall  
Hyperreflexia  
Tremor  
Vertigo

Date:11/03/97ISR Number: 3025733-3Report Type:Periodic      Company Report #A0054028  
Age:      Gender:Unknown      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral	Health	Wellbutrin	PS		ORAL
300MG / PER			Professional				
DAY / ORAL							

Date:11/03/97ISR Number: 3025734-5Report Type:Periodic      Company Report #A0054059  
Age:      Gender:Male      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Health	Wellbutrin	PS		ORAL
ORAL		Nausea Vomiting	Professional	Venlafaxine Hydrochloride Lithium Salt	SS C		

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Date:11/03/97ISR Number: 3025737-0Report Type:Periodic Company Report #A0054092  
 Age:52 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG / TWICE PER DAY / ORAL		Influenza Like Illness	Health Professional	Wellbutrin	PS		ORAL

Date:11/03/97ISR Number: 3025740-0Report Type:Periodic Company Report #A0054112  
 Age: Gender:Unknown I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Anxiety Feeling Jittery	Health Professional Company Representative	Wellbutrin	PS		ORAL

Date:11/03/97ISR Number: 3025742-4Report Type:Periodic Company Report #A0054113  
 Age: Gender:Unknown I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Anxiety Feeling Jittery	Health Professional Company Representative	Wellbutrin	PS		ORAL

Date:11/03/97ISR Number: 3025745-XReport Type:Periodic Company Report #A0054153  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Dyspnoea	Health	Wellbutrin	PS		ORAL

Professional  
Company  
Representative

Date:11/03/97ISR Number: 3025748-5Report Type:Periodic Company Report #A0054157  
Age:39 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Breast Pain	Health	Wellbutrin	PS		ORAL
100 MG /			Professional				
TWICE PER DAY							
/ ORAL				..	C		

Date:11/03/97ISR Number: 3025750-3Report Type:Periodic Company Report #A0054204  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Health	Wellbutrin	PS		ORAL
ORAL			Professional				

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Date:11/03/97ISR Number: 3025752-7Report Type:Periodic Company Report #A0054205  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG /TWICE	Agitation	Consumer	Wellbutrin	PS		ORAL
	PER DAY /	Cough					
	ORAL	Headache					
		Nervousness					

Date:11/03/97ISR Number: 3025754-0Report Type:Periodic Company Report #A0054222  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	ORAL	Alopecia	Health	Wellbutrin	PS		ORAL
			Professional				

Date:11/03/97ISR Number: 3025757-6Report Type:Periodic Company Report #A0054280  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	ORAL	Dermatitis	Health	Wellbutrin	PS		ORAL
			Professional				
			Company Representative				

Date:11/03/97ISR Number: 3025760-6Report Type:Periodic Company Report #A0054281  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	ORAL	Dermatitis	Health	Wellbutrin	PS		ORAL
			Professional				

Date:11/03/97ISR Number: 3025762-XReport Type:Periodic Company Report #A0054300  
Age:33 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Adnexa Uteri Pain Pelvic Pain Tenderness	Consumer	Wellbutrin	PS		ORAL
75 MG / AT NIGHT / ORAL				Wellbutrin	SS		ORAL
				Temazepam	C		

Date:11/03/97ISR Number: 3025764-3Report Type:Periodic Company Report #A0054302  
Age:25 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG / TWICE PER DAY / ORAL		Tinnitus	Health Professional	Wellbutrin	PS		ORAL

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Freedom Of Information (FOI) Report

Date:11/03/97ISR Number: 3025767-9Report Type:Periodic Company Report #A0054303  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Health	Wellbutrin	PS		ORAL
150 MG. ORAL			Professional				

Date:11/03/97ISR Number: 3025770-9Report Type:Periodic Company Report #A0054304  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Faeces	Consumer	Wellbutrin	PS		ORAL
100 MG / FOUR		Colitis Ulcerative					
TIMES PER		Faecal Occult Blood					
DAY/ ORAL		Positive		Clonazepam	C		
				Mesalazine	C		

Date:11/03/97ISR Number: 3025772-2Report Type:Periodic Company Report #A0054361  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphagia	Consumer	Wellbutrin	PS		ORAL
150 MG / ORAL		Glossodynia					

Date:11/03/97ISR Number: 3025775-8Report Type:Periodic Company Report #A0054369  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Health	Wellbutrin	PS		ORAL
ORAL	MON	Rash Macular	Professional				

Date:11/03/97ISR Number: 3025777-1Report Type:Periodic Company Report #A0054445  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Health	Wellbutrin	PS		ORAL
ORAL			Professional Company Representative				

Date:11/03/97ISR Number: 3025779-5Report Type:Periodic Company Report #A0054446  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Wellbutrin	PS		ORAL
150 MG /		Drug Ineffective					
TWICE PER		Tobacco Abuse					
DAY/ ORAL				Blood Pressure Medication	C		

Date:11/03/97ISR Number: 3025781-3Report Type:Periodic Company Report #A0054447  
Age:38 YR Gender:Female I/FU:I

Outcome	PT	Report Source
Life-Threatening	Grand Mal Convulsion	Health Professional



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			Company Representative	Product	Role	Manufacturer	Route
Dose	Duration			Wellbutrin	PS		
150 MG / FOUR							
TIMES PER DAY							

Date:11/03/97ISR Number: 3025783-7Report Type:Periodic Company Report #A0054450  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Health	Wellbutrin	PS		ORAL
150 MG / PER		Dry Mouth	Professional				
DAY/ ORAL		Paraesthesia		Venlafaxine			
		Visual Disturbance		Hydrochloride	C		

Date:11/03/97ISR Number: 3025784-9Report Type:Periodic Company Report #A0054510  
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flushing	Consumer	Wellbutrin	PS		ORAL
150 MG /		Insomnia					
TWICE PER				Nicotine Patch	SS		
DAY/ ORAL				Prempro	C		
TOPICAL							

Date:11/03/97ISR Number: 3025785-0Report Type:Periodic Company Report #A0054528  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

VARIABLE DOSE	Constipation	Consumer	Wellbutrin	PS	ORAL
/ ORAL	Drug Ineffective				
	Elevated Mood		Imipramine	C	
	Fatigue		Hydrochloride		
	Sedation				

Date:11/03/97ISR Number: 3025786-2Report Type:Periodic Company Report #A0054542  
 Age:57 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dry Mouth	Consumer	Wellbutrin	PS		ORAL
TWICE PER DAY		Tongue Coated					
/ ORAL		Tongue Disorder					

Date:11/03/97ISR Number: 3025789-8Report Type:Periodic Company Report #A0054544  
 Age:42 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dizziness	Health	Wellbutrin	PS		ORAL
TWICE PER DAY		Nausea	Professional				
/ ORAL	4 MON	Vomiting					

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Date:11/03/97ISR Number: 3025791-6Report Type:Periodic Company Report #A0054545  
 Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
ORAL	Tremor	Health	Wellbutrin	PS		ORAL
		Professional				

Date:11/03/97ISR Number: 3025792-8Report Type:Periodic Company Report #A0054565  
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
ORAL	Rash Pruritic	Health	Wellbutrin	PS		ORAL
		Professional				

Date:11/03/97ISR Number: 3025793-XReport Type:Periodic Company Report #A0054657  
 Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
ORAL	Dermatitis	Health	Wellbutrin	PS		ORAL
		Professional Company Representative				

Date:11/03/97ISR Number: 3025794-1Report Type:Periodic Company Report #A0054658  
 Age:11 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
150 MG / PER DAY/ ORAL	Tic	Consumer	Wellbutrin	PS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased	Consumer	Wellbutrin	PS		ORAL
150MG / TWICE		Hypertension					
PER DAY /		Migraine					
ORAL				Imitrex	SS		
				Thyroxine	C		
				Verapamil	C		
				Estratest	C		
				Prinzide	C		
				Beclomethasone			
				Dipropion	C		
				Salmeterol			
				Xinafoate	C		
				Fluoxetine			
				Hydrochloride	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Micturition Urgency	Health	Wellbutrin	PS		ORAL
150MG / TWICE		Pollakiuria	Professional				
PER DAY /							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/97ISR Number: 3025800-4Report Type:Periodic Company Report #A0054697  
 Age: Gender:Unknown I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /TWICE PER DAY/ ORAL		Drug Effect Decreased	Health Professional Company Representative	Wellbutrin	PS		ORAL

Date:11/03/97ISR Number: 3025801-6Report Type:Periodic Company Report #A0054716  
 Age:43 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG / PER DAY / ORAL		Dizziness Dry Mouth Dysphonia Dyspnoea Hypervigilance Insomnia Nausea Polydipsia	Consumer	Wellbutrin Fluvoxamine Maleate Alesse	PS C C		ORAL

Date:11/03/97ISR Number: 3025803-XReport Type:Periodic Company Report #A0054728  
 Age:19 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG / THREE TIMES PER DAY/ ORAL	4 YR	Amnesia Anorexia Asthenia Coordination Abnormal Depression Disorientation Dissociation	Consumer	Wellbutrin Lithium Salt Buspirone Hydrochloride Carbamazepine	PS C C C C		ORAL

Dizziness  
 Dysphemia  
 Fatigue  
 Feeling Jittery  
 Headache  
 Impaired Healing  
 Influenza  
 Muscle Spasms  
 Nervousness  
 Thinking Abnormal  
 Vision Blurred

Date:11/03/97ISR Number: 3025806-5Report Type:Periodic Company Report #A0054744  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG / PER		Anaphylactic Reaction	Health	Wellbutrin	PS		ORAL
DAY/ ORAL		Face Oedema	Professional				
		Pruritus	Company				
		Urticaria	Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/97ISR Number: 3025807-7Report Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #A0054745

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Pruritic	Health	Wellbutrin	PS		ORAL
150MG / TWICE			Professional				
PER DAY/ ORAL				Temazepam	C		
				Lorazepam	C		

Date:11/03/97ISR Number: 3025809-0Report Type:Periodic  
Age:69 YR Gender:Male I/FU:I

Company Report #A0054747

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Health	Wellbutrin	PS		ORAL
ORAL			Professional	Wellbutrin	SS		ORAL
450MG / PER							
DAY / ORAL							

Date:11/03/97ISR Number: 3025811-9Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0054755

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Effect Decreased	Health	Wellbutrin	PS		ORAL
150MG / TWICE			Professional				
PER DAY /			Company				
ORAL			Representative				

Date:11/03/97ISR Number: 3025812-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0054756

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Effect Decreased	Health	Wellbutrin	PS		ORAL
150MG / TWICE			Professional				
PER DAY /			Company				
ORAL			Representative				

Date:11/03/97ISR Number: 3025813-2Report Type:Periodic Company Report #A0054772  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Extrapyramidal Disorder	Health	Wellbutrin	PS		ORAL
ORAL			Professional				
			Company				
			Representative				

Date:11/03/97ISR Number: 3025814-4Report Type:Periodic Company Report #A0054794  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea	Health	Wellbutrin	PS		ORAL
100MG / THREE		Rectal Haemorrhage	Professional				
TIMES PER DAY							
/ ORAL				Doxazosin Mesylate	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/97ISR Number: 3025815-6Report Type:Periodic Company Report #A0054796  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Wellbutrin	PS		ORAL
150MG / TWICE		Drug Ineffective					
PER DAY/ ORAL		Dry Mouth					

Date:11/03/97ISR Number: 3025816-8Report Type:Periodic Company Report #A0054806  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Health	Wellbutrin	PS		ORAL
ORAL		Fibromyalgia	Professional				

Date:11/03/97ISR Number: 3025817-XReport Type:Periodic Company Report #A0054819  
 Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Health	Wellbutrin	PS		ORAL
150MG / TWICE		Convulsion	Professional				
PER DAY/ ORAL		Depression	Company Representative				

Date:11/03/97ISR Number: 3025818-1Report Type:Periodic Company Report #A0054822  
 Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion	Health	Wellbutrin	PS		ORAL
200MG / TWICE			Professional				
PER DAY /							

ORAL

Company  
Representative

Date:11/03/97ISR Number: 3025819-3Report Type:Periodic Company Report #A0054824  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Wellbutrin	PS		ORAL
100 MG /		Urticaria					
TWICE PER DAY							
/ ORAL				Vanquish	C		
				Paracetamol	C		

Date:11/03/97ISR Number: 3025820-XReport Type:Periodic Company Report #A0054831  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Wellbutrin	PS		ORAL
ORAL			Professional Company Representative				



150MG /TWICE  
Drug Ineffective  
Consumer  
Wellbutrin  
PS  
ORAL  
PER DAY /  
Vaginal Haemorrhage  
ORAL

Conjugated Estrogens  
C  
Date:11/03/97ISR Number: 3025825-9Report Type:Periodic Company Report #A0054971  
Age: Gender:Unknown I/FU:I  
Outcome Dose Duration PT Report Source Product Role Manufacturer Route  
Convulsion Health Wellbutrin PS ORAL  
ORAL  
Professional Company Representative

Date:11/03/97ISR Number: 3025826-0Report Type:Periodic Company Report #A0055045  
Age:16 YR Gender:Female I/FU:I  
Outcome Dose Duration PT Report Source Product Role Manufacturer Route  
Alopecia Health Wellbutrin PS ORAL  
150MG / TWICE  
PER DAY /  
ORAL  
Professional  
Fluoxetine Hydrochloride  
C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/97ISR Number: 3025827-2Report Type:Periodic Company Report #A0055056  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Health	Wellbutrin	PS		ORAL
150MG / ORAL			Professional Company Representative				

Date:11/03/97ISR Number: 3025828-4Report Type:Periodic Company Report #A0055066  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Health	Wellbutrin	PS		ORAL
150MG / TWICE		Menstrual Disorder	Professional Company Representative				
PER DAY /							
ORAL							

Date:11/03/97ISR Number: 3025829-6Report Type:Periodic Company Report #A0055078  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Wellbutrin	PS		ORAL
ORAL			Professional Company Representative				

Date:11/03/97ISR Number: 3025830-2Report Type:Periodic Company Report #A0055079  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

ORAL                      Dermatitis                      Health                      Wellbutrin                      PS                      ORAL

Professional  
Company  
Representative

Date:11/03/97ISR Number: 3025831-4Report Type:Periodic                      Company Report #A0055092  
Age:53 YR    Gender:Male                      I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Abdominal Pain	Consumer	Wellbutrin	PS		ORAL
TWICE PER		Anxiety					
DAY / ORAL		Flatulence					
		Insomnia					

Date:11/03/97ISR Number: 3025832-6Report Type:Periodic                      Company Report #A0055101  
Age:                      Gender:Female                      I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Amenorrhoea	Health	Wellbutrin	PS		ORAL
			Professional Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/97ISR Number: 3025833-8Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0055133

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Wellbutrin	PS		ORAL
ORAL		Skin Odour Abnormal		..	C		

Date:11/03/97ISR Number: 3025834-XReport Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #A0055134

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Wellbutrin	PS		ORAL
ORAL		Skin Odour Abnormal		Hormones	C		
				Sedapap	C		
				Vitamin	C		
				Lorazepam	C		
				Diphenhydramine Hcl	C		

Date:11/03/97ISR Number: 3025835-1Report Type:Periodic  
Age:30 YR Gender: I/FU:I

Company Report #A0055135

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin	PS		ORAL
150 MG /		Convulsion	Professional				
TWICE PER DAY			Company				
/ ORAL			Representative				

Date:11/03/97ISR Number: 3025836-3Report Type:Periodic  
Age:66 YR Gender:Male I/FU:I

Company Report #A0055137

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG /	Cough	Consumer	Wellbutrin	PS	ORAL
TWICE PER DAY	Dyspnoea				
/ ORAL	Insomnia				
150 MG / ORAL			Zyban	SS	ORAL
			Fosinopril Sodium	C	
			Perphenazine	C	
			Cimetidine	C	
			Amoxapine	C	

Date:11/03/97ISR Number: 3025837-5Report Type:Periodic Company Report #A0055138  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER		Disorientation	Consumer	Wellbutrin	PS		ORAL
DAY / ORAL		Drug Ineffective					
		Memory Impairment		Carisoprodol	C		
		Nervousness					
		Paranoia					
		Psychomotor Hyperactivity					
		Psychotic Disorder					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/97ISR Number: 3025838-7Report Type:Periodic Company Report #A0055164  
 Age:45 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG/TWICE PER DAY/ ORAL		Dysgeusia Epistaxis Feeling Abnormal Tongue Coated	Consumer	Wellbutrin	PS		ORAL

Date:11/03/97ISR Number: 3025839-9Report Type:Periodic Company Report #A0055181  
 Age:35 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG /TWICE PER DAY / 100MG TWICE PER DAY/ORAL 100 MG / TWICE PER DAY /		Condition Aggravated Dermatitis Drug Effect Decreased Hypersensitivity Pruritus	Health Professional Company Representative	Wellbutrin    Wellbutrin	PS    SS		ORAL
				Valproic Acid	C		

Date:11/03/97ISR Number: 3025840-5Report Type:Periodic Company Report #A0055186  
 Age:44 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY		Condition Aggravated Crying	Consumer	Wellbutrin	PS		ORAL

/ ORAL	Depressed Mood				
	Insomnia		Estratest	C	
			Medroxyprogesterone		
			Ace.	C	

Date:11/03/97ISR Number: 3025841-7Report Type:Periodic Company Report #A0055195  
 Age:60 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Condition Aggravated	Health	Wellbutrin	PS		ORAL
150 MG /						
Initial or Prolonged	Convulsion	Professional				
TWICE PER DAY						
	Cough	Company				
/ ORAL						
	Grand Mal Convulsion	Representative	Sertraline			
	Loss Of Consciousness		Hydrochloride	C		
	Urinary Incontinence		Nortriptyline	C		
			Salmeterol Xinafoate	C		
			Flunisolide	C		
			Aspirin	C		
			Conjugated Estrogens	C		
			Clonidine	C		
			Doxepin	C		
			Prednisone	C		
			Buspirone			
			Hydrochloride	C		
			Corticosteroid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/97ISR Number: 3025843-0Report Type:Periodic Company Report #A0055252  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Wellbutrin	PS		ORAL
ORAL			Professional Company Representative				

Date:11/03/97ISR Number: 3025844-2Report Type:Periodic Company Report #A0055254  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Wellbutrin	PS		ORAL
100MG /TWICE		Nausea					
PER DAY /							
ORAL				Nicotine Patch	SS		
21MG /							
TOPICAL	1	MON					

Date:11/03/97ISR Number: 3025845-4Report Type:Periodic Company Report #A0055258  
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hair Texture Abnormal	Health	Wellbutrin	PS		ORAL
75 MG / PER			Professional				
DAY / ORAL							

Date:11/03/97ISR Number: 3025846-6Report Type:Periodic Company Report #A0055272  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lethargy	Consumer	Wellbutrin	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Clonazepam	C		

Date:11/03/97ISR Number: 3025847-8Report Type:Periodic Company Report #A0055303  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Myalgia	Health	Wellbutrin	PS		ORAL
150 MG /		Nausea	Professional				
TWICE PER DAY							
/ ORAL	5 WK						

Date:11/03/97ISR Number: 3025848-XReport Type:Periodic Company Report #A0055320  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fall	Health	Wellbutrin	PS		ORAL
150 MG / PER		Hyperhidrosis	Professional				
DAY / ORAL		Loss Of Consciousness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/97ISR Number: 3025849-1Report Type:Periodic Company Report #A0055431  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Wellbutrin	PS		ORAL
ORAL		Face Oedema	Professional Company Representative				

Date:11/03/97ISR Number: 3025850-8Report Type:Periodic Company Report #A0055432  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Wellbutrin	PS		ORAL
ORAL			Professional Company Representative				

Date:11/03/97ISR Number: 3025851-XReport Type:Periodic Company Report #A0052521  
 Age:22 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bulimia Nervosa	Health	Wellbutrin	PS		ORAL
Other		Grand Mal Convulsion	Professional				
150 MG TWICE				Wellbutrin	SS		ORAL
PER DAY ORAL							
225 MF PER							
DAY ORAL							

Date:11/05/97ISR Number: 100000187Report Type:Expedited (15-DaCompany Report #A0056490  
 Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death	Convulsion	Consumer	Zyban	PS	ORAL
150 MG /PER					
Hospitalization -	Dizziness				
DAY/ ORAL/					
Initial or Prolonged	Lethargy				
TAB					
	Loss Of Consciousness		Warfarin Sodium	C	
	Viral Infection				

Date:11/05/97ISR Number: 1999981-5Report Type:Expedited (15-DaCompany Report #A0056490  
Age:72 YR Gender:Male I/FU:U

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Consumer	Zyban	PS		ORAL
Hospitalization -		Dizziness		Warfarin Sodium	C		
Initial or Prolonged		Sedation					
		Syncope					

Date:11/05/97ISR Number: 3002688-9Report Type:Direct Company Report #  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Oedema Peripheral	Health	Zyban	PS		
150 MG BID		Pruritus	Professional				
		Rash Maculo-Papular					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/05/97ISR Number: 3005353-7Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Blister		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged	Rash Generalised Stevens-Johnson Syndrome					

Date:11/10/97ISR Number: 100000223Report Type:Expedited (15-DaCompany Report #A0055651  
 Age:40 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Amnesia	Consumer	Zyban	PS		ORAL
150 MG /TWICE						
Initial or Prolonged	Dysarthria					
PER DAY	Speech Disorder Syncope		Salbutamol Sulphate Prednisone Nisoldipine Beclomethasone Hydrochlorothiazide Theophyline Ipratropium Bromide	C C C C C C C		

Date:11/10/97ISR Number: 3005792-4Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Psoriasis		Zyban	PS		
150 MG QD 1 WK			Dovonex Ultravate Claritin Hydroxyzine Triphasil	C C C C C		

Date:11/17/97ISR Number: 3000899-XReport Type:Expedited (15-DaCompany Report #A0056815  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY/ORAL		Body Temperature  Decreased  Dyspnoea Hypersensitivity Hypotension Oedema Peripheral Po2 Decreased Pruritus Urticaria	Consumer	Zyban	PS		ORAL

Date:11/17/97ISR Number: 3000902-7Report Type:Expedited (15-DaCompany Report #A0056717  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150 MG/  TWICE PER DAY		Circulatory Collapse  Disorientation	Consumer	Zyban	PS		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/17/97ISR Number: 3001171-4Report Type:Expedited (15-DaCompany Report #A001-002-001429

Age:82 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Aricept	PS		ORAL
5 MG BID (2			Professional				
IN 1 D), PER							
ORAL							
				Wellbutrin	SS		ORAL
75 MG BID (2							
IN 1 D),PER							
ORAL							
				Accupril	C		
				Paxil	C		
				Aspirin	C		

Date:11/17/97ISR Number: 3001223-9Report Type:Expedited (15-DaCompany Report #A0055651

Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Aphasia	Health	Zyban	PS		ORAL
150 MG /							
Initial or Prolonged		Confusional State	Professional				
TWICE PER DAY							
Other		Dizziness					
/ ORAL							
		Dysarthria		Salbutamol Sulphate	C		
		Dysphemia		Prednisone	C		
		Feeling Jittery		Nisoldipine	C		
		Headache		Beclomethasone			
				Dipropion	C		
				Hydrochlorothiazide	C		
				Theophylline	C		
				Ipratropium Bromide	C		
				Salbutamol Sulphate	C		
				Beclomethasone			
				Dipropion	C		
				Alprazolam	C		
				Oxygen	C		

Date:11/18/97ISR Number: 3001455-XReport Type:Expedited (15-DaCompany Report #A0055872

Age:59 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Agranulocytosis	Health	Wellbutrin	PS		ORAL
ORAL; TAB						
Initial or Prolonged	Leukopenia	Professional	Fluoxetine Hydrochloride	C		

Date:11/19/97ISR Number: 3001611-0Report Type:Expedited (15-DaCompany Report #MPI-97418

Age:9 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Aggression	Health	Methylphenidate	PS		ORAL
55 MG						
Initial or Prolonged	Agitation	Professional	Clonidine Hcl	SS		ORAL
0.1 MG						
	Convulsion		Bupropion	SS		ORAL
150 MG						
	Disorientation					

Date:11/19/97ISR Number: 3001886-8Report Type:Expedited (15-DaCompany Report #MPI-97418

Age:9 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Aggression	Health	Methylphenidate	PS		ORAL
55MG PO						
Initial or Prolonged	Agitation	Professional	Clonidine Hcl	SS		ORAL
0.1MG PO						

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

150MG PO Bupropion SS ORAL

Date:11/20/97ISR Number: 3002248-XReport Type:Expedited (15-DaCompany Report #A0056001  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional Company Representative	Wellbutrin	PS		ORAL

Date:11/24/97ISR Number: 3005401-4Report Type:Direct Company Report #  
 Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Mydriasis	Health Professional	Wellbutrin	PS		ORAL

Date:11/25/97ISR Number: 3006144-3Report Type:Direct Company Report #  
 Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Angioneurotic Oedema		Zyban	PS		
(THERAPY		Arthralgia					
DURATION 2		Pyrexia					
WKS)	2 WK	Urticaria					
PRN				Advil Prn	SS		

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
I.CAVERN		Drug Ineffective	Consumer	Caverject	PS		
		Penile Pain	Company Representative	Wellbutrin St. John'S Wert	SS C		

Date:11/26/97ISR Number: 3002376-9Report Type:Expedited (15-DaCompany Report #A0054799

Age:69 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY  / ORAL		Abdominal Pain Upper  Gastritis  Haematochezia  Nausea	Health  Professional	Zyban    Verapamil Hydrochloride Indomethacin Vitamin Vicodin	PS    C C C C		ORAL

Date:11/26/97ISR Number: 3002381-2Report Type:Expedited (15-DaCompany Report #A0055872

Age:54 YR Gender:Female I/FU:F

Outcome  
Hospitalization -  
Initial or Prolonged

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Other

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG/TWICE		Agranulocytosis	Health	Wellbutrin	PS		ORAL
PER DAY/ORAL		Leukopenia	Professional				
				Fluoxetine			
				Hydrochloride	C		
				Trazodone	C		
				Minoxidil	C		
				Phenoxymethylpenicil			
				lin K	C		

Date:11/26/97ISR Number: 3002382-4Report Type:Expedited (15-DaCompany Report #A0054301  
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Deep Vein Thrombosis	Foreign	Wellbutrin	PS		ORAL
75MG/THREE			Study				
Initial or Prolonged			Health				
TIMES PER			Professional				
DAY/ ORAL							

Date:12/01/97ISR Number: 3006230-8Report Type:Direct Company Report #  
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Accommodation Disorder		Zyban	PS		ORAL
150 MG BID PO							
Initial or Prolonged		Balance Disorder					
		Dizziness					
		Electrocardiogram					
		Abnormal					
		Headache					
		Nausea					
		Vision Blurred					
		Vomiting					

Date:12/01/97ISR Number: 3050739-8Report Type:Periodic Company Report #A0055439  
Age:49 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY  / ORAL	Myocardial Infarction	Health  Professional	Zyban	PS		ORAL
			Human Int / Long Insulin	C		

Date:12/01/97ISR Number: 3050767-2Report Type:Periodic Company Report #A0055476  
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL Initial or Prolonged	Pancytopenia	Health  Professional Company Representative	Zyban	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/01/97ISR Number: 3050885-9Report Type:Periodic  
Age:76 YR Gender:Female I/FU:I

Company Report #A0055715

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban	PS		ORAL
Other		Dermatitis	Professional				
150 MG TWICE							
PER DAY/							
ORAL							

Date:12/01/97ISR Number: 3050924-5Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #A0055749

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban	PS		ORAL
Life-Threatening		Abnormal Behaviour	Professional				
150 MG/ PER							
DAY/ ORAL							

Date:12/01/97ISR Number: 3050931-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0055753

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban	PS		ORAL
Other		Agitation	Professional				
150 MG/		Anxiety					
TWICE PER		Derealisation					
DAY/ ORAL		Drug Level Above		Wellbutrin	SS		ORAL
150 MG/		Therapeutic					
THREE TIMES							
PER DAY/							
ORAL							

Date:12/01/97ISR Number: 3051356-6Report Type:Periodic Company Report #A0053959  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema	Health	Zyban	PS		ORAL
150 MG TWICE							
PER DAY/		Urticaria	Professional				
ORAL				Oxpentifylline	C		

Date:12/01/97ISR Number: 3051424-9Report Type:Periodic Company Report #A0054056  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Consumer	Zyban	PS		ORAL
300MG / SEE							
TEXT / ORAL		Feeling Abnormal					
		Feeling Drunk		Antibiotics	C		
		Medication Error					
		Tremor					
		Vision Blurred					

Date:12/01/97ISR Number: 3051746-1Report Type:Periodic Company Report #A0054114  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion	Health	Zyban	PS		ORAL
150 MG/TWICE							
Initial or Prolonged		Dizziness	Professional				
PER DAY/ORAL							
Other		Faecal Incontinence		Baclofen	C		
		Loss Of Consciousness		Misoprostol	C		
		Nausea		Omeprazole	C		
		Urinary Incontinence		Nabumetone	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Zolpidem Tartrate C  
 Venlafaxine  
 Hydrochloride C  
 Semisodium Valproate C  
 Desamethasone C  
 Clarithromycin C

Date:12/01/97ISR Number: 3051800-4Report Type:Periodic Company Report #A0054881  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG ORAL	Anaphylactic Reaction	Health	Zyban	PS		ORAL
Other		Dyspnoea Throat Tightness Tongue Disorder	Professional Company Representative	Oestradiol	C		

Date:12/01/97ISR Number: 3052148-4Report Type:Periodic Company Report #A0053088  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG /	Chest Discomfort	Health	Zyban	PS		ORAL
Initial or Prolonged	TWICE PER DAY	Costochondritis	Professional				
/ ORAL				Sertraline Hydrochloride	C		
				Thyroxine Sodium	C		
				Dexfenfluramine	C		
				Atorvastatin Calcium	C		

Date:12/01/97ISR Number: 3052150-2Report Type:Periodic Company Report #A0053159  
 Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other ORAL	Dyspnoea	Health	Zyban	PS	ORAL	
	Pruritus Urticaria	Professional				
Date:12/01/97ISR Number: 3052152-6Report Type:Periodic Company Report #A0053181						
Age:58 YR Gender:Female I/FU:F						
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Disorientation	Health	Zyban	PS		ORAL
150 MG / PER	Disturbance In Attention	Professional				
DAY / ORAL	Dizziness		Lisinopril	C		
	Feeling Abnormal		Diazepam	C		
			Paracetamol	C		

Date:12/01/97ISR Number: 3052155-1Report Type:Periodic Company Report #A0053466						
Age:54 YR Gender:Female I/FU:F						
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Cardiac Disorder	Health	Zyban	PS		ORAL
150 MG / ORAL						
Initial or Prolonged	Chest Pain	Professional	Digoxin	C		
	Convulsion		Conjugated Estrogens	C		
			Medroxyprogesterone			
			Ace.	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Ipratropium Bromide C  
 Salbutamol Sulphate C  
 Fluticasone  
 Propionate C  
 Triamcinolone  
 Acetonide C  
 Diphenhydramine Hcl C

Date:12/01/97ISR Number: 3052175-7Report Type:Periodic Company Report #A0056234  
 Age:60 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / PER Initial or Prolonged DAY / ORAL Disability Other	Mania	Health  Professional  Company Representative	Zyban   Lamotrigine Loarazepam Thiothixene Temazepam Lisinopril Ibuprofen Oestradiol	PS   C C C C C C		ORAL

Date:12/01/97ISR Number: 3052191-5Report Type:Periodic Company Report #A0056289  
 Age:43 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY  / ORAL	5 WK Dermatitis Dry Mouth Face Oedema Oral Mucosal Eruption Stevens-Johnson Syndrome Tongue Oedema	Consumer	Zyban	PS		ORAL

Date:12/01/97ISR Number: 3053025-5Report Type:Periodic Company Report #A0053833  
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Dizziness	Health	Zyban	PS		ORAL
150 MG TWICE							
Other		Pruritus	Professional				
PER DAY/							
		Swelling	Company				
ORAL		Urticaria	Representative				
		Vomiting					

Date:12/01/97ISR Number: 3053053-XReport Type:Periodic Company Report #A0053894  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Dermatitis	Health	Zyban	PS		ORAL
150 MG ORAL							
Initial or Prolonged		Urticaria	Professional				
			Company				
			Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/01/97ISR Number: 3053387-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0056510

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / ORAL	Hypersensitivity	Consumer	Zyban	PS		ORAL
Initial or Prolonged	Pruritus Rash Erythematous Swelling Urticaria		Decongestant	C		

Date:12/01/97ISR Number: 3053388-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0056511

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / ORAL	Hypersensitivity	Consumer	Zyban	PS		ORAL
Initial or Prolonged						

Date:12/01/97ISR Number: 3053553-2Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #A0054981

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG/TWICE	Agitation	Consumer	Zyban	PS		ORAL
PER DAY/ORAL	Excitability Overdose					

Date:12/01/97ISR Number: 3053602-1Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #A0056364

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG /	Agitation	Health	Zyban	PS		ORAL
TWICE PER	Hypertension	Professional				

DAY/ ORAL                      Nausea                      Company  
                                 Psychomotor Hyperactivity                      Representative                      Multiple Medication                      C

Date:12/01/97ISR Number: 3054081-0Report Type:Periodic                      Company Report #A0056760  
Age:                      Gender:Unknown                      I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose                      Duration Hospitalization - 150 MG; ORAL Initial or Prolonged Other	Syncope	Health  Professional Company Representative	Zyban	PS		ORAL

Date:12/01/97ISR Number: 3054192-XReport Type:Periodic                      Company Report #A0055144  
Age:44 YR                      Gender:Male                      I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose                      Duration Hospitalization - 150 MG/ Initial or Prolonged TWICE PER Other DAY/ ORAL	Grand Mal Convulsion	Health  Professional Company Representative	Zyban	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/01/97ISR Number: 3054235-3Report Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #A0055214

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Health	Zyban	PS		ORAL
150 MG/ TWICE		Convulsion	Professional				
PER DAY/		Dizziness					
ORAL	2 WK	Haematoma		Ciprofloxacin	C		
		Laceration		Loestrin	C		
		Loss Of Consciousness		Nicotine	C		
		Syncope					
		Tooth Injury					
		Urinary Tract Infection					

Date:12/01/97ISR Number: 3055015-5Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0055397

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eyelid Oedema	Health	Zyban	PS		ORAL
150 MG TWICE		Hypersensitivity	Professional				
PER DAY ORAL		Joint Swelling		Vitamin E	C		
		Rash Pruritic		Ascorbic Acid	C		
				Cyanocobalamin	C		
				Aspirin	C		

Date:12/01/97ISR Number: 3058639-4Report Type:Periodic  
Age:35 YR Gender:Male I/FU:I

Company Report #A0055974

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Hypersensitivity	Health	Zyban	PS		ORAL
150 MG ORAL		Oedema	Professional	Acyclovir	C		
Other		Pruritus					
		Rash Erythematous					

Date:12/02/97ISR Number: 3007014-7Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Urticaria		Zyban	PS		

Date:12/03/97ISR Number: 3006683-5Report Type:Direct  
Age:44 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Depression		Wellbutrin	PS	Burroughs Wellcome	ORAL
150 MG QID PO 1 WK							
Initial or Prolonged		Hypomania		Gabapentin	C		
		Mania					
		Suicidal Ideation					

Date:12/04/97ISR Number: 3007173-6Report Type:Direct  
Age:15 YR Gender:Female I/FU:I

Company Report #

Outcome  
Hospitalization -  
Initial or Prolonged  
Required  
Intervention to

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Prevent Permanent  
Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG BID PO		Angioneurotic Oedema		Wellbutrin	PS	Glaxo Wellcome	ORAL
1 MG BID PO		Arthralgia		Tenex	SS		ORAL
100 MG BID		Chest Pain		Doxycycline	C		
		Convulsion					
		Cyanosis					
		Diarrhoea					
		Dyspnoea					
		Face Oedema					
		Loss Of Consciousness					
		Nausea					
		Oedema Peripheral					
		Pain In Extremity					
		Rash Pruritic					
		Throat Tightness					

Date:12/04/97ISR Number: 3007606-5Report Type:Direct  
Age:35 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG 1 TAB Initial or Prolonged PO Q AM 16		Asthenia		Wellbutrin Sr	PS		ORAL
DOSES		Dizziness					
(SAMPLE)		Dyskinesia					
		Hypoaesthesia					
		Paraesthesia					

Date:12/10/97ISR Number: 3006294-1Report Type:Expedited (15-DaCompany Report #A0057764  
Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL	Chest Pain  Dermatitis  Hypersensitivity Oedema Oedema Peripheral Pruritus	Consumer	Zyban	PS	ORAL
			Hydrocortisone Warfarin Sodium	C C	

Date:12/10/97ISR Number: 3006295-3Report Type:Expedited (15-DaCompany Report #A0057792  
Age:58 YR Gender:Female I/FU:I

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ PER Initial or Prolonged DAY/ ORAL	Blood Sodium Decreased  Dehydration	Consumer	Zyban	PS		ORAL
	Polyuria Vomiting Weight Decreased		Oxpentifylline Nabumetone Claritin-D Lansoprazole Conjugated Estrogens Dyazide Simvastatin Alprazolam Amitriptyline Hcl Fluticasone Propionate	C C C C C C C C C C C		

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Freedom Of Information (FOI) Report

Date:12/10/97ISR Number: 3006296-5Report Type:Expedited (15-DaCompany Report #A0057783  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Foreign	Wellbutrin	PS		ORAL
150 MG/ THREE Life-Threatening TIMES PER DAY			Study				
ORAL			Health				
			Professional	Venlafaxine Hydrechloride	C		
				Lorazepam	C		
				Temazepam	C		

Date:12/11/97ISR Number: 3007417-0Report Type:Expedited (15-DaCompany Report #A0054873  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG ORAL		Anxiety	Consumer	Zyban	PS		ORAL
Initial or Prolonged		Asthenia		Theophylline	C		
		Cardiac Failure		Salmeterol Xinafoate	C		
		Congestive		Salbutamol Sulphate	C		
		Dizziness		Lactulose	C		
		Dry Mouth		Frusemide	C		
		Dysgeusia		Alprazolam	C		
		Headache					
		Insomnia					
		Nausea					
		Pneumonia					

Date:12/11/97ISR Number: 3007632-6Report Type:Direct Company Report #  
Age:5 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100 MG TID	1 DAY	Balance Disorder		Wellbutrin Sr	PS		
		Coordination Abnormal					

Date:12/15/97ISR Number: 3006983-9Report Type:Direct  
Age:26 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Iron Decreased	Other	Clozapine	PS		
12.5 MG BID		Microcytic Anaemia		Effexor	SS		
				Wellbutrin	SS		
				Klonopin	SS		

Date:12/15/97ISR Number: 3008811-4Report Type:Expedited (15-DaCompany Report #A0057391  
Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Drug Interaction	Health	Wellbutrin	PS		ORAL
UNK / TWICE							
Other		Headache	Professional				
PER DAY /		Hypersensitivity	Company				
ORAL			Representative	Trazodone	SS		ORAL
50 MG /							
SINGLE DOSE /							
ORAL							

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Freedom Of Information (FOI) Report

Date:12/17/97ISR Number: 3011549-0Report Type:Direct  
 Age:49 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blister		Zyban	PS		
AS DIRECTED	3 WK	Erythema Multiforme Urticaria					

Date:12/17/97ISR Number: 3011763-4Report Type:Direct  
 Age:19 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Bupropion	PS		ORAL
150MG PO BID Required Intervention to Prevent Permanent Impairment/Damage		Tremor					

Date:12/18/97ISR Number: 3009376-3Report Type:Expedited (15-DaCompany Report #A0057968  
 Age:34 YR Gender:Female I/FU:I

Company Report #A0057968

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ ORAL		Gait Disturbance Hypersensitivity Joint Swelling Pharyngeal Oedema Urticaria	Consumer	Wellbutrin	PS		ORAL

Date:12/18/97ISR Number: 3009383-0Report Type:Expedited (15-DaCompany Report #A0057990  
 Age:44 YR Gender:Female I/FU:I

Company Report #A0057990

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY ORAL	Arthralgia Bronchospasm Myalgia Oedema Pharyngeal Oedema Serum Sickness Urticaria	Health Professional	Zyban  Buspirone	PS  C	ORAL
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Date:12/19/97ISR Number: 3008836-9Report Type:Expedited (15-DaCompany Report #A0057263  
Age:16 YR Gender:Female I/FU:I

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG / THREE Initial or Prolonged TIMES PER DAY ORAL	Polyarthritits Serum Sickness Urticaria	Health Professional	Wellbutrin	PS		ORAL

Date:12/23/97ISR Number: 3014108-9Report Type:Expedited (15-DaCompany Report #A0056978  
Age:43 YR Gender:Male I/FU:I

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150 MG / Hospitalization - TWICE PER DAY Initial or Prolonged / ORAL Other	Pancreatitis	Health Professional Company Representative	Zyban	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/29/97ISR Number: 3014415-XReport Type:Expedited (15-DaCompany Report #A0058368

Age:75 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG/SINGLE	Blindness Unilateral	Consumer	Zyban	PS		ORAL
Initial or Prolonged DOSE/ORAL	Embolism					
	Rectal Haemorrhage		Terazosin Hydrochloride	C		
			Pravastatin Sodium	C		
			Aspirin	C		

Date:12/29/97ISR Number: 3014418-5Report Type:Expedited (15-DaCompany Report #A0054873

Age:64 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG/ UNK/	Anxiety	Health	Zyban	PS	Zyban	ORAL
Initial or Prolonged ORAL	Asthenia	Professional				
	Cardiac Failure		Theophylline	C		
	Congestive		Salmeterol Xinafoate	C		
	Dizziness		Salbutamol Sulphate	C		
	Dry Mouth		Lactulose	C		
	Dysgeusia		Frusemide	C		
	Headache		Alprazolam	C		
	Insomnia					
	Nausea					
	Pneumonia					
	Respiratory Failure					

Date:01/07/98ISR Number: 3015994-9Report Type:Expedited (15-DaCompany Report #A0057797

Age:66 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / PER	Blood Pressure Increased	Consumer	Zyban	PS		ORAL
Initial or Prolonged DAY/ ORAL	Dehydration					

Hypersomnia  
Malaise  
Nausea  
Vision Blurred

Conjugated Estrogens C  
Ibuprofen C

Date:01/08/98ISR Number: 3015442-9Report Type:Expedited (15-DaCompany Report #97030194-1  
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Nicoderm Cq	PS		
TRANSDERMAL	21 MILLIGRAMS	Feeling Abnormal					
1.0 DAILY		Loss Of Consciousness					
TRANSDERMAL		Movement Disorder		Zyban	SS		ORAL
150		Nervousness					
MILLIGRAMS							
2.0 DAILY							
ORAL							

Date:01/08/98ISR Number: 3016021-XReport Type:Expedited (15-DaCompany Report #8-97286-020L  
Age:43 YR Gender:Female I/FU:F

Outcome  
Other  
Required  
Intervention to

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Prevent Permanent  
Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
20MG THREE TIMES A DAY		Cardiac Valve Disease	Health	Pondimin	PS		ORAL
ORAL		Mitral Valve Incompetence	Professional				
30MG DAILY		Palpitations		Fastin	SS		ORAL
ORAL				Ionamin	SS		ORAL
30MG DAILY				Redux	SS		ORAL
ORAL				Wellbutrin	SS		
100 MG BID				Ionamin	C		
				Ionamin	C		
				Redux	C		

Date:01/12/98ISR Number: 3016979-9Report Type:Expedited (15-DaCompany Report #A0058622  
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/PER Initial or Prolonged DAY/ORAL		Lung Neoplasm Malignant	Consumer	Zyban	PS		ORAL
Other				Betaxolol Hydrochloride	C		
				Lorazepam	C		
				Theophylline	C		

Date:01/13/98ISR Number: 3016394-8Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Haemorrhage		Wellbutrin	PS		
300 MG QD							
		Shock		Zoloft	SS		
75 MG QD							

Date:01/13/98ISR Number: 3016645-XReport Type:Expedited (15-DaCompany Report #A0057378  
Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Coma	Study	Retrovir	PS		ORAL
300 MG /							
Initial or Prolonged		Intentional Misuse	Health				
TWICE PER DAY							
Other		Suicide Attempt	Professional				
/ ORAL							
				Epivir	SS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
800 MG /				Indinavir Sulfate	SS		ORAL
THREE TIMES							
PER DA / ORAL							
				Placebo	SS		
				Wellbutrin	SS		ORAL
SINGLE DOSE /							
ORAL							
				Trazodone	C		
				Bupropion			
				Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/15/98ISR Number: 3017125-8Report Type:Expedited (15-DaCompany Report #A0057990  
Age:44 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150 MG / Hospitalization - TWICE PER DAY Initial or Prolonged / ORAL Other	Arthralgia Bronchospasm Dyspnoea Feeling Hot Myalgia Oedema Serum Sickness Throat Tightness Urticaria	Health Professional	Zyban     Buspirone Hydrochloride	PS     C		ORAL

Date:01/15/98ISR Number: 3017130-1Report Type:Expedited (15-DaCompany Report #A0058368  
Age:75 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged SINGLE DOSE / ORAL	Blindness Unilateral Embolism Rectal Haemorrhage	Consumer	Zyban     Terazosin Hydrochloride Pravastatin Sodium Aspirin Allopurinol	PS     C C C C		ORAL

Date:01/15/98ISR Number: 3017133-7Report Type:Expedited (15-DaCompany Report #A0057783  
Age:47 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death 150 MG /	Completed Suicide	Foreign	Wellbutrin	PS		ORAL

Life-Threatening  
THREE TIMES  
  
PER DAY

Study  
  
Health  
  
Professional  
Venlafaxine  
Hydrochloride C  
Lorazepam C  
Temazepam C

Date:01/15/98ISR Number: 3017147-7Report Type:Expedited (15-DaCompany Report #A0057496  
Age:61 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG TWICE PER DAY ORAL	Heart Rate Decreased Hypertension	Health Professional	Zyban  Nadolol Hydroxychloroquine	PS  C C		ORAL

Date:01/15/98ISR Number: 3017151-9Report Type:Expedited (15-DaCompany Report #A0058633  
Age:62 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL	Blood Pressure Increased Cardiomegaly Hypertension Palpitations	Consumer	Zyban  Clonazepam Theophylline Hypericum Doxazosin	PS  C C C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Alprazolam C

Date:01/15/98ISR Number: 3017155-6Report Type:Expedited (15-DaCompany Report #A0058742  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL	1 WK		Health	Wellbutrin	PS		ORAL
Initial or Prolonged		Petechiae	Professional	Tetracycline	C		
		Stomatitis Haemorrhagic		Phenobarbitone	C		
		Thrombocytopenia					

Date:01/15/98ISR Number: 3077789-XReport Type:Periodic Company Report #WAES 97070306  
 Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gout	Consumer	Crixivan	PS	Merck Research Laboratories Div	
		Nervousness				Merck Co Inc	ORAL
		Tinnitus					
800 MG/Q8H/PO				Bupropion Hcl Unk	SS		
				Zoloft Unk	SS		
				Biaxin	C		
				DiFlucan	C		
				Floxin	C		
				Indocin	C		
				Lomotil	C		
				Restoril	C		
				Sporanox	C		
				Viramune	C		
				Xanax	C		
				Zerit	C		
				Zovirax	C		
				Zyloprim	C		
				Composition			
				Unspecified	C		
				Lamivudine	C		

Date:01/20/98ISR Number: 3017239-2Report Type:Expedited (15-DaCompany Report #A0059083  
Age:17 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL	Bradycardia	Health	Wellbutrin	PS		ORAL
Initial or Prolonged	Tachycardia	Professional Company Representative				

Date:01/20/98ISR Number: 3017242-2Report Type:Expedited (15-DaCompany Report #A0059063  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 75 MG/ORAL1	Liver Function Test	Health	Wellbutrin	PS		ORAL
Hospitalization - YEARS 1 YR	Abnormal	Professional				
Initial or Prolonged	Nausea Oedema Pancreatitis Pleural Effusion		Sertraline	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/20/98ISR Number: 3017286-0Report Type:Expedited (15-DaCompany Report #A0059062

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Zyban	PS		ORAL
ORAL		Smoker					

Date:01/20/98ISR Number: 3017288-4Report Type:Expedited (15-DaCompany Report #A0059118

Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Pressure Increased	Consumer	Zyban	PS		ORAL
ORAL		Dehydration					
Initial or Prolonged		Nausea					
		Vision Blurred					

Date:01/20/98ISR Number: 3018004-2Report Type:Expedited (15-DaCompany Report #A0057927

Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Condition Aggravated	Consumer	Wellbutrin	PS		ORAL
150 MG/ORAL		Dyspnoea		Vitamin E	C		
Initial or Prolonged		Oedema Peripheral		Multivitamin	C		
				Prempro	C		
				Prednisone	C		
				Ticlopidine	C		
				Metoclopramine	C		
				Nabumetone	C		
				Losartan Potassium	C		
				Terazosin	C		
				Hydrochloride	C		
				Diltiazem	C		
				Hydrochloride	C		
				Omeprazole	C		
				Insulin	C		

Date:01/20/98ISR Number: 3018609-9Report Type:Direct  
Age:58 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anorexia		Wiellbutrin	PS	Sr	
QD X 3 D THEN		Hyperhidrosis					
BID		Insomnia		Mivacor	C		
		Palpitations		Claritin	C		
				Premarin	C		
				Vasotec	C		

Date:01/20/98ISR Number: 3018611-7Report Type:Direct  
Age:40 YR Gender:Male I/FU:I

Company Report #

Outcome	PT
Hospitalization -	Angioneurotic Oedema
Initial or Prolonged	Arthralgia
	Asthenia
	Eyelid Oedema
	Fatigue
	Pyrexia
	Rash Generalised



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Rash Pruritic Tachycardia Urticaria	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Trimethoprim/Sulfame thoxazole	PS		ORAL
PO 5 DAYS							
PERIOD TO							
ADMISSION	5 DAY			Bupropion	SS		

Date:01/20/98ISR Number: 3083294-7Report Type:Periodic Company Report #A0053314  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aortic Valve Stenosis	Health	Wellbutrin Tablet	PS		ORAL
Other			Professional	Sertraline Hydrochloride (Formulation Unknown)	SS		ORAL
ORAL							
ORAL							

Date:01/20/98ISR Number: 3083775-6Report Type:Periodic Company Report #A0053886  
Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion	Health	Wellbutrin			
Hospitalization - Initial or Prolonged			Professional	(Formulation Unknown)	PS		ORAL
100 MG / FOUR							
TIMES PER DAY							
ORAL	1 MON						

Date:01/20/98ISR Number: 3083799-9Report Type:Periodic Company Report #A0054330  
Age:39 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ ORAL	Convulsion Loss Of Consciousness Psychotic Disorder	Health Professional	Wellbutrin Fluoxetine Hydrochloride Olanzapine	PS C C		ORAL

Date:01/20/98ISR Number: 3083884-1Report Type:Periodic Company Report #A0054792  
Age:55 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration 150 MG ORAL	Postmenopausal Haemorrhage	Consumer	Wellbutrin Tablet	PS		ORAL

Date:01/20/98ISR Number: 3083924-XReport Type:Periodic Company Report #A0050599  
Age:12 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration 75 MG TWICE PER DAY ORAL	Drug Interaction Dysphagia Erythema Multiforme Joint Swelling	Health Professional	Wellbutrin Tablet Clarithromycin (Formulation Unknown)	PS SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/20/98ISR Number: 3083927-5Report Type:Periodic  
Age:31 YR Gender:Male I/FU:I

Company Report #A0050600

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - ORAL	Duration Dermatitis	Health	Wellbutrin Tablet	PS		ORAL
Initial or Prolonged Other	Mucosal Inflammation Pyrexia Stevens-Johnson Syndrome	Professional Company Representative				

Date:01/20/98ISR Number: 3083949-4Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0050649

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other	Duration Asthma Pneumonia	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL

150 MG / IN  
THE MORNING /  
ORAL

Trazodone  
Hydrochloride C  
Semisodium Valproate C  
Prednisone C

Date:01/20/98ISR Number: 3084003-8Report Type:Periodic  
Age:18 YR Gender:Female I/FU:I

Company Report #A0051411

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Duration Convulsion	Health	Wellbutrin Tablet	PS		ORAL
450 MG / PER DAY / ORAL		Professional Company Representative				

Date:01/20/98ISR Number: 3084007-5Report Type:Periodic Company Report #A0051433  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 75 MG/PER		Chest Pain	Consumer	Wellbutrin	PS		ORAL
Initial or Prolonged DAY/ORAL		Hallucination					
		Hypertension		Alprazolam	C		
		Insomnia		Unspecified			
		Paranoia		Medication	C		
				Trazodone	C		

Date:01/20/98ISR Number: 3084109-3Report Type:Periodic Company Report #A0051790  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 200 MG /		Convulsion	Company	Wellbutrin Tablet	PS		ORAL
			Representative				

TWICE PER DAY

/ ORAL

Date:01/20/98ISR Number: 3084125-1Report Type:Periodic Company Report #A0051931  
Age:52 YR Gender:Female I/FU:I

Outcome	PT
Other	Abdominal Pain Upper Asthma

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Hospitalization - 100 MG ORAL Initial or Prolonged	Convulsion	Health	Wellbutrin	PS	ORAL
		Professional Company Representative	Bupropion Hydrochloride Disulfiram	C C	

Date:01/20/98ISR Number: 3084203-7Report Type:Periodic Company Report #A0052872  
 Age: Gender:Female I/FU:I

Outcome Dose Hospitalization - 100 MG / Initial or Prolonged THREE TIMES  PER DAY /  ORAL	Duration	PT Vomiting	Report Source Consumer	Product Wellbutrin Tablet	Role PS	Manufacturer	Route ORAL
	1	MON					

Date:01/20/98ISR Number: 3084244-XReport Type:Periodic Company Report #A0052930  
 Age: Gender: I/FU:I

Outcome Dose Other  ORAL	Duration	PT Anorexia Coordination Abnormal Facial Palsy  Muscle Rigidity Musculoskeletal Stiffness	Report Source Health Professional	Product Wellbutrin (Formulation Unknown)	Role PS	Manufacturer	Route ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/20/98ISR Number: 3085328-2Report Type:Periodic Company Report #A0049086  
 Age:61 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL Initial or Prolonged	Convulsion	Health  Professional Company Representative	Wellbutrin	PS		ORAL

Date:01/20/98ISR Number: 3085330-0Report Type:Periodic Company Report #A0049102  
 Age:60 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100MG/FOUR Initial or Prolonged TIMES PER DAY/ORAL	Grand Mal Convulsion	Health  Professional Company Representative	Wellbutrin	PS		ORAL

Date:01/20/98ISR Number: 3085350-6Report Type:Periodic Company Report #A0049148  
 Age:47 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 900 MG/PER Initial or Prolonged DAY/ORAL Other	Convulsion  Overdose	Health  Professional Company Representative	Wellbutrin  Alprazolam	PS  C		ORAL

Date:01/20/98ISR Number: 3085390-7Report Type:Periodic Company Report #A0049329  
 Age:20 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Hospitalization - Initial or Prolonged	Agitation Convulsion	Health Professional Company Representative	Wellbutrin Caffeine	PS C
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Date:01/20/98ISR Number: 3085469-XReport Type:Periodic Company Report #A0049701  
 Age:16 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin	PS		ORAL
Other		Euphoric Mood	Professional				
800MG/ORAL		Hallucination Overdose					

Date:01/20/98ISR Number: 3085643-2Report Type:Periodic Company Report #A0055690  
 Age:9 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin	PS		ORAL
Hospitalization - 150 MG/ORAL		Abnormal Behaviour	Professional	Clonidine	C		
Initial or Prolonged		Aggression	Other	Methylphenidate	C		
		Agitation					
		Disorientation					
		Impulse-Control Disorder					
		Memory Impairment					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/20/98ISR Number: 3085658-4Report Type:Periodic Company Report #A0056325  
 Age:23 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL Other	Myocardial Infarction	Health Professional Company Representative	Wellbutrin	PS		ORAL

Date:01/20/98ISR Number: 3085733-4Report Type:Periodic Company Report #A0057417  
 Age:26 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL	Confusional State Disorientation Hallucination	Health Professional Company Representative	Wellbutrin	PS		ORAL

Date:01/20/98ISR Number: 3085746-2Report Type:Periodic Company Report #A0057671  
 Age:35 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100 MG/ORAL Initial or Prolonged Other	Convulsion Overdose	Health Professional	Wellbutrin Fluoxetine Hydrochloroide (Formulation Unknwon) Ethanol (Formulation Unknown)	PS SS SS		ORAL

Date:01/20/98ISR Number: 3085884-4Report Type:Periodic Company Report #A0050029  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Myocardial Infarction	Health	Wellbutrin	PS		ORAL
Initial or Prolonged			Professional Company Representative	Diltiazem Hydrochloride Glibenclamide Ranitidine Hydrochloride	C C C		

Date:01/20/98ISR Number: 3085885-6Report Type:Periodic Company Report #A0050061  
 Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100MG/TWICE PER DAY/ORAL		Bruxism Pain In Jaw	Health Professional	Wellbutrin	PS		ORAL

Date:01/20/98ISR Number: 3085900-XReport Type:Periodic Company Report #A0050131  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG/PER DAY/ORAL	6 MON	Arrhythmia Chest Pain Convulsion	Consumer	Wellbutrin Lithium Salt Lamotrigine Thyroxine Sodium	PS C C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Estrogen C  
 Dalmane C  
 Thiothixene C

Date:01/27/98ISR Number: 3019377-7Report Type:Expedited (15-DaCompany Report #8-98009-010H  
 Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cardiac Valve Disease	Health	Redux	PS		ORAL
15 MG TWICE		Cardiomegaly	Professional				
DAILY ORAL		Dyspnoea		Wellbutrin	SS		
150 MG TWICE							
DAILY				Provera	C		
10 MG				Synthroid	C		
0.125 MG				Wellbutrin	C		
300 MG							

Date:01/27/98ISR Number: 3021414-0Report Type:Direct Company Report #  
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis		Zyban	PS	Glaxo	
Other		Disability		Depo-Provera	C		
1 TABLET BID	2 DAY	Eyelid Oedema		Benadryl	C		
		Immobile					
		Joint Swelling					
		Oedema Mouth					
		Pain					
		Rash Erythematous					

Date:01/27/98ISR Number: 3030775-8Report Type:Direct Company Report #  
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia		Zyban	PS		
1 TABLET BID	14 DAY						
		Dermatitis		Depo-Provera	C		
		Eyelid Oedema		Benadryl	C		
		Joint Swelling					
		Oedema					
		Oedema Mouth					
		Pain					

Date:01/27/98ISR Number: 3109823-2Report Type:Direct Company Report #  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Libido Increased		Wellbutrin	PS		ORAL
150 G TID							

Date:01/28/98ISR Number: 3020961-5Report Type:Expedited (15-DaCompany Report #8-98016-002S  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Ventricular Hypertrophy	Health	Pondimin	PS		ORAL
ORAL	6 MON						
			Professional	Phentermine	SS		ORAL
ORAL	6 MON			Wellbutrin	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/98ISR Number: 3021045-2Report Type:Expedited (15-DaCompany Report #8-98016-002S  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	6	MON	Ventricular Hypertrophy	Health	PS		ORAL
ORAL			Professional	Wellbutrin Phentermine	SS SS		
6	MON						

Date:01/28/98ISR Number: 3021583-2Report Type:Direct Company Report #  
 Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG PO BID		Blister	Wellbutrin	PS		ORAL
			Dermatitis Bullous Mucosa Vesicle Mucosal Haemorrhage Petechiae Rosacea	Tetracycline	C		

Date:01/28/98ISR Number: 3021588-1Report Type:Direct Company Report #  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG PO BID		Drug Hypersensitivity	Zyban	PS	Glaxo/Wellcome	
Hospitalization - Initial or Prolonged			Pharyngeal Oedema Respiratory Distress Urticaria				

Date:01/28/98ISR Number: 3021674-6Report Type:Direct Company Report #  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Chest Discomfort Zyban PS Glaxo  
 150 MG BID 2 WK  
 Initial or Prolonged Erythema Multiforme  
 Face Oedema  
 Urticaria  
 White Blood Cell Count  
 Increased

Date:01/29/98ISR Number: 3112043-9Report Type:Periodic Company Report #8-97174-018L  
 Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Health	Redux	PS		ORAL
15-30 MG		Fatigue	Professional				
DAILY ORAL							
				Wellbutrin	SS		ORAL
25 MG DAILY							
ORAL							

Date:01/30/98ISR Number: 3021134-2Report Type:Expedited (15-DaCompany Report #A0059505  
 Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Euphoric Mood	Consumer	Zyban	PS		ORAL
150 MG PER		Feeling Abnormal					
DAY ORAL	6 DAY	Neoplasm Malignant		..	C		
		Psychomotor Hyperactivity					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/30/98ISR Number: 3021151-2Report Type:Expedited (15-DaCompany Report #A0058742  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Haemorrhagic Disorder	Health	Wellbutrin	PS		ORAL
Hospitalization -		Mouth Haemorrhage	Professional				
Initial or Prolonged		Petechiae					
Other		Platelet Count Decreased		Tetracycline	C		
		Thrombocytopenia		Phenobarbitone	C		

Date:02/02/98ISR Number: 3021856-3Report Type:Direct Company Report #  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Decreased Activity		Zyban	PS		ORAL
		Flashback		Lovazepam	C		
		Flushing		Pindolol	C		
		Insomnia					
		Panic Attack					
		Post-Traumatic Stress Disorder					
		Psychiatric Symptom					

Date:02/02/98ISR Number: 3022150-7Report Type:Expedited (15-DaCompany Report #A0058605  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Anorectal Disorder	Health	Zyban	PS		ORAL
Initial or Prolonged		Constipation	Professional				
Other		Drug Interaction		Omeprazole	SS		

Date:02/02/98ISR Number: 3022152-0Report Type:Expedited (15-DaCompany Report #A0057797  
Age:66 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG PER Initial or Prolonged DAY	Blood Pressure Increased Dehydration Hypersomnia Malaise Nausea Visual Disturbance	Health Professional	Zyban Conjugated Estrogens Ibuprofen	PS C C		ORAL

Date:02/02/98ISR Number: 3088615-7Report Type:Periodic Company Report #A0057121  
Age:44 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG TWICE PER DAY ORAL	Depression Drug Ineffective Feeling Abnormal	Consumer	Wellbutrin Tablet- Controlled Release Hyzaar Berocca	PS C C		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/02/98ISR Number: 3091420-9Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #A0057691

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG TWICE PER DAY ORAL 10 MG PER DAY ORAL	Hepatitis A	Health Professional Company Representative	Wellbutrin Tablet- Controlled Release  Simvastatin Tablet	PS  SS		ORAL  ORAL

Date:02/02/98ISR Number: 3091432-5Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0057795

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150 MG PER DAY ORAL	Arthralgia	Consumer	Wellbutrin  Imipramine Conjugated Estrogens	PS  C C		ORAL
Duration 30 DAY						

Date:02/02/98ISR Number: 3107304-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0058281

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG/TWICE Initial or Prolonged PER DAY/ORAL	Anaphylactic Reaction Blood Pressure Decreased Cognitive Disorder Dermatitis Dyspnoea Hypoaesthesia Oral Oedema Peripheral Urticaria	Health Professional	Wellbutrin  Diphenhydramine Hcl Sertraline Hydrochloride	PS  SS C		ORAL

Date:02/02/98ISR Number: 3107306-7Report Type:Periodic Company Report #A0054447  
Age:38 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150MG/FOUR	Fall	Health	Wellbutrin	PS		
TIMES PER DAY 75 MG / SEE	Grand Mal Convulsion	Professional				
TEXT/ ORAL	Periorbital Haematoma	Company Representative	Wellbutrin	SS		ORAL
			Desipramine	C		

Date:02/02/98ISR Number: 3114037-6Report Type:Periodic Company Report #A0055443  
Age:49 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ TWICE	Blood Creatine	Health	Wellbutrin	PS		ORAL
Initial or Prolonged PER DAY/ ORAL	Phosphokinase Increased	Professional				
Other	Blood Lactate Dehydrogenase Increased Hepatic Enzyme Increased Muscle Disorder Rhabdomyolysis	Company Representative	Glipizide Metformin Multivitamin Insulin	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/02/98ISR Number: 3114038-8Report Type:Periodic Company Report #A0055506  
 Age:82 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	75 MG/ UNK/	Coordination Abnormal	Health	Wellbutrin	PS		ORAL
ORAL			Professional				
			Company Representative	Ludiomil Lithium Salt Thyroxine	C C C		

Date:02/02/98ISR Number: 3114039-XReport Type:Periodic Company Report #A0055507  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG/ TWICE	Insomnia	Health	Wellbutrin	PS		ORAL
PER DAY/ ORAL		Road Traffic Accident	Professional				
		Sedation	Company Representative	Aspirin Nicotinic Acid Atorvastatin Calcium	C C C		

Date:02/02/98ISR Number: 3114040-6Report Type:Periodic Company Report #A0055603  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	50 MG/ IN THE	Face Oedema	Health	Wellbutrin	PS		ORAL
MORNING/ ORAL		Fatigue	Professional				
		Malaise Mucosal Inflammation Pruritus Pyrexia Rash Macular Stevens-Johnson Syndrome	Company Representative	Amitriptyline Hcl Fluticasone Propionate Claritin-D Vitamin E Pravastatin Sodium Zolpidem Tartrate	C C C C C C C		

Date:02/02/98ISR Number: 3114041-8Report Type:Periodic Company Report #A0056413  
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain	Consumer	Wellbutrin	PS		ORAL
150 MG/	SEE	Chills					
TEXT/	ORAL	Cognitive Disorder		Venlafaxine			
		Depersonalisation		Hydrochloride	C		
		Insomnia		Caffeine	C		
		Neck Pain		Antioxidants	C		
		Overdose					
		Pain					
		Tremor					

Date:02/02/98ISR Number: 3114042-XReport Type:Periodic Company Report #A0056647  
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Face Oedema	Health	Wellbutrin	PS		ORAL
150 MG / UNK/		Oedema Peripheral	Professional				
Initial or Prolonged		Pruritus	Company	Fiorinal	C		
ORAL		Urticaria	Representative				
Other							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/03/98ISR Number: 3022709-7Report Type:Direct  
Age:36 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	100 MG PO BID	Convulsion		Wellbutrin	PS		ORAL

Date:02/03/98ISR Number: 3028904-5Report Type:Direct  
Age:76 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG PO QD	Tachycardia	Health	Wellbutrin	PS		
	500MG PO Q12		Professional	Robaxin	SS		

H

Prinivil	C
Prilosec	C
Cipro	C
Imdur	C
K-Dur	C
Ativan	C
Lorcet 10/650	C

Date:02/05/98ISR Number: 3024334-0Report Type:Direct  
Age:52 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - TAKE ONE		Dermatitis		Zyban	PS		ORAL
Initial or Prolonged TABLET BY		Drug Hypersensitivity					
Other MOUTH EVERY MORNING FOR 3 DAYS THEN 1		Dyspnoea					
		Pyrexia					

Date:02/06/98ISR Number: 3026429-4Report Type:Expedited (15-DaCompany Report #A0057792  
Age:58 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150 MG/ PER	Confusional State	Consumer	Zyban	PS		ORAL
Initial or Prolonged DAY / ORAL	Dehydration					
	Depressed Level Of Consciousness		Oxpentifylline	C		
	Hyponatraemia		Nabumetone	C		
	Polyuria		Claritin-D	C		
	Vomiting		Lansoprazole	C		
	Weight Decreased		Conjugated Estrogens	C		
			Dyazide	C		
			Simvastatin	C		
			Alprazolam	C		
			Amitriptyline Hcl	C		
			Fluticasone			
			Propionate	C		
			Aspirin	C		

Date:02/09/98ISR Number: 3027036-XReport Type:Expedited (15-DaCompany Report #A0060104  
Age:68 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150 MG TWICE	Abdominal Pain	Consumer	Zyban	PS		ORAL
Initial or Prolonged PER DAY ORAL	Cholelithiasis					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/10/98ISR Number: 3027179-0Report Type:Expedited (15-DaCompany Report #A0059063  
 Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Malaise	Health	Wellbutrin	PS		ORAL
Hospitalization - DAY / ORAL		Pancreatic Enzymes	Professional				
Initial or Prolonged Other		Increased Pancreatitis Pulmonary Oedema		Sertraline Hydrochloride	C		

Date:02/11/98ISR Number: 3029072-6Report Type:Expedited (15-DaCompany Report #1998-02-0042  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death UNKNOWN ORAL		Completed Suicide	Health	Albuterol	PS		
Other AER INH		Drug Toxicity	Professional				
				Bupropion Hydrochloride	SS		
				Hydrocodone	SS		
				Ethanol	SS		
				Phenylpropanolamine	SS		
				Chlorpheniramine	SS		

Date:02/11/98ISR Number: 3029514-6Report Type:Direct Company Report #  
 Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 SR BID		Rectal Haemorrhage		Wellbutrin	PS		
				Flovent	C		
				Serevent	C		
				Conbivent	C		
				Theophylline	C		

Date:02/13/98ISR Number: 3029818-7Report Type:Direct  
Age:16 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG BID PO Initial or Prolonged MONTHS		Duodenal Ulcer Haematemesis MON		Wellutrin  Pancrease Vit C Vit E	PS  C C C	Glaxo	ORAL

Date:02/17/98ISR Number: 3030644-3Report Type:Expedited (15-DaCompany Report #A0060394  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY  (TABLET)		Anxiety Chest Pain Disturbance In Attention  Dyspnoea Palpitations Poor Peripheral Circulation Tongue Oedema Tremor	Consumer	Zyban	PS		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/98ISR Number: 3030646-7Report Type:Expedited (15-DaCompany Report #A0059505  
Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Euphoric Mood	Consumer	Zyban	PS		ORAL
150 MG PER		Psychomotor Hyperactivity					
DAY ORAL							
(TABLET)							

Date:02/19/98ISR Number: 3032437-XReport Type:Expedited (15-DaCompany Report #A0059062  
Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Zyban	PS		ORAL
ORAL		Nausea		Cardovascular Medication	C		

Date:02/19/98ISR Number: 3032439-3Report Type:Expedited (15-DaCompany Report #A0060149  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Blood Pressure Increased	Health	Wellbutrin	PS		ORAL
150 MG /		Dermatitis	Professional				
TWICE PER		Heart Rate Increased	Company				
DAY/		Oedema Peripheral Tongue Oedema	Representative				

Date:02/23/98ISR Number: 3036055-9Report Type:Expedited (15-DaCompany Report #A0060690  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 150MG/TWICE Initial or Prolonged PER DAY/ ORAL	Dermatitis Diarrhoea Erythema Multiforme Nausea Pruritus Renal Impairment Vomiting	Health	Zyban	PS	ORAL
		Professional			

Date:02/26/98ISR Number: 3036550-2Report Type:Expedited (15-DaCompany Report #A0060825  
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY / ORAL	8 DAY	Alcohol Interaction	Health	Wellbutrin	PS		ORAL
		Asthenia	Professional				
		Clonic Convulsion	Company				
		Confusional State	Representative	Ethanol	C		
		Difficulty In Walking					
		Encephalopathy					
		Hallucination					
		Syncope					
		Urinary Incontinence					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/98ISR Number: 3043863-7Report Type:Expedited (15-DaCompany Report #A0060881  
Age:47 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150MG / TWICE	Haematuria	Health	Zyban	PS		ORAL
Initial or Prolonged PER DAY /	Subarachnoid Haemorrhage	Professional				
ORAL	Vascular Injury					

Date:02/26/98ISR Number: 3043866-2Report Type:Expedited (15-DaCompany Report #A0060760  
Age:48 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150 MG / PER	Convulsion	Consumer	Zyban	PS		ORAL
Initial or Prolonged DAY / ORAL	Insomnia					
	Spinal Compression		Mirtazapine	C		
	Fracture		Alprazolam	C		

Date:02/26/98ISR Number: 3116007-0Report Type:Periodic Company Report #1997-000731(0)  
Age:47 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Erectile Dysfunction	Consumer	Viracept (Nelfinavir			
750 MG TID		Health	Mesyate)	PS		
		Professional	Wellbutrin	SS		
			Zerit	C		
			3tc	C		

Date:02/27/98ISR Number: 3042833-2Report Type:Direct Company Report #  
Age:50 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

150 MG BID Rash Pruritic Wellbutrin PS  
 Urticaria

Date:02/27/98ISR Number: 3043945-XReport Type:Direct Company Report #  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis Exfoliative		Wellbutrin	PS		ORAL
150 MG BID		Rash Macular					

Date:03/02/98ISR Number: 3038990-4Report Type:Expedited (15-DaCompany Report #A0011982  
 Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Anaphylactic Reaction	Study	Wellbutrin	PS		ORAL
150 MG /		Cyanosis	Health				
TWICE PER		Diarrhoea	Professional				
DAY/ ORAL		Dyspnoea		Verapamil	C		
		Nausea		Chlorpheniramine	C		
		Oedema Peripheral		Diphenhydramine	C		
		Petechiae		Hydrocortisone	C		
		Pruritus		Amoxicillin + K			
		Urticaria		Clavulanat	C		
		Vomiting					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/98ISR Number: 3038994-1Report Type:Expedited (15-DaCompany Report #A0041335  
Age:66 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150 MG /TWICE PER DAY/ ORAL	Anaphylactic Reaction Cyanosis Diarrhoea Dyspnoea Nausea Oedema Mouth Oedema Peripheral Periorbital Oedema Petechiae Pruritus Urticaria Vomiting	Study Health Professional	Bupropion Hydrochloride Verapamil Chloropheniramine Augmentin Diphenhydramine Hydrocortisone	PS  C C C C C		ORAL

Date:03/02/98ISR Number: 3129598-0Report Type:Periodic Company Report #A0056494  
Age:46 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL	Convulsion	Health Professional Company Representative	Zyban Clarithromycin Humibid Dm Guaiphenesin	PS  C C C		ORAL

Date:03/02/98ISR Number: 3129600-6Report Type:Periodic Company Report #A0056729  
Age:58 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability ORAL	Depersonalisation Depression Suicidal	Consumer	Zyban	PS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE		Dysgeusia	Consumer	Zyban	PS		ORAL
Initial or Prolonged PER DAY/ORAL		Dyspnoea					
		Tremor		Aspirin	C		
				Quinapril			
				Hydrochloride	C		
				Diltiazem			
				Hydrochloride	C		
				Ranitidine			
				Hydrochloride	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150 MG/ORAL		Decreased Appetite	Consumer	Zyban	PS		ORAL
		Nausea		Conjugated Estrogens	C		
				Thyroxine Sodium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/98ISR Number: 3129607-9Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0056996

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL		Grand Mal Convulsion	Consumer	Zyban	PS		ORAL
				Prempro	C		

Date:03/02/98ISR Number: 3132927-5Report Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #9719922

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 30.00 MG TOTAL:DAILY		Anxiety Cardiovascular Disorder	Health Professional	Zoloft Redux	PS SS		ORAL ORAL
				Ambien	SS		ORAL
				Wellbutrin	SS		ORAL

Date:03/02/98ISR Number: 3133750-8Report Type:Periodic  
Age:65 YR Gender:Female I/FU:I

Company Report #A0058109

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG/TWICE PER DAY/ORAL		Dermatitis Exfoliative Pruritus Urticaria	Health Professional	Zyban	PS		ORAL
				Triamterene	C		
				Simvastatin	C		
				Potassium Chloride	C		
				Aspirin	C		

Date:03/02/98ISR Number: 3133751-XReport Type:Periodic  
Age:37 YR Gender:Male I/FU:I

Company Report #A0058045

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Angioneurotic Oedema	Health	Zyban	PS		ORAL
150 MG/TWICE							
		Dermatitis	Professional				
PER DAY	ORAL						
		Hypersensitivity					
		Pruritus					
		Urticaria					

Date:03/02/98ISR Number: 3134710-3Report Type:Periodic Company Report #A0057956  
Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Hospitalization -		Arrhythmia	Health	Zyban	PS		ORAL
150 MG/ORAL							
Initial or Prolonged		Hypertension	Professional				
			Company Representative				

Date:03/02/98ISR Number: 3138605-0Report Type:Periodic Company Report #A0055397  
Age:50 YR Gender:Male I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Arthritis	Health	Zyban	PS		ORAL
150 MG /							
TWICE PER DAY		Dermatitis	Professional				
		Eyelid Oedema					
/ ORAL		Hypersensitivity		Vitamin E	C		
		Pruritus		Ascorbic Acid	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Cyanocobalamin C  
 Aspirin C  
 Atenolol C  
 Atorvastatin Calcium C

Date:03/02/98ISR Number: 3138607-4Report Type:Periodic  
 Age:49 YR Gender:Male I/FU:F

Company Report #A0055439

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG/ TWICE	Myocardial Infarction	Health	Zyban	PS		ORAL
Hospitalization - PER DAY / Initial or Prolonged			Professional				
Other				Human Int/Long Insulin	C		

Date:03/02/98ISR Number: 3138609-8Report Type:Periodic  
 Age:45 YR Gender:Female I/FU:F

Company Report #A0055780

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged		Anxiety	Health	Zyban	PS		ORAL
TWICE PER Other		Hypertension	Professional				
DAY/ ORAL				Atenolol Nifedipine	C C		

Date:03/02/98ISR Number: 3138611-6Report Type:Periodic  
 Age:40 YR Gender:Male I/FU:F

Company Report #A0055781

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG /		Dermatitis	Health	Zyban	PS		ORAL

Professional  
Dry Mouth  
Dysphagia  
Mouth Ulceration  
TWICE PER DAY  
/ ORAL

Date:03/02/98ISR Number: 3138612-8Report Type:Periodic Company Report #A0056353  
Age:40 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY / ORAL	Convulsion	Health Professional Company Representative	Zyban	PS		ORAL

Date:03/02/98ISR Number: 3138614-1Report Type:Periodic Company Report #A0056449  
Age:59 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG / PER DAY / ORAL	Abdominal Discomfort Vomiting	Consumer	Zyban	PS		ORAL
			Salbutamol Sulphate	C		
			Ipratropium Bromide	C		
			Zafirlukst	C		
			Conjugated Estrogens	C		
			Guaiphenesin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/98ISR Number: 3138616-5Report Type:Periodic  
Age:37 YR Gender:Female I/FU:F

Company Report #A0056510

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY/ ORAL	Hypersensitivity Pruritus Rash Erythematous Swelling Urticaria	Consumer	Zyban   Decongestant	PS   C		ORAL

Date:03/02/98ISR Number: 3138618-9Report Type:Periodic  
Age:27 YR Gender:Male I/FU:F

Company Report #A0056581

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150 MG / Other TWICE PER DAY / ORAL	Dyspnoea Oropharyngeal Swelling Rash Erythematous Urticaria	Health Professional Company Representative	Zyban	PS		ORAL

Date:03/02/98ISR Number: 3139909-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0058975

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG/ ORAL	Mucosal Inflammation Pruritus Rash Erythematous Urticaria	Consumer	Zyban Tablet	PS		ORAL

Date:03/02/98ISR Number: 3139913-XReport Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0058600

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG/ TWICE	Flushing	Health	Zyban Tablet	PS		ORAL
PER DAY /			Professional				
ORAL			Company				
			Representative				

Date:03/02/98ISR Number: 3139916-5Report Type:Periodic Company Report #A0059506  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG /	Anorexia	Consumer	Zyban Tablet	PS		ORAL
TWICE PER		Paranoia					
DAY/ ORAL		Suicide Attempt					

Date:03/02/98ISR Number: 3139917-7Report Type:Periodic Company Report #A0059708  
Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG/ TWICE	Hypoaesthesia	Consumer	Zyban Tablet	PS		ORAL
Initial or Prolonged	PER DAY /	Pain In Extremity					
ORAL				Metoprolol Tartrate	C		
				Simvastatin	C		
				Atorvastatin Calcium	C		
				Losartan Potassium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/98ISR Number: 3139919-0Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #A0059766

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY  / ORAL	Dyspnoea  Eyelid Oedema  Insomnia  Oedema Peripheral Pruritus Urticaria	Consumer	Zyban Tablet	PS		ORAL
			Glipizide	C		
			Theophylline	C		
			Dyazide	C		

Date:03/02/98ISR Number: 3139921-9Report Type:Periodic  
Age:60 YR Gender:Male I/FU:I

Company Report #A0059794

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged SINGLE DOSE /  ORAL	Bronchospasm  Dyspnoea	Consumer	Zyban Tablet	PS		ORAL
			Salbutamol Sulphate	C		
			Combivent	C		
			Theophylline	C		

Date:03/02/98ISR Number: 3146677-2Report Type:Periodic  
Age:85 YR Gender:Male I/FU:I

Company Report #9715884

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other TABLETS,  50.00 MG  TOTAL:DAILY:0  RAL	Aortic Valve Stenosis  Cardiac Failure  Congestive  Drug Interaction	Health  Professional	Zoloft	PS		ORAL

100.00 MG  
TOTAL:DAILY:0  
RAL

Wellbutrin

SS

ORAL

Date:03/02/98ISR Number: 3147151-XReport Type:Periodic Company Report #9706189  
Age:17 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Health	Zoloft	PS		ORAL
TABLETS,		Asthenia	Professional				
200.00 MG		Confusional State					
TOTAL:DAILY:0		Tremor					
RAL		Visual Disturbance		Wellbutrin	SS		ORAL
ORAL				Desyrel	C		

Date:03/02/98ISR Number: 3147733-5Report Type:Periodic Company Report #A0059990  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Crying	Health	Zyban	PS		ORAL
150 MG ORAL	7 DAY	Depression	Professional	Nicotine	C		
		Psychotic Disorder	Company				
		Tremor	Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/98ISR Number: 3150053-6Report Type:Periodic  
Age:33 YR Gender:Male I/FU:I

Company Report #9728129

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Dizziness	Consumer	Zoloft	PS		ORAL
ORAL		Drug Ineffective		Prozac	SS		
ORAL		Erectile Dysfunction		Wellbutrin	SS		ORAL
ORAL				Parnate	SS		ORAL
				Unknown			
				Antidepressants	C		

Date:03/02/98ISR Number: 3266458-9Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #A0058677

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Grand Mal Convulsion	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG TWICE		Insomnia					
Initial or Prolonged		Speech Disorder					
PER DAY ORAL		Tremor					

Date:03/02/98ISR Number: 3266463-2Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0058688

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/02/98ISR Number: 3266466-8Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #A0058629

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG TWICE		Dermatitis	Consumer	Zyban Tablet- Zyban	PS		ORAL
Initial or Prolonged PER DAY ORAL		Hypersensitivity					

Date:03/02/98ISR Number: 3266468-1Report Type:Periodic Company Report #A0058434  
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150 MG PER Disability DAY ORAL		Amnesia	Health	Zyban Tablet- Zyban	PS		ORAL
		Convulsion	Professional				
		Loss Of Consciousness Road Traffic Accident					

Date:03/02/98ISR Number: 3266471-1Report Type:Periodic Company Report #A0058441  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Constipation	Health	Zyban Tablet- Zyban	PS		ORAL
.2 MG TWICE PER DAY		Drug Interaction Insomnia Orthostatic Hypotension	Professional	Clonidine (Formulation Unknown)	SS		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/98ISR Number: 3266473-5Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #A0058329

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL	Arthralgia	Consumer	Zyban Tablet-Zyban	PS		ORAL
	Diarrhoea					
	Difficulty In Walking		Dristan	C		
	Hypersensitivity		Famotidine	C		
	Joint Swelling					
	Musculoskeletal Stiffness					
	Oedema					
	Pruritus					
	Rash Erythematous					
	Urticaria					

Date:03/02/98ISR Number: 3266477-2Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #A0058318

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL	Dizziness	Consumer	Zyban Tablet- Zyban	PS		ORAL
	Face Oedema					
	Hyperhidrosis		Diltiazem			
	Hypersensitivity		Hydrochloride	C		
	Syncope		Paroxetine			
	Tinnitus		Hydrochloride	C		
	Vomiting		Lisinopril	C		

Date:03/02/98ISR Number: 3266484-XReport Type:Periodic  
Age:31 YR Gender:Male I/FU:I

Company Report #A0058298

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG ORAL 2 WK Initial or Prolonged	Anaphylactic Reaction	Health	Zyban Tablet- Zyban	PS		ORAL
	Bronchospasm	Professional				
	Dizziness					
	Eyelid Oedema					

Face Oedema  
 Hypersensitivity  
 Laryngeal Oedema  
 Pain  
 Periorbital Oedema  
 Swelling  
 Urticaria

Date:03/04/98ISR Number: 3040452-5Report Type:Expedited (15-DaCompany Report #8-97307-005S  
 Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Health	Pondimin	PS		ORAL
20-40MG DAILY		Aortic Valve Incompetence	Professional				
ORAL		Cardiac Valve Disease		Phentermine	SS		ORAL
ORAL		Disturbance In Attention		Phentermine	SS		ORAL
ORAL		Fibromyalgia		Prozac	SS		
		Herpes Simplex		Wellbutrin	SS		
		Mitral Valve Incompetence		Phentermine	C		
		Mitral Valve Prolapse		Phentermine	C		
		Oedema		Prozac	C		
		Skin Disorder		Wellbutrin	C		
				Xanax	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/04/98ISR Number: 3147732-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0059950

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG ORAL		Convulsion	Consumer	Zyban	PS		ORAL
Initial or Prolonged		Dry Mouth Nervousness Pancreatitis					

Date:03/05/98ISR Number: 3050356-XReport Type:Direct  
 Age:2 DY Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - PO		Complications Of Maternal		Wellbutrin	PS		ORAL
Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage		Exposure To Therapeutic Drugs Irritability Screaming Tachycardia Tachypnoea					

Date:03/05/98ISR Number: 3050361-3Report Type:Direct  
 Age:37 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Angioneurotic Oedema Arthropathy Urticaria		Wellbutrin	PS		

Date:03/06/98ISR Number: 3145638-7Report Type:Periodic  
 Age:33 YR Gender:Male I/FU:I

Company Report #8-97344-007H

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG TWICE		Drug Ineffective	Consumer	Effexor	PS		ORAL

Drug Withdrawal Syndrome

DAILY TO 18+

Paraesthesia

MG DAILY ORAL

Wellbutrin

SS

150 MG TWICE

DAILY

Date:03/06/98ISR Number: 3145923-9Report Type:Periodic

Company Report #8-97154-001B

Age:35 YR Gender:Male

I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 350 MG DAILY Initial or Prolonged ORAL		Coordination Abnormal	Health Professional	Effexor	PS		ORAL
20 MG				Ambien	SS		
50-100 MG				Wellbutrin	SS		
DAILY				Lithium	C		
				Risperidone	C		
				Thyroid	C		
				Xanax	C		
				Ambien	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/09/98ISR Number: 3049047-0Report Type:Expedited (15-DaCompany Report #A0061492  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG/ TWICE	Convulsion	Consumer	Zyban	PS		ORAL
PER DAY/ ORAL	4	WK	Disturbance In Attention				
		Fall		Paracetamol	C		
		Head Injury		Darvocet-N	C		
		Insomnia		Nicotine	C		

Date:03/09/98ISR Number: 3049050-0Report Type:Expedited (15-DaCompany Report #A0061472  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Medication Error	Health	Wellbutrin	PS		ORAL
ORAL		Hospitalization - Initial or Prolonged	Status Epilepticus	Professional Company Representative			

Date:03/09/98ISR Number: 3049566-7Report Type:Expedited (15-DaCompany Report #A0061053  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Death	Health	Wellbutrin	PS		ORAL
ORAL			Professional Company Representative	Beta-Blocker	C		

Date:03/10/98ISR Number: 3049124-4Report Type:Direct Company Report #  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other 150MG PO BID Face Oedema Zyban PS ORAL  
 Oedema Peripheral  
 Pruritus  
 Urticaria

Date:03/10/98ISR Number: 3049152-9Report Type:Direct Company Report #  
 Age:88 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 75MG BID		Blood Creatinine		Bupropion	PS		
Initial or Prolonged 7/30/96 24H		Increased					
INTRAVENOUS	2MG IV	Blood Urea Increased Q 4 HR		Morphine Sulfate	SS		
PRN 2 DOSES		Confusional State					
		Hip Fracture		Feso4	C		
		Lethargy		Levothyroxine	C		
		Mental Impairment		Metoprolrol	C		
				Hydrocortisone	C		
				Ofloxacin	C		

Date:03/10/98ISR Number: 3056531-2Report Type:Direct Company Report #  
 Age: Gender:Male I/FU:I

Outcome PT  
 Other Extrapyramidal Disorder  
 Paraesthesia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Restlessness

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
75MG PO TID			Bupropion	PS		ORAL
			Cyclobenzaprine	C		
			Thothixene	C		
			Lisinopril	C		
			Trazodone	C		
			Ritalin	C		
			Etodolac	C		
			Sinemet	C		

Date:03/11/98ISR Number: 3056366-0Report Type:Direct  
 Age:62 YR Gender:Female I/FU:I Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dermatitis Pyrexia Stevens-Johnson Syndrome	Health Professional	Zyban	PS		

Date:03/11/98ISR Number: 3128844-7Report Type:Expedited (15-DaCompany Report #A0069088  
 Age:65 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG/ TWICE PER DAY/ ORAL		Chest Pain Choking Convulsion Thirst Tremor	Health Professional	Zyban  Simvastatin	PS  C		ORAL

Date:03/12/98ISR Number: 3053266-7Report Type:Expedited (15-DaCompany Report #A0060690  
 Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 150MG/ TWICE	Anaemia	Health	Zyban	PS	ORAL
Initial or Prolonged PER DAY	Asthenia	Professional			
Disability	Blood Creatinine Increased				
Other	Blood Urea Increased				
	Dermatitis				
	Diarrhoea				
	Erythema Multiforme				
	Gastritis				
	Haematochezia				
	Hyperglycaemia				
	Nausea				
	Oedema				
	Pain				
	Pneumonia				
	Proteinuria				
	Pruritus				
	Pyrexia				
	Renal Impairment				
	Urticaria				
	Vomiting				



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/12/98ISR Number: 3127339-4Report Type:Periodic  
Age:11 YR Gender:Male I/FU:I

Company Report #JAUSA-28403

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Leukopenia	Health Professional	Risperdal	PS		
				Clonidine (Clonidine)	SS		
				Wellbutrin (Amfebutamone)	SS		

Date:03/12/98ISR Number: 3148265-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #JAUSA-29679

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tremor		Risperdal (Risperidone) Janssen Tablet	PS	Janssen	ORAL
ORAL				Wellbutrin (Amfebutamone)	SS		
				Depakote	SS		

Date:03/12/98ISR Number: 3148304-7Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #JAUSA-29749

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation Chest Pain Tachycardia	Consumer	Risperdal (Risperidone) Janssen Tablets	PS	Janssen	ORAL
3MG 2 DAILY							
ORAL, AN							
OCASSINAL 1MG							
IS ADDED							
DAILY WHEN				Wellbutrin			

75MG 2 DAILY (Amfebutamone) SS ORAL  
 ORAL, 75MG 3  
 DAILY 6 DAY Claritin-D C

Date:03/16/98ISR Number: 3056168-5Report Type:Expedited (15-DaCompany Report #A0061756  
 Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Influenza Like Illness	Health Professional	Wellbutrin	PS		ORAL
150MG/TWICE							
PER DAY/ORAL/							
TAB				Pseudoephedrine Hcl	SS		ORAL
ORAL				Cough Medication	C		

Date:03/16/98ISR Number: 3056172-7Report Type:Expedited (15-DaCompany Report #A0061791  
 Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Blood Pressure Increased Dizziness Dyspnoea Heart Rate Increased Loss Of Consciousness Monoparesis

Freedom Of Information (FOI) Report

Pain In Extremity

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG/ TWICE PER DAY/ ORAL/ TAB		Consumer	Zyban	PS		ORAL

Date:03/16/98ISR Number: 3056176-4Report Type:Expedited (15-DaCompany Report #A0059505  
Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG/PER DAY/ORAL/ TAB		Anxiety Brain Neoplasm Drug Ineffective Euphoric Mood Psychomotor Hyperactivity Small Cell Lung Cancer Stage Unspecified	Health Professional	Zyban	PS		ORAL

Date:03/16/98ISR Number: 3057431-4Report Type:Expedited (15-DaCompany Report #A0061467  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Complications Of Maternal Exposure To Therapeutic Drugs Drug Withdrawal Convulsions	Health Professional Company Representative	Wellbutrin	PS		ORAL

Date:03/17/98ISR Number: 3055919-3Report Type:Expedited (15-DaCompany Report #COU980009  
Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abdominal Pain Upper	Health	Coumadin	PS		ORAL
UNKNOWN-2.5							
Hospitalization -		Arthralgia	Professional				
MG; ORAL							
Initial or Prolonged		Cardiac Failure		Wellbutrin	SS		ORAL
300 MG; ORAL							
		Congestive		Albuterol	C		
		Constipation		Atrovent	C		
		Dizziness		Axid	C		
		Drug Interaction		Beclovent	C		
		International Normalised		Colchicine	C		
		Ratio Increased		Diamox	C		
		Prothrombin Time		Digoxin	C		
		Prolonged		Isordil	C		
		Rhinorrhoea		Lisinopril	C		
				Procan	C		

Date:03/18/98ISR Number: 3056376-3Report Type:Direct Company Report #  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Burning Sensation		Wellbutrin Sr	PS	Glaxo-Wellcome	
I BID							
		Hypoaesthesia					
		Paraesthesia					
		Skin Exfoliation					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/19/98ISR Number: 3057789-6Report Type:Direct  
 Age:25 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
IPO BID		Convulsion		Zyban	PS		ORAL

Date:03/20/98ISR Number: 3059526-8Report Type:Expedited (15-DaCompany Report #A0058622  
 Age:68 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / PER Initial or Prolonged DAY/ ORAL Other		No Adverse Drug Effect	Consumer	Zyban	PS		ORAL
				Betaxolol Hydrochloride	C		
				Lorazepam	C		
				Theophylline	C		

Date:03/20/98ISR Number: 3059528-1Report Type:Expedited (15-DaCompany Report #A0062268  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / UNK Initial or Prolonged / ORAL		Unevaluable Event	Consumer	Zyban	PS		ORAL

Date:03/20/98ISR Number: 3059530-XReport Type:Expedited (15-DaCompany Report #A0062113  
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY		Abdominal Pain Haemorrhage	Health Professional	Wellbutrin	PS		ORAL

/ ORAL	Hypotension					
	Lethargy			Fluoxetine		
	Retching			Hydrochloride	C	
	Splenic Rupture			Oxaprozin	C	
	Vomiting			Sinemet	C	

Date:03/20/98ISR Number: 3059533-5Report Type:Expedited (15-DaCompany Report #A0062241  
 Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Congenital Anomaly 150 MG / TWICE PER DAY	Complications Of Maternal Exposure To Therapeutic Drugs	Health Professional	Wellbutrin	PS		ORAL
/ ORAL	Exomphalos Multiple Cardiac Defects					

Date:03/20/98ISR Number: 3059534-7Report Type:Expedited (15-DaCompany Report #A0057927  
 Age:43 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / UNK Initial or Prolonged / ORAL	Asthma Dyspnoea Oedema Peripheral	Health Professional	Wellbutrin	PS		ORAL
			Vitamin E	C		
			Multivitamin	C		
			Prempro	C		
			Prednisone	C		
			Ticlopidine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Metoclopramide	C
Nabumetone	C
Losartan Potassium	C
Terazosin	
Hydrochloride	C
Diltiazem	
Hydrochloride	C
Omeprazole	C
Insulin	C

Date:03/23/98ISR Number: 3057561-7Report Type:Direct  
 Age:30 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis		Wellbutrin	PS		ORAL
150 MG POQ		Pruritus					
DAILY				Prozac	C		

Date:03/23/98ISR Number: 3057611-8Report Type:Direct  
 Age:38 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Fracture	Consumer	Wellbutrin	PS		
150 MG TWICE		Grand Mal Convulsion					
Hospitalization -		Joint Dislocation					
A DAY		Laceration					
Initial or Prolonged							
Disability							
Required							
Intervention to							
Prevent Permanent							
Impairment/Damage							

Date:03/24/98ISR Number: 3060328-7Report Type:Expedited (15-DaCompany Report #8-98005-018H  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Valve Disease	Consumer	Pondimin	PS		ORAL
20 MG TWICE		Headache					
DAILY ORAL		Hypoaesthesia		Buspar	SS		
20 MG THREE							
TIMES DAILY,A							
NUMBER OF							
YEARS							
30MG DAILY				Phentermine	SS		
THREE TIMES				Redux	SS		
PER DAY	1	MON					
THREE TIMES A				Wellbutrin	SS		
DAY "A NUMBER							
OF YEARS"							

Date:03/25/98ISR Number: 3129547-5Report Type:Periodic Company Report #8-98005-011H  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arrhythmia	Consumer	Redux	PS		ORAL
THREE TIMES		Drug Ineffective					
PER DAY ORAL	1	MON		Buspar	SS		
THREE TIMES		Headache					
PER DAY "A		Hypoaesthesia					
NUMBER OF		Paraesthesia Oral					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

YEARS "

2 TIMES PER

DAY

THREE TIMES

PER DAY ORAL

THREE TIMES

PER DAY

"NUMBER OF

YEARS "

Phentermine

SS

Pondimin Tablet

SS

ORAL

Wellbutrin

SS

Premarin

C

Date:03/25/98ISR Number: 3130531-6Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #8-97353-005B

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Consumer	Redux	PS		ORAL
Other		Amnesia					
15 MG TWICE		Condition Aggravated		Wellbutrin	SS		
DAILY ORAL		Diarrhoea					
150 MG TWICE		Dyspnoea		Dilacor Xl	C		
DAILY		Headache		Toprol	C		
		Nervousness					
		Neurosis					
		Palpitations					
		Tinnitus					

Date:03/25/98ISR Number: 3138469-5Report Type:Periodic  
Age:40 YR Gender:Female I/FU:F

Company Report #8-97296-001J

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Consumer	Redux	PS		ORAL
TWO TIMES		Drug Withdrawal Syndrome					
DAILY ORAL	6	MON		Wellbutrin	SS		
				Depakote	C		

Date:03/25/98ISR Number: 3148868-3Report Type:Periodic Company Report #8-97305-041H  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain	Health	Redux	PS		ORAL
15 MG DAILY		Drug Withdrawal Syndrome	Professional				
ORAL	1	YR		Desyrel	SS		ORAL
50 MG AT		Dyspnoea					
BEDTIME ORAL		Palpitations					
2 DAILY ORAL		Vasodilatation		Neurontin	SS		ORAL
2 DAILY ORAL				Ultram	SS		ORAL
400 DAILY				Wellbutrin	SS		ORAL
ORAL				Baclofen	C		
				Desyrel	C		
				Neurontin	C		
				Ultram	C		
				Vicodin	C		
				Wellbutrin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/98ISR Number: 3058427-9Report Type:Expedited (15-DaCompany Report #A0061715  
 Age:46 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Myocardial Infarction	Health	Zyban	PS		ORAL
150 MG / ORAL						
Hospitalization -	Vasospasm	Professional				
Initial or Prolonged		Company				
Other		Representative				

Date:03/30/98ISR Number: 3064847-9Report Type:Direct Company Report #  
 Age:50 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
2 TAB BID	Abdominal Pain		Wellbutrin	PS		
	Arthralgia		Prilosecv	C		
	Blister		Librax	C		
	Chromaturia		Lopressor	C		
	Diarrhoea		Tylenol #4	C		
	Dizziness		Vallium	C		
	Dry Mouth					
	Dysuria					
	Faeces Discoloured					
	Fatigue					
	Haematuria					
	Stomatitis					

Date:03/31/98ISR Number: 3063180-9Report Type:Direct Company Report #  
 Age:37 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Convulsion		Wellbutrin	PS		ORAL
100MG PO BID						
50MG PO HS			Nortriptyline	SS		ORAL

Date:03/31/98ISR Number: 3063240-2Report Type:Direct  
Age:37 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - PO 300 MG TID	Abdominal Pain		Wellbutrin	PS		ORAL
Initial or Prolonged ( FOR SOME TIME)	Constipation Vomiting					

Date:03/31/98ISR Number: 3067119-1Report Type:Direct  
Age:54 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other 2 TABLETS	Hypoaesthesia		Wellbutrin	PS	Glaxo Wellcome	
DAILY	3 WK					
	Oedema Peripheral Pain In Extremity Rash Erythematous Skin Exfoliation					

Date:04/01/98ISR Number: 3058889-7Report Type:Expedited (15-DaCompany Report #8-98009-010H  
Age:18 YR Gender:Female I/FU:F

Outcome	PT
Other	Cardiac Murmur Cardiac Valve Disease

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cardiomegaly Dyspnoea Ventricular Hypertrophy	Health	Redux	PS		ORAL
15 MG TWICE			Professional				
DAILY				Wellbutrin	SS		ORAL
150 MG TWICE							
DAILY				Provera	C		
				Synthroid	C		
				N/A	C		

Date:04/01/98ISR Number: 3063181-0Report Type:Expedited (15-DaCompany Report #8-98016-002S  
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Health	Pondimin	PS		ORAL
20-60 MG			Professional				
DAILY ORAL		Dysuria					
		Insomnia		Wellbutrin	SS		ORAL
400 MG DAILY							
ORAL		Ventricular Hypertrophy					
				Ionamin	SS		ORAL
15 MG DAILY							
ORAL				Loestrin 1/20	C		

Date:04/02/98ISR Number: 3059986-2Report Type:Expedited (15-DaCompany Report #A0062683  
Age:25 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Facial Palsy	Health	Zyban	PS		ORAL
150							

Date:04/02/98ISR Number: 3059987-4Report Type:Expedited (15-DaCompany Report #A0061890  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Clonic Convulsion	Health	Wellbutrin	PS		ORAL
UNK/ORAL			Professional Company Representative				

Date:04/02/98ISR Number: 3071857-4Report Type:Direct Company Report #  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain		Zyban	PS	Glaxo Wellcome	ORAL
150 MG ORALLY							

Date:04/02/98ISR Number: 3071950-6Report Type:Direct Company Report #  
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dermatitis		Bupropion	PS		ORAL
100MG /TI							
Initial or Prolonged		Pruritus					
TID/ORAL		Urticaria		Loraxepam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/03/98ISR Number: 3062720-3Report Type:Expedited (15-DaCompany Report #A0061756

Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Influenza Like Illness	Health	Wellbutrin	PS		ORAL
150 MG /			Professional				
TWICE PER							
DAY/ ORAL/							
TAB							
ORAL				Pseudoephedrine Hcl	SS		ORAL
				Cough Medication	C		

Date:04/06/98ISR Number: 3062918-4Report Type:Expedited (15-DaCompany Report #A0062864

Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous	Consumer	Zyban	PS		ORAL
150MG/ TWICE							
PER DAY/ ORAL							

Date:04/06/98ISR Number: 3062920-2Report Type:Expedited (15-DaCompany Report #A0062949

Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Asthenia	Consumer	Zyban	PS		ORAL
150MG/TWICE							
Hospitalization -		Collapse Of Lung					
PER DAY/ORAL							
Initial or Prolonged		Dyspnoea		Oxygen	C		
		Epistaxis		Dilantin	C		
		Insomnia		Blood Pressure			
		Nervousness		Medication	C		
				Nebulizer	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aortic Valve Incompetence	Consumer	Redux	PS		ORAL
THREE TIMES		Arrhythmia					
PER DAY ORAL	1	MON					
THREE TIMES		Feeling Abnormal		Buspar	SS		
PER DAY		Headache					
NUMBER OF		Hypoaesthesia					
YEARS		Hypoaesthesia Oral					
30 MG DAILY				Phentermine	SS		ORAL
ORAL							
20 MG 2 TO 3				Pondimin	SS		ORAL
TIMES DAILY							
ORAL							
THREE TIMES				Wellbutrin	SS		
PER DAY							
NUMBER OF							
YEARS							
				Buspar	C		
				Phentermine	C		
				Pondimin	C		
				Wellbutrin	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/08/98ISR Number: 3072565-6Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Mania		Bupropion	PS		ORAL
100 MG PO BID				Aripiprozole	SS		ORAL
30 MG PO QAM							
(NO LATER							
THEN 3/28)							

Date:04/09/98ISR Number: 3062490-9Report Type:Expedited (15-DaCompany Report #A0063085  
 Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea	Consumer	Zyban	PS		ORAL
300 MG/TWICE		Vomiting					
PE DAY/ORAL				Glibenclamide	C		
				Atorvastatin Calcium	C		
				Glucophage	C		
				Omeprazole	C		

Date:04/09/98ISR Number: 3062492-2Report Type:Expedited (15-DaCompany Report #A0063016  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Zyban	PS		ORAL
150 MG/ORAL			Professional				

Date:04/10/98ISR Number: 3073150-2Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Pancreatitis		Zyban	PS		
Initial or Prolonged							

Date:04/13/98ISR Number: 3063810-1Report Type:Expedited (15-DaCompany Report #A0063001  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphasia	Health	Wellbutrin	PS		ORAL
30		Overdose	Professional				
TABLET/SINGLE							
DOSE/ORAL							

Diazepam C

Date:04/13/98ISR Number: 3063861-7Report Type:Expedited (15-DaCompany Report #A0061492  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Convulsion	Consumer	Zyban	PS		ORAL
150 MG /		Cyanosis					
TWICE PER DAY							
/ ORAL							
TOPICAL 21 MG/							
DAY / TOPICAL							
		Disturbance In Attention		Nicotine Patch	SS		
		Fall					
		Head Injury		Paracetamol	C		
		Insomnia		Darvocet-N	C		
		Loss Of Consciousness					
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/14/98ISR Number: 3064324-5Report Type:Expedited (15-DaCompany Report #001-0991-980613

Age:78 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening PER ORAL	Angioneurotic Oedema	Health	Rezulin	PS		
Hospitalization - 300 MG	Dyspnoea	Professional	Zyban	SS		ORAL
Initial or Prolonged (DAILY) PER	Eyelid Oedema					
ORAL	Face Oedema					
	Pharyngeal Oedema		Amaryl Cardizem Compazine	C C C		

Date:04/15/98ISR Number: 3064474-3Report Type:Expedited (15-DaCompany Report #M98-093 (2)

Age:45 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 0.3 MG QHS	Anaphylactic Reaction	Health	Baycol	PS		ORAL
Hospitalization - ORAL	Angioneurotic Oedema	Professional				
Initial or Prolonged 150 MG BID	Pyrexia		Zyban	SS		ORAL
Required ORAL	Scrotal Oedema					
Intervention to Prevent Permanent Impairment/Damage	Urticaria		Prevacid Wellbutrin Xyban	C C C		

Date:04/15/98ISR Number: 3065424-6Report Type:Expedited (15-DaCompany Report #A0062572

Age:17 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL	Overdose	Health	Wellbutrin	PS		ORAL
Initial or Prolonged Other	Speech Disorder Suicide Attempt	Professional	Ethanol	C		

Date:04/15/98ISR Number: 3065444-1Report Type:Expedited (15-DaCompany Report #A0057797

Age:66 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/PER Initial or Prolonged DAY/ORAL	Blood Pressure Increased Dehydration Dizziness Hypersomnia Malaise Nausea Sedation Visual Disturbance	Health Professional	Zyban Conjugated Estrotens Ibuprofen	PS C C		ORAL

Date:04/15/98ISR Number: 3065446-5Report Type:Expedited (15-DaCompany Report #A0059118

Age:66 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/PER Initial or Prolonged DAY/ORAL	Blood Pressure Increased Dehydration Dizziness Hypersomnia Malaise Nausea Visual Disturbance	Health Professional	Zyban Conjugated Estrogens Ibuprofen	PS C C		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/20/98ISR Number: 3065289-2Report Type:Expedited (15-DaCompany Report #A0063509  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blindness Dyspnoea Hypersensitivity Pruritus Urinary Incontinence Urticaria	Consumer	Wellbutrin	PS		ORAL

Date:04/20/98ISR Number: 3065925-0Report Type:Expedited (15-DaCompany Report #A0063649  
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150 MG		Dermatitis	Health	Zyban	PS		ORAL
Initial or Prolonged (TABLET)		Stevens-Johnson Syndrome	Professional Company Representative				

Date:04/20/98ISR Number: 3065961-4Report Type:Expedited (15-DaCompany Report #A0063323  
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death 150 MG/PER		Agitation	Health	Wellbutrin	PS		ORAL
DAY/ORAL		Anxiety	Professional				
		Completed Suicide Diarrhoea Irritability Paranoia		Semisodium Valproate	C		

Date:04/21/98ISR Number: 3071889-6Report Type:Direct Company Report #  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - PO BID		Atrial Fibrillation		Zyban	PS		ORAL
Initial or Prolonged		Labile Blood Pressure Palpitations Ventricular Tachycardia		Digoxin Cardazen	C C		

Date:04/21/98ISR Number: 3071930-0Report Type:Direct Company Report #  
 Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG SR PO		Agitation		Wellbutrin	PS		ORAL
QD AM							

Date:04/21/98ISR Number: 3071940-3Report Type:Direct Company Report #  
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 3 WK		Urticaria		Wellbutrin Sr	PS		
				Prednisone	SS		
				Vitamin B	C		
				Vitamin C Supplements	C		



Other		Dermatitis		Zyban	PS	ORAL
PO	5	DAY				
		Pruritus				

Date:04/24/98ISR Number: 3068779-1Report Type:Expedited (15-DaCompany Report #A0063654  
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY		Condition Aggravated	Health	Zyban	PS		ORAL
/ ORAL		Drug Interaction	Professional				
		International Normalised Ratio Increased Phlebitis Prothrombin Time Prolonged		Warfarin Sodium Nabumetone	SS C		

Date:04/24/98ISR Number: 3072261-5Report Type:Direct Company Report #  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage		Grand Mal Convulsion		Bupropion	PS		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/24/98ISR Number: 3072353-0Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash Pruritic		Zyban	PS		ORAL
150 MG PO BID				Diazepam	C		
				Premarin	C		

Date:04/28/98ISR Number: 3072016-1Report Type:Direct  
 Age:25 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Arthralgia		Zyban	PS		ORAL
150MG PO BID		Dermatitis					
		Nausea					
		Oedema Peripheral					
		Pruritus					

Date:04/28/98ISR Number: 3072092-6Report Type:Direct  
 Age:47 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Blister	Health	Wellbutrin	PS		
100 MG BID		Pruritus	Professional				
		Rash Erythematous					
		Rash Maculo-Papular					

Date:04/28/98ISR Number: 3072573-5Report Type:Direct  
 Age:43 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Colitis	Consumer	Wellbutrin	PS		ORAL
2 QD	4 WK						
Required		Diarrhoea		Estradiol	C		

Intervention to  
Prevent Permanent  
Impairment/Damage

Date:04/28/98ISR Number: 3073809-7Report Type:Direct  
Age:48 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain Upper		Zyban	PS		ORAL
1PO BID		Burning Sensation Dermatitis Pyrexia					

Date:04/28/98ISR Number: 3153994-9Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0059397

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150 MG/TWICE		Anaphylactic Reaction	Health	Wellbutrin	PS		ORAL
Initial or Prolonged PER DAY/ORAL		Dyspnoea	Professional				
Other		Urticaria	Company Representative				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/28/98ISR Number: 3153996-2Report Type:Periodic Company Report #A0059398  
 Age:31 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening ORAL	Anaphylactic Reaction	Health	Wellbutrin	PS		ORAL
Hospitalization - Initial or Prolonged Other	Urticaria	Professional Company Representative	Lysine	C		

Date:04/28/98ISR Number: 3153997-4Report Type:Periodic Company Report #A0059419  
 Age:37 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100 MG/TWICE Initial or Prolonged PER DAY/ORAL	Confusional State Convulsion	Health Professional	Wellbutrin	PS		ORAL
	Disorientation Joint Stiffness Tremor		Pethidine Hydrochloride	C		

Date:04/28/98ISR Number: 3153999-8Report Type:Periodic Company Report #A0059754  
 Age:75 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL	Balance Disorder Dizziness	Consumer	Wellbutrin	PS		ORAL
Disability Other	Dysarthria Dyspraxia Headache Nausea Transient Ischaemic Attack		Aspirin Lithium Salt	C C		

Date:04/28/98ISR Number: 3154000-2Report Type:Periodic Company Report #A0060059  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin	PS		ORAL
ORAL		Overdose	Professional Company Representative				

Date:04/28/98ISR Number: 3154002-6Report Type:Periodic Company Report #A0060316  
Age:16 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Duodenal Ulcer	Health	Wellbutrin	PS		ORAL
150 MG/TWICE							
Hospitalization -		Duodenitis	Professional				
PER DAY/ORAL							
Initial or Prolonged		Gastritis		Ibuprofen	SS		
Other		Gastrointestinal Haemorrhage		Pancrelipase Vitamin E Ascorbic Acid	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/28/98ISR Number: 3154004-XReport Type:Periodic Company Report #A0060770  
 Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG/TWICE	Angioneurotic Oedema	Health	Wellbutrin	PS		ORAL
PER DAY/ORAL		Urticaria	Professional				
			Company Representative				

Date:04/28/98ISR Number: 3160400-7Report Type:Periodic Company Report #A0057121  
 Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150MG TWICE	Aggression Antisocial Behaviour Depression	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
PER DAY ORAL		Disturbance In Attention					
		Drug Ineffective		Hyzaar	C		
		Feeling Abnormal		Berocca	C		
		Irritability		Valsartan	C		
		Mood Altered					
		Sleep Disorder					

Date:04/28/98ISR Number: 3160491-3Report Type:Periodic Company Report #A0058281  
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged	150 MG / TWICE PER DAY/ ORAL	Anaphylactic Reaction Blood Pressure Decreased Dermatitis	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
Other		Dyspnoea					
		Hypoaesthesia Oral					
		Oedema Peripheral		Diphenhydramine Hcl			

Urticaria

(Formulation

Unknown)

SS

UNK/ UNK/ UNK

Dephenhydramine Hcl

Injection

SS

INTRAVENOUS

UNK/ UNK/

INTRAVENOUS

Sertraline

Hydrochloride

C

Date:04/28/98ISR Number: 3315033-6Report Type:Periodic

Company Report #A0076885

Age:14 YR Gender:Male

I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Health Professional	Wellbutrin Tablet-Controlled Released			
	PS		ORAL				

Date:04/28/98ISR Number: 3315035-XReport Type:Periodic

Company Report #A0077287

Age: Gender:Female

I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ORAL			Representative				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/29/98ISR Number: 3073029-6Report Type:Expedited (15-DaCompany Report #A0064080  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	400 ORAL PER DAY	Abdominal Distension	Health	Wellbutrin	PS		ORAL
		Dyspnoea	Professional				
		Dysuria		Fenfluramine Hcl	SS		ORAL
		Fall		Loestrin	C		
		Insomnia		Tramadol			
		Ventricular Hypertrophy		Hydrochloride	C		
				Naproxen	C		

Date:04/30/98ISR Number: 3073820-6Report Type:Direct Company Report #  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	1 BID PO; WAS TITRATED 1 QD X 3 DAYS, THEN 1 BID	Paraesthesia Oral	Health	Wellbutrin Sr	PS		ORAL
		Tongue Blistering	Professional				
		Tongue Oedema					

Date:04/30/98ISR Number: 3073937-6Report Type:Direct Company Report #  
 Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	100MG PO Q8H	Dermatitis		Bupropion	PS	Glaxo	ORAL

Date:05/01/98ISR Number: 3072979-4Report Type:Direct Company Report #  
 Age:38 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening			Abdominal Pain Upper		Wellbutrin	PS	Glaxo Welcome	
1 PILL A DAY			Blindness					
			Depression					
			Dry Mouth					
			Hallucination					
			Headache					
			Hostility					
			Irritability					
			Memory Impairment					
			Nervousness					
			Paranoia					
			Speech Disorder					
			Suicidal Ideation					

Date:05/01/98ISR Number: 3073610-4Report Type:Expedited (15-DaCompany Report #A0064472  
Age:70 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL			Hypernatraemia	Health	Wellbutrin	PS		ORAL
Initial or Prolonged				Professional				



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/01/98ISR Number: 3073637-2Report Type:Expedited (15-DaCompany Report #A0063323

Age:67 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abnormal Behaviour	Health	Wellbutrin	PS		ORAL
150 MG / PER DAY / ORAL		Agitation	Professional				
		Anxiety		Semisodium Valproate	C		
		Completed Suicide		Lansoprazole	C		
		Diarrhoea		Ascorbic Acid	C		
		Irritability		Vitamin B Complex	C		
		Mood Altered		Tranlycypromine			
		Paranoia		Sulphate	C		

Date:05/01/98ISR Number: 3080321-8Report Type:Direct

Age:28 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Urticaria		Bupropion Hcl	PS		

Date:05/01/98ISR Number: 3080564-3Report Type:Direct

Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG OD 3	Anxiety		Zyban	PS	Glaxo Wellcome	ORAL
DAYS		Decreased Appetite					
		Depressed Mood		Atenolol	C		
		Depression		Levothyroxine	C		
		Emotional Disorder		Glucotrol	C		
		Suicidal Ideation					
		Weight Decreased					

Date:05/01/98ISR Number: 3250183-4Report Type:Periodic

Age:49 YR Gender:Female I/FU:I

Company Report #JAUSA-31616

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Chest Pain Dizziness Dyspnoea Palpitations	Consumer	Hismanal (Astemizole), Janssen, Tablet 10 Mg	PS	Janssen	ORAL
10 MG 1 DAILY							
ORAL				Wellbutrin (Amfebutamone)	SS		ORAL
150 MG DAILY							
ORAL							

Date:05/04/98ISR Number: 3073449-XReport Type:Expedited (15-DaCompany Report #A0063965  
Age:3 MON Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 450 MG QD		Complications Of Maternal	Health	Wellbutrin	PS		ORAL
Initial or Prolonged ORAL TAB		Exposure To Therapeutic	Professional				
400 MG QD		Drugs		Lamictal	SS		ORAL
ORAL TAB		Jaundice					
				Buspirone Hydrochloride	C		
				Nabumetone	C		
				Thyroxine Sodium	C		
				Clonazepam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/04/98ISR Number: 3085634-1Report Type:Direct  
 Age:28 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - ONE QD THEN	Angioneurotic Oedema		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged INCREASE BIDI	Dermatitis					
PO	Oedema Peripheral					

Date:05/04/98ISR Number: 3090392-0Report Type:Expedited (15-DaCompany Report #A0065467  
 Age:80 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Death 150 MG	Deep Vein Thrombosis	Health	Wellbutrin	PS		ORAL
Life-Threatening Hospitalization -	Grand Mal Convulsion	Professional	Venlafaxine Hydrochloride	C		
Initial or Prolonged	Intentional Misuse		Alprazolam	C		
	Loss Of Consciousness		Thyroxine Sodium	C		
	Pneumonia		Nifedipine	C		
	Respiratory Failure		Sinemet	C		

Date:05/05/98ISR Number: 3088479-1Report Type:Direct  
 Age:75 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening Hospitalization -	Convulsion		Bupropion	PS		ORAL
75MG BID; 75 MG TID, ORAL						
Initial or Prolonged			Terazosin	C		
			Quinapril	C		
			Levothyroxine	C		
			Risperidone	C		

Date:05/06/98ISR Number: 3080753-8Report Type:Direct  
Age:58 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG BID PO		Grand Mal Convulsion	Health	Wellbutrin Sr	PS		ORAL
Initial or Prolonged 20 MG QAM			Professional	Prozac	SS		ORAL

Date:05/06/98ISR Number: 3080902-1Report Type:Direct  
Age:73 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 5MG PO QD	2 MON	Asthenia		Aricept	PS		ORAL
Initial or Prolonged 150MG PO QD	2 MON	Convulsion		Wellbutrin	SS		ORAL
		Grand Mal Convulsion Sleep Attacks Tremor					

Date:05/12/98ISR Number: 3074854-8Report Type:Direct  
Age:67 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 3X DAILY -		Dermatitis	Health	Prilosec	PS	Astra-Merck	ORAL
ORAL - 20MG		Pruritus	Professional				
2 X ORAL				Zyban	SS	Glaxo Welcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/13/98ISR Number: 3075607-7Report Type:Direct  
Age:45 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG PO BID 2 WK Initial or Prolonged	Convulsion Tremor		Zyban	PS		ORAL

Date:05/13/98ISR Number: 3075612-0Report Type:Direct  
Age:56 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG PO BID	Blister Feeling Hot Paraesthesia Pruritus Rash Erythematous		Bupropion Sr	PS	Glaxo Wellcome	ORAL

Date:05/13/98ISR Number: 3079000-2Report Type:Expedited (15-DaCompany Report #A0063654  
Age:46 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY Disability / ORAL Other 15 MG / PER DAY / UNK	Drug Interaction International Normalised Ratio Increased Phlebitis Prothrombin Time Prolonged	Health Professional	Zyban Warfarin Sodium Nabumetone	PS SS C		ORAL

Date:05/13/98ISR Number: 3079004-XReport Type:Expedited (15-DaCompany Report #A0065005  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health	Wellbutrin	PS		ORAL
UNK / UNK /			Professional				
ORAL			Company Representative				

Date:05/13/98ISR Number: 3079009-9Report Type:Expedited (15-DaCompany Report #A0064936  
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Angina Pectoris	Consumer	Zyban	PS		ORAL
150 MG /		Tachycardia					
Initial or Prolonged							
TWICE PER DAY							
/ ORAL				Blood Pressure Medication	C		

Date:05/14/98ISR Number: 3079126-3Report Type:Direct Company Report #  
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Anger	Health	Zyban	PS	Glaxo	ORAL
150 MG PO BID		Suicidal Ideation	Professional	Voltaren-Xr	C		
				Xanax	C		
				Methotrexate	C		
				Prednisone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/15/98ISR Number: 3079683-7Report Type:Expedited (15-DaCompany Report #A0060104  
Age:68 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL	Abdominal Pain	Health	Zyban	PS		ORAL
	Cholelithiasis	Professional	Sertraline Hydrochloride Metoprolol Succinate	C C		

Date:05/15/98ISR Number: 3079686-2Report Type:Expedited (15-DaCompany Report #A0064987  
Age:69 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG ORAL Initial or Prolonged TWICE PER DAY	Abdominal Pain	Consumer	Zyban	PS		ORAL
	Colitis					
	Condition Aggravated Diarrhoea Faecal Incontinence Fatigue Nervousness Weight Decreased White Blood Cell Count Increased					

Date:05/15/98ISR Number: 4456559-1Report Type:Direct Company Report #USP 51193  
Age:37 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Medication Error			Propulsid Restoril Colace Wellbutrin Pepcid	PS SS SS SS SS		

Date:05/18/98ISR Number: 3080005-6Report Type:Expedited (15-DaCompany Report #A0065101  
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	100 MG/SINGLE	Myocardial Infarction	Health	Wellbutrin	PS		ORAL
	DOSE/ORAL		Professional				

Date:05/18/98ISR Number: 3080008-1Report Type:Expedited (15-DaCompany Report #A0063649  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ ORAL		Conjunctivitis	Health	Zyban	PS		ORAL
Initial or Prolonged Other		Dermatitis Dermatitis Exfoliative Disability Dyspnoea Erythema Multiforme Eye Irritation Mouth Ulceration Stevens-Johnson Syndrome Urticaria	Professional Company Representative				



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/18/98ISR Number: 3080092-5Report Type:Expedited (15-DaCompany Report #A0062008

Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bundle Branch Block Left	Health	Wellbutrin	PS		ORAL
75 MG/ TWICE		Fatigue	Professional				
PER DAY/ ORAL		Heart Rate Irregular	Company				
		Suicidal Ideation	Representative				
		Ventricular Extrasystoles					
		Ventricular Tachycardia					

Date:05/18/98ISR Number: 3080093-7Report Type:Expedited (15-DaCompany Report #A0061053

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Death	Health	Wellbutrin	PS		ORAL
ORAL TAB		Drug Interaction	Professional	Beta-Blocker	C		
			Company				
			Representative				

Date:05/19/98ISR Number: 3079928-3Report Type:Direct

Company Report #

Age:10 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Lipase Increased	Health	Clonidine	PS		ORAL
0.1 MG BID			Professional	Wellbutrin	SS		ORAL
75 MG PO QD							

Date:05/22/98ISR Number: 3081788-1Report Type:Expedited (15-DaCompany Report #A0064080

Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death	Abdominal Distension	Health	Wellbutrin	PS	ORAL
400 MG / PER					
DAY / ORAL	Dyspnoea	Professional			
	YR				
20 MG / PER	Dysuria		Fenfluramine	SS	ORAL
DAY / ORAL	Insomnia				
15 MG / PER	Ventricular Hypertrophy		Phentermine	SS	ORAL
DAY / ORAL					
			Loestrin	C	
			Naproxen	C	

Date:05/22/98ISR Number: 3081861-8Report Type:Expedited (15-DaCompany Report #A0064577  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban	PS		ORAL
Other		Blood Creatine	Health				
150MG ORAL		Phosphokinase Increased	Professional	Cyclosporin	C		
		Myalgia	Company	Pyridostigmine			
		Myositis	Representative	Bromide	C		
				Doxepin	C		
				Alprazolam	C		

Date:05/26/98ISR Number: 3082953-XReport Type:Expedited (15-DaCompany Report #A0065467  
Age:80 YR Gender:Female I/FU:I

Outcome  
Death  
Life-Threatening  
  
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hospitalization -  
Initial or Prolonged

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG		Deep Vein Thrombosis	Health	Wellbutrin	PS		ORAL
		Grand Mal Convulsion	Professional	Venlafaxine			
		Loss Of Consciousness		Hydrochloride	C		
		Overdose		Alprazolam	C		
		Pneumonia		Thyroxine Sodium	C		
		Respiratory Failure		Nifedipine	C		
				Sinemet	C		

Date:05/26/98ISR Number: 3082955-3Report Type:Expedited (15-DaCompany Report #20711  
Age:71 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY			Blood Cholesterol	Health	Zyban	PS		ORAL
			Increased	Professional				
			Hypertension		Diltiazem			
			Palpitations		Hydrochloride	C		
6 YR					Losartan Potassium	C		

Date:05/26/98ISR Number: 3082957-7Report Type:Expedited (15-DaCompany Report #A0063931  
Age:34 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY ORAL			Agitation	Consumer	Zyban	PS		ORAL
			Confusional State					
			Convulsion		Unknown	C		
			Disorientation					
			Fatigue					
			Feeling Jittery					
			Hypertension					
			Insomnia					
			Irritability					

Motor Dysfunction  
Mouth Ulceration  
Pruritus  
Syncope  
Urticaria  
Visual Disturbance

Date:05/26/98ISR Number: 3084464-4Report Type:Direct  
Age:76 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Tremor	Health	Zyban	PS	Glaxo Wellcome	
150 MG I QD X			Professional				
3 DAYS THEN 1							

BID;NDC#

00173-00556-0

2

Date:05/28/98ISR Number: 3161424-6Report Type:Periodic  
Age:53 YR Gender:Female I/FU:F

Company Report #8-97353-005B

Outcome	PT
Other	Abdominal Pain Amnesia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Condition Aggravated	Report Source	Product	Role	Manufacturer	Route
15 MG TWICE	DAILY ORAL	Arrhythmia Chest Pain	Consumer	Redux	PS		ORAL
150 MG TWICE	DAILY	Depression Diarrhoea Dyspnoea		Wellbutrin	SS		
		Emotional Disorder Headache Nervousness Neurosis Oedema Peripheral Palpitations Syncope Tinnitus		Dilacor Xl Toprol	C C		

Date:05/28/98ISR Number: 3166234-1Report Type:Periodic Company Report #A0063808  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE	PER DAY/ ORAL	Dermatitis	Consumer	Zyban	PS		ORAL
		Pharyngeal Oedema Swelling Urticaria		Erythromyin	C		

Date:05/29/98ISR Number: 3085227-6Report Type:Expedited (15-DaCompany Report #A0064752  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE	PER DAY/ORAL	Dysarthria	Health Professional	Wellbutrin	PS		ORAL

Date:05/29/98ISR Number: 3085239-2Report Type:Expedited (15-DaCompany Report #A0065468  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Clonic Convulsion	Health	Zyban	PS		ORAL
150 MG ORAL							
Initial or Prolonged		Hyperreflexia Tremor	Professional				

Date:05/29/98ISR Number: 3085243-4Report Type:Expedited (15-DaCompany Report #A0065568  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Health	Zyban	PS		ORAL
ORAL							
		Connective Tissue Disorder Dermatitis Genital Ulceration Mouth Ulceration	Professional				

Date:05/29/98ISR Number: 3085249-5Report Type:Expedited (15-DaCompany Report #A0062683  
Age:25 YR Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Facial Palsy	Health	Zyban	PS		ORAL
150 MG ORAL							
			Professional				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3085478-0Report Type:Expedited (15-DaCompany Report #A0062175

Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG TWICE		Chest Pain	Health	Zyban	PS		ORAL
Initial or Prolonged PER DAY ORAL		Conversion Disorder	Professional				
		Coordination Abnormal		Digoxin	C		
		Crying		Atenolol	C		
		Depression					
		Fear					
		Pneumonia Mycoplasmal					
		Tuberculosis					

Date:05/29/98ISR Number: 3162219-XReport Type:Periodic Company Report #A0061513

Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150 MG/PER		Condition Aggravated	Consumer	Zyban	PS		ORAL
DAY/ORAL		Tremor					
				Conjugated Estrogens	C		
				Triamter + Hclthiazide	C		

Date:05/29/98ISR Number: 3162220-6Report Type:Periodic Company Report #A0061581

Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150 MG/PER		Anaphylactic Reaction	Health	Zyban Tablet-Zyban	PS		ORAL
Other DAY/ORAL	3 WK	Dysphagia	Professional				
		Pain		Loracarbef	C		
		Pruritus					
		Swelling					
		Urticaria					

Date:05/29/98ISR Number: 3162221-8Report Type:Periodic Company Report #A0061585  
Age:56 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150 MG/PER	Agitation	Consumer	Zyban	PS		ORAL
Initial or Prolonged DAY/ORAL	Cerebrovascular Accident					
	Excitability		Thyroxine Sodium	C		
	Restlessness		Hydrochlorothiazide	C		
			Nortriptyline Hcl	C		
			Alprazolam	C		

Date:05/29/98ISR Number: 3162222-XReport Type:Periodic Company Report #A0061638  
Age:60 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150 MG/ORAL	Convulsion	Health	Zyban Tablet-Zyban	PS		ORAL
Initial or Prolonged Other		Professional	Ipratropium Bromide	C		
			Orciprenaline			
			Sulphate	C		
			Nebulizer	C		
			Theophylline	C		
			Beclomethasone			
			Dipropion	C		
			Omeprazole	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Doxazosin Mesylate C  
 Oxygen C  
 Psyllium Husk C  
 Prednisone C  
 Cephazolin Sodium C

Date:05/29/98ISR Number: 3162223-1Report Type:Periodic Company Report #A0061681  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Convulsion	Consumer	Zyban Tablet-Zyban	PS		ORAL
Initial or Prolonged		Gingivitis Libido Decreased		Diclofenac Sodium Tramadol Hydrochloride Multivitamin	C C C		

Date:05/29/98ISR Number: 3162224-3Report Type:Periodic Company Report #A0065042  
 Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE		Angina Pectoris	Consumer	Zyban Tablet-Zyban	PS		ORAL
Initial or Prolonged PER DAY/ORAL		Increased Appetite		Multiple Medication Pravastatin Sodium	C C		

Date:05/29/98ISR Number: 3162225-5Report Type:Periodic Company Report #A0065208  
 Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG/ORAL		Hypersensitivity	Consumer	Zyban Tablet-Zyban	PS		ORAL

Date:05/29/98ISR Number: 3162226-7Report Type:Periodic Company Report #A0057442  
Age:61 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL	Anxiety Blood Pressure Increased Chest Pain Hypoaesthesia Nervousness Oedema Peripheral Paraesthesia Tachycardia	Health Professional	Zyban Conjugated Estrogens	PS C		ORAL

Date:05/29/98ISR Number: 3162227-9Report Type:Periodic Company Report #A0058434  
Age:57 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150 MG/PER Disability DAY/ORAL	Amnesia Convulsion Loss Of Consciousness Road Traffic Accident	Health Professional	Zyban Tablet-Zyban	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3162228-0Report Type:Periodic  
Age:65 YR Gender:Female I/FU:F

Company Report #A0058617

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban	PS		ORAL
Other		Dermatitis					
150 MG/TWICE		Herpes Simplex	Professional				
PER DAY/ORAL		Urticaria		Vitamin	C		
				Ibuprofen	C		

Date:05/29/98ISR Number: 3162229-2Report Type:Periodic  
Age:47 YR Gender:Female I/FU:F

Company Report #A0058629

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban Tablet-Zyban	PS		ORAL
Hospitalization -		Dermatitis					
150 MG/TWICE		Hypersensitivity	Professional				
Initial or Prolonged							
PER DAY/ORAL							

Date:05/29/98ISR Number: 3162230-9Report Type:Periodic  
Age:70 YR Gender:Female I/FU:F

Company Report #A0058920

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet-Zyban	PS		ORAL
Hospitalization -		Accommodation Disorder					
150 MG/TWICE		Dizziness					
Initial or Prolonged		Dysgraphia					
PER DAY/ORAL		Face Oedema					
		Headache					
		Insomnia					
		Tinnitus					
		Tremor					

Date:05/29/98ISR Number: 3162231-0Report Type:Periodic  
Age:48 YR Gender:Female I/FU:F

Company Report #A0058975

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Mucosal Inflammation	Health	Zyban Tablet-Zyban	PS		ORAL
150 MG/ORAL		Pruritus	Professional				
		Rash Erythematous					
		Urticaria					

Date:05/29/98ISR Number: 3162232-2Report Type:Periodic Company Report #A0059110  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema	Health	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE		Fatigue	Professional				
PER DAY/ORAL		Mental Impairment		Claritin-D	C		
		Tachycardia		Cisapride	C		
		Tremor		Triamcinolone			
		Urticaria		Acetonide	C		
				Ovral	C		

Date:05/29/98ISR Number: 3163038-0Report Type:Periodic Company Report #A0061047  
Age: Gender: I/FU:I

Outcome	PT	Report Source
Hospitalization -	Angioneurotic Oedema	Health
Initial or Prolonged	Hypersensitivity	Professional
	Urticaria	Company

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Representative

Dose	Duration	Product	Role	Manufacturer	Route
150 MG / UNK		Zyban Tablet-Zyban	PS		ORAL
/ ORAL					

Date:05/29/98ISR Number: 3163206-8Report Type:Periodic Company Report #A0060935  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY		Arthralgia	Consumer	Zyban Tablet-Zyban	PS		ORAL
/ ORAL		Urticaria					

Date:05/29/98ISR Number: 3163208-1Report Type:Periodic Company Report #A0060931  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY		Anaphylactic Reaction	Health	Zyban Tablet-Zyban	PS		ORAL
Other / ORAL		Dermatitis	Professional				
		Dyspnoea	Company				
		Oedema	Representative	Ranitidine			
		Pruritus		Hydrochloride	C		
				Metoprolol Tartrate	C		

Date:05/29/98ISR Number: 3163209-3Report Type:Periodic Company Report #A0060908  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 150 MG / ORAL Initial or Prolonged	Convulsion	Health	Zyban Tablet-Zyban	PS	ORAL
		Professional Company Representative			
Date:05/29/98ISR Number: 3163211-1Report Type:Periodic		Company Report #A0060907			
Age:	Gender:Male	I/FU:I			
Outcome	PT	Report Source	Product	Role	Manufacturer
Dose	Duration				Route
Hospitalization - 150 MG / ORAL Initial or Prolonged Other	Convulsion	Health	Zyban Tablet-Zyban	PS	ORAL
		Professional Company Representative			
Date:05/29/98ISR Number: 3163212-3Report Type:Periodic		Company Report #A0060861			
Age:35 YR	Gender:Female	I/FU:I			
Outcome	PT	Report Source	Product	Role	Manufacturer
Dose	Duration				Route
Other	Agitation	Health	Zyban Tablets -Zyban	PS	ORAL
150 MG /	Anxiety	Professional			
TWICE PER DAY	Disturbance In Attention				
/ ORAL	Insomnia		Oral Contraceptive	C	
	Nervousness				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3164388-4Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #A0057814

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG TWICE	Agitation	Consumer	Zyban	PS		ORAL
PER DAY/	ORAL	Bronchitis					
		Cough		Hrt	C		
		Depression		Amitriptyline Hcl	C		
		Difficulty In Walking		Decongestant	C		
		Dyspraxia					
		Hallucination					
		Headache					
		Nervousness					
		Speech Disorder					
		Tremor					

Date:05/29/98ISR Number: 3164391-4Report Type:Periodic  
Age:75 YR Gender:Male I/FU:I

Company Report #A0058368

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG/SINGLE	Blindness	Health	Zyban	PS		ORAL
Initial or Prolonged	DOSE ORAL	Rectal Haemorrhage	Professional				
		Thrombosis		Terazosin			
				Hydrochloride	C		
				Pravastatin Sodium	C		
				Aspirin	C		
				Allopurinol	C		

Date:05/29/98ISR Number: 3164458-0Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #A0063074

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG/TWICE	Agitation	Consumer	Zyban	PS		ORAL
Initial or Prolonged	PER DAY/ORAL	Anger					

Disorientation  
Drug Toxicity  
Feeling Abnormal  
Nervousness

Captopril C  
Isosorbide Dinitrate C  
Simvastatin C  
Thyroxine Sodium C  
Lansoprazole C  
Alprazolam C  
Paracetamol C

Date:05/29/98ISR Number: 3164461-0Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0063111

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL	Anaphylactic Reaction Anaphylactic Shock Arthralgia Dermatitis Hypersensitivity Musculoskeletal Stiffness Pneumatic Compression Therapy Pruritus Swelling Syncope Urticaria	Consumer	Zyban  Ibuprofen	PS  C		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3164464-6Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0063297

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Health	Zyban	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL		Hypersensitivity	Professional				
		Oedema Peripheral	Company	Atenolol	C		
		Periorbital Oedema	Representative	Medroxyprogesterone			
		Pruritus		Ace.	C		
		Pyrexia					
		Urticaria					

Date:05/29/98ISR Number: 3164530-5Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #A0063779

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Agitation	Consumer	Zyban	PS		ORAL
TABLET, 150							
Initial or Prolonged		Memory Impairment					
MG / TWICE							
PER DAY /		Tremor					
ORAL							

Date:05/29/98ISR Number: 3164533-0Report Type:Periodic  
Age:62 YR Gender:Male I/FU:I

Company Report #A0063660

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Abdominal Pain	Health	Zyban	PS		ORAL
TABLET, 150							
Initial or Prolonged		Pancreatitis Acute	Professional				
MG / ORAL							
				Temazepam	C		

Date:05/29/98ISR Number: 3164536-6Report Type:Periodic Company Report #A0063622  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Tinnitus	Health	Zyban	PS		ORAL
TABLET, 150							
Professional							
MG / TWICE							
PER DAY /							
ORAL							

Date:05/29/98ISR Number: 3164540-8Report Type:Periodic Company Report #A0063591  
Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Dermatitis	Consumer	Zyban	PS		ORAL
TABLET, 150							
Dyspnoea							
MG / TWICE							
Hypersensitivity							
PER DAY /							
Muscle Spasms							
ORAL							
Muscular Weakness							
Oesophageal Disorder							
Swelling							

Date:05/29/98ISR Number: 3164543-3Report Type:Periodic Company Report #A0064074  
Age:27 YR Gender:Male I/FU:I

Outcome	PT
Life-Threatening	Anaphylactic Reaction
Hospitalization -	Chest Pain
Initial or Prolonged	Dyspepsia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pain In Extremity

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
TABLET, 150		Consumer	Zyban	PS		ORAL
MG / TWICE						
PER DAY /						
ORAL						

Date:05/29/98ISR Number: 3164546-9Report Type:Periodic Company Report #A0064342  
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Dermatitis	Consumer	Zyban	PS		ORAL
TABLET, 150							
Initial or Prolonged		Diarrhoea					
MG / TWICE							
PER DAY /		Dysphonia					
ORAL		Face Oedema					
		Nausea					
		Oedema					
		Periorbital Oedema					
		Pruritus					
		Pyrexia					
		Urticaria					

Date:05/29/98ISR Number: 3164551-2Report Type:Periodic Company Report #A0064347  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Serum Sickness	Health	Zyban	PS		ORAL
TABLET, 150							
Initial or Prolonged			Professional				
MG /ORAL			Company	Ibuprofen	C		

Representative

Date:05/29/98ISR Number: 3164558-5Report Type:Periodic Company Report #A0062747  
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Amnesia	Health	Zyban	PS		ORAL
Other		Epileptic Aura	Professional				
MG / TWICE		Feeling Abnormal	Company				
PER DAY /		Grand Mal Convulsion	Representative				
ORAL		Head Injury		Conjugated Estrogens	C		
		Headache		Ethanol	C		
		Laceration					

Date:05/29/98ISR Number: 3164561-5Report Type:Periodic Company Report #A0062744  
 Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Arthralgia	Health	Zyban	PS		ORAL
TABLET, 150		Dermatitis	Professional				
MG / TWICE							
PER DAY /							
ORAL	2 WK			Omeprazole	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3164565-2Report Type:Periodic Company Report #A0062673  
 Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaphylactic Reaction	Health	Zyban	PS		ORAL
TABLET, 150		Chills	Professional				
MG / TWICE		Flushing					
PER DAY /		Heart Rate Increased					
ORAL		Hypotension					
		Pruritus					
		Swelling					
		Tongue Oedema					
		Tremor					
		Urticaria					

Date:05/29/98ISR Number: 3164568-8Report Type:Periodic Company Report #A0062647  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Rash Generalised	Health	Zyban	PS		ORAL
TABLET, 150			Professional				
Initial or Prolonged			Company				
MG / ORAL			Representative				

Date:05/29/98ISR Number: 3164569-XReport Type:Periodic Company Report #A0062644  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion	Health	Zyban	PS		ORAL
TABLET, 150			Professional				
Initial or Prolonged			Company				
MG / ORAL			Representative				

Date:05/29/98ISR Number: 3164572-XReport Type:Periodic Company Report #A0062615  
Age:56 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - TABLET, 150 Initial or Prolonged MG / TWICE PER DAY / ORAL	Amnesia Choking Diarrhoea Dizziness Dry Mouth Insomnia Mental Impairment Pneumonia Weight Decreased	Consumer	Zyban          Librium Cyclobenzaprine Hcl Anticoagulant Diltiazem Hydrochloride Simvastatin Cimetidine	PS          C C C  C C C		ORAL

Date:05/29/98ISR Number: 3164578-0Report Type:Periodic Company Report #A0063053  
Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - TABLET, 150 Initial or Prolonged MG / ORAL	Chest Pain Dyspnoea	Health Professional	Zyban	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3164580-9Report Type:Periodic  
 Age:60 YR Gender:Male I/FU:I

Company Report #A0063042

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - TABLET, 150		Chest Pain	Health	Zyban	PS		ORAL
Initial or Prolonged MG / PER DAY		Dizziness	Professional				
/ ORAL	4 WK	Dyspnoea		Nifedipine	C		

Date:05/29/98ISR Number: 3164584-6Report Type:Periodic  
 Age:28 YR Gender:Female I/FU:I

Company Report #A0062867

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other TABLET, 150		Convulsion	Consumer	Zyban	PS		ORAL
MG / TWICE		Diarrhoea	Company				
PER DAY /		Disorientation	Representative				
ORAL		Syncope					
		Vomiting					

Date:05/29/98ISR Number: 3164585-8Report Type:Periodic  
 Age:59 YR Gender:Female I/FU:I

Company Report #A0064892

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - TABLET, 150		Chest Pain	Health	Zyban	PS		ORAL
Initial or Prolonged MG / TWICE		Convulsion	Professional				
PER DAY /		Muscle Twitching					
ORAL							

Date:05/29/98ISR Number: 3164587-1Report Type:Periodic Company Report #A0064923  
Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - TABLET, 150 Initial or Prolonged MG / TWICE PER DAY / ORAL	Delirium Fall Laceration Vision Blurred	Consumer	Zyban     Vitamin	PS     C		ORAL

Date:05/29/98ISR Number: 3164590-1Report Type:Periodic Company Report #A0064852  
Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - TABLET, 150 Initial or Prolonged MG / ORAL	Coronary Artery Disease Myocardial Infarction	Consumer	Zyban	PS		ORAL

Date:05/29/98ISR Number: 3164593-7Report Type:Periodic Company Report #A0061438  
Age:75 YR Gender:Male I/FU:F

Outcome	PT
Other	Blood Pressure Decreased Chest Discomfort Drug Interaction Ecchymosis Excoriation Fall Head Injury Malaise



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Syncope

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
TABLET, 150		Health Professional	Zyban	PS		ORAL
MG / PER DAY			Aspirin	C		
/ ORAL			Digoxin	C		
			Diclofenac	C		
			Amlodipine	C		

Date:05/29/98ISR Number: 3165439-3Report Type:Periodic Company Report #A0063910  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS		ORAL
150 MG; PER							
DAY; ORAL							

Date:05/29/98ISR Number: 3165442-3Report Type:Periodic Company Report #A0063911  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS		ORAL
150 MG;		Pruritus					
TWICE PER		Urticaria					
DAY; ORAL	2 WK						

Date:05/29/98ISR Number: 3165444-7Report Type:Periodic Company Report #A0063912  
 Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bronchitis	Consumer	Zyban	PS		ORAL
150 MG; TWICE		Chest Discomfort					
PER DAY;		Dyspnoea					
ORAL							

Date:05/29/98ISR Number: 3165446-0Report Type:Periodic Company Report #A0063913  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS		ORAL
150 MG;							
ORAL							

Date:05/29/98ISR Number: 3165448-4Report Type:Periodic Company Report #A0063914  
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS		ORAL
150 MG;		Joint Swelling					
THREE TIMES		Oedema Peripheral					
PER DAY;		Pruritus					
ORAL		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3165449-6Report Type:Periodic Company Report #A0063920  
 Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS		ORAL
150 MG;		Pruritus					
TWICE PER		Tenderness					
DAY;	ORAL						

Date:05/29/98ISR Number: 3165450-2Report Type:Periodic Company Report #A0063921  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS		ORAL
150 MG; PER		Dizziness					
DAY;	ORAL	Drug Ineffective					
		Emotional Disorder					
		Headache					
		Hypoaesthesia					

Date:05/29/98ISR Number: 3165451-4Report Type:Periodic Company Report #A0063922  
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Consumer	Zyban	PS		ORAL
150 MG;		Dermatitis					
TWICE PER		Pruritus					
DAY;	ORAL	Urticaria					

Date:05/29/98ISR Number: 3165452-6Report Type:Periodic Company Report #A0063923  
 Age:43 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG;		Face Oedema	Consumer	Zyban	PS		ORAL
ORAL	5 DAY	Pruritus					
		Rash Erythematous Swelling		Nicotine	C		
		Urticaria					

Date:05/29/98ISR Number: 3165453-8Report Type:Periodic Company Report #A0063924  
Age:54 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG;		Blood Pressure Increased	Consumer	Zyban	PS		ORAL
TWICE PER DAY;	ORAL	Dizziness					
		Dry Mouth					
		Nervousness		Diltiazem			
		Rash Pruritic		Hydrochloride	C		
				Amlodipine	C		
				Nitroglycerin	C		
				Oxpentifylline	C		
				Triazolam	C		
				Aspirin	C		
				Fluoxetine			
				Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3165454-XReport Type:Periodic  
Age:30 YR Gender:Male I/FU:I

Company Report #A0063925

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS		ORAL
150 MG;		Tachycardia					
TWICE PER							
DAY; ORAL				Vitamin	C		

Date:05/29/98ISR Number: 3165455-1Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0063926

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Consumer	Zyban	PS		ORAL
150 MG;		Disturbance In Attention					
TWICE PER		Dry Mouth					
DAY; ORAL		Insomnia		Oestradiol	C		
		Mood Swings					
		Nervousness					

Date:05/29/98ISR Number: 3165456-3Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0063927

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS		ORAL
150 MG; PER							
DAY; ORAL							

Date:05/29/98ISR Number: 3165457-5Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #A0063929

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Petit Mal Epilepsy	Health Professional	Zyban	PS		ORAL
150MG;	TWICE						
PER DAY;							
ORAL						Multiple Medication	C

Date:05/29/98ISR Number: 3165458-7Report Type:Periodic Company Report #A0063944  
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bone Pain	Consumer	Zyban	PS		ORAL
150 MG;		Cough					
TWICE PER		Dermatitis					
DAY; ORAL		Dry Mouth		Aspirin	C		
		Parosmia		Thiamine			
		Pruritus		Hydrochloride	C		
				Ascorbic Acid	C		
				Simvastatin	C		
				Inhaler	C		

Date:05/29/98ISR Number: 3165459-9Report Type:Periodic Company Report #A0063947  
Age:67 YR Gender:Female I/FU:I

Outcome	PT
	Difficulty In Walking
	Dizziness
	Insomnia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG;		Consumer	Zyban	PS		ORAL
TWICE PER						
DAY;	ORAL					
			Bayer	C		
			Centrum Silver	C		
			Hyzaar	C		

Date:05/29/98ISR Number: 3165461-7Report Type:Periodic Company Report #A0063948  
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Burning Sensation	Consumer	Zyban	PS		ORAL
150 MG;		Dermatitis					
TWICE PER		Pruritus					
DAY;	ORAL	Urticaria					
				Hormones	C		
				Fiorinal	C		

Date:05/29/98ISR Number: 3165463-0Report Type:Periodic Company Report #A0063953  
 Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS		ORAL
150 MG;	PER	Anxiety					
DAY;	ORAL	Dyspnoea					

Date:05/29/98ISR Number: 3165465-4Report Type:Periodic Company Report #A0063954  
 Age:40 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG;		Abnormal Dreams	Consumer	Zyban	PS		ORAL
TWICE PER		Dermatitis					
DAY;	ORAL	Dry Skin					

Date:05/29/98ISR Number: 3165467-8Report Type:Periodic Company Report #A0063961  
Age:55 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG;		Headache	Consumer	Zyban	PS		ORAL
TWICE PER		Paraesthesia					
DAY;	ORAL	Tremor					
				Oestradiol	C		
				Medroxyprogesterone			
				Ace.	C		

Date:05/29/98ISR Number: 3166154-2Report Type:Periodic Company Report #A0063872  
Age:32 YR Gender:Female I/FU:I

Outcome	PT
	Aggression
	Agitation
	Crying
	Depression
	Drug Ineffective



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Drug Withdrawal Syndrome  
Emotional Disorder

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG /		Consumer	Zyban	PS		ORAL
TWICE PER DAY						
/ ORAL						
			Ibuprofen	C		
			Medroxyprogesterone	C		
			Ace.			

Date:05/29/98ISR Number: 3166155-4Report Type:Periodic Company Report #A0063874  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS		ORAL
150 MG / PER		Vomiting					
DAY / ORAL				Oral Contraceptive	C		

Date:05/29/98ISR Number: 3166157-8Report Type:Periodic Company Report #A0063876  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Face Oedema	Consumer	Zyban	PS		ORAL
150 MG/TWICE		Influenza Like Illness					
PER DAY/ ORAL		Oedema Peripheral		Vitamin	C		
		Pain					
		Pyrexia					
		Urticaria					

Date:05/29/98ISR Number: 3166160-8Report Type:Periodic Company Report #A0063878  
Age:35 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Insomnia	Consumer	Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL							

Oral Contraceptive	C
Decongestant	C

Date:05/29/98ISR Number: 3166161-XReport Type:Periodic Company Report #A0063879  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Burning Sensation	Consumer	Zyban	PS		ORAL
150 MG / ORAL		Dizziness		Wellbutrin	SS		ORAL
Formication							
Hypoaesthesia							

Date:05/29/98ISR Number: 3166165-7Report Type:Periodic Company Report #A0063882  
 Age:46 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Anxiety	Health	Zyban	PS		ORAL
Panic Disorder							
Professional							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3166167-0Report Type:Periodic Company Report #A0063883  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amenorrhoea	Health	Zyban	PS		ORAL
150 MG / ORAL			Professional Company Representative				

Date:05/29/98ISR Number: 3166168-2Report Type:Periodic Company Report #A0063884  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amenorrhoea	Health	Zyban	PS		ORAL
150 MG / ORAL			Professional Company Representative				

Date:05/29/98ISR Number: 3166170-0Report Type:Periodic Company Report #A0063885  
 Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Zyban	PS		ORAL
ORAL							

Date:05/29/98ISR Number: 3166172-4Report Type:Periodic Company Report #A0063886  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban	PS		ORAL
150 MG /PER		Dysgeusia					
DAY/ ORAL				Me-Prednisolone Na Succ.	C		

Diazepam

C

Date:05/29/98ISR Number: 3166174-8Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #A0063888

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER		Tremor	Consumer	Zyban	PS		ORAL
DAY / ORAL	5 DAY			Benazepril	C		
				Alprazolam	C		

Date:05/29/98ISR Number: 3166176-1Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #A0063900

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Depression	Consumer	Zyban	PS		ORAL
TWICE PER				Diphenhydramine Hcl	C		
DAY/ ORAL	35 DAY			Diazepam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3166178-5Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #A0063901

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS		ORAL
150 MG /		Dry Mouth					
TWICE PER		Emotional Disorder					
DAY/ ORAL		Hallucination, Auditory					
		Insomnia		Tamoxifen	C		
		Paranoia		Diuretic	C		
		Pruritus		Potassium Salt	C		

Date:05/29/98ISR Number: 3166179-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0063902

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS		ORAL
150 MG /		Asthenia					
TWICE PER DAY		Chills					
/ ORAL		Dehydration					
		Night Sweats					
		Pyrexia					

Date:05/29/98ISR Number: 3166181-5Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #A0063903

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS		ORAL
150 MG /		Chest Discomfort					
TWICE PER		Condition Aggravated					
DAY/ ORAL		Dizziness					
		Feeling Cold		Paroxetine			
		Hyperhidrosis		Hydrochloride	C		
		Insomnia		Clonazepam	C		

Nausea  
Rhinorrhoea  
Vomiting

Warfarin Sodium C  
Prednisone C  
Methotrexate C  
Torasemide C  
Potassium Chloride C  
Folic Acid C  
Zolpidem Tartrate C  
Vicodin Es C

Date:05/29/98ISR Number: 3166183-9Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0063904

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Accommodation Disorder	Consumer	Zyban	PS		ORAL
		Communication Disorder Constipation Disturbance In Attention Feeling Abnormal Hyperhidrosis Insomnia		Thyroxine Sodium	C		

Date:05/29/98ISR Number: 3166184-0Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #A0063905

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY /		Dyspepsia Insomnia	Consumer	Zyban	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL

Claritin-D

C

Date:05/29/98ISR Number: 3166186-4Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #A0063907

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ PER		Faeces Discoloured	Consumer	Zyban	PS		ORAL
DAY / ORAL							

Date:05/29/98ISR Number: 3166189-XReport Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0063908

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Anxiety	Consumer	Zyban	PS		ORAL
TWICE PER DAY		Tremor					
/ ORAL							

Date:05/29/98ISR Number: 3166191-8Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #A0063909

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Drug Ineffective	Consumer	Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL	2 WK						

Date:05/29/98ISR Number: 3166194-3Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0063963

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG;	PER DAY;	Headache	Consumer	Zyban	PS		ORAL
				Omeprazole	C		
				Theophylline	C		
				Antihistamine	C		

Date:05/29/98ISR Number: 3166196-7Report Type:Periodic Company Report #A0063966  
 Age:44 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG;	TWICE PER DAY;	Keratoconjunctivitis	Consumer	Zyban	PS		ORAL
		Sicca					
		Urticaria					
				Verapamil			
				Hydrochloride	C		
				Conjugated Estrogens	C		

Date:05/29/98ISR Number: 3166198-0Report Type:Periodic Company Report #A0063967  
 Age:42 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG;	TWICE PER DAY;	Confusional State	Consumer	Zyban	PS		ORAL
		Disorientation					
		Dissociation					
		Insomnia					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3166200-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0063968

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS		ORAL
150 MG;	ORAL 12 DAY	Drug Ineffective					
		Dry Throat					
		Insomnia					
		Tremor					

Date:05/29/98ISR Number: 3166202-XReport Type:Periodic  
 Age:70 YR Gender:Female I/FU:I

Company Report #A0063969

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS		ORAL
150MG;	TWICE	Diarrhoea					
		Pruritus					
		Weight Decreased					
				Felodipine	C		
PER DAY;							
ORAL							

Date:05/29/98ISR Number: 3166204-3Report Type:Periodic  
 Age:32 YR Gender:Female I/FU:I

Company Report #A0063970

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS		ORAL
150 MG;		Abdominal Pain					
		Diarrhoea					
		Myalgia					
		Nausea					
TWICE PER							
DAY; ORAL							

Date:05/29/98ISR Number: 3166206-7Report Type:Periodic  
 Age:46 YR Gender:Male I/FU:I

Company Report #A0063971

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction	Consumer	Zyban	PS		ORAL
150 MG;							
TWICE PER							
DAY;	ORAL	2	WK				

Date:05/29/98ISR Number: 3166208-0Report Type:Periodic Company Report #A0063972  
 Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS		ORAL
150 MG;		Anorexia					
TWICE PER		Anxiety					
DAY;	ORAL	Depression		Ibuprofen	C		

Date:05/29/98ISR Number: 3166210-9Report Type:Periodic Company Report #A0063988  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS		ORAL
150 MG;		Insomnia					
TWICE PER		Pruritus					
DAY;	ORAL			Conjugated Estrogens	C		
				Omeprazole	C		
				Maxzide	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3166212-2Report Type:Periodic  
Age:65 YR Gender:Male I/FU:I

Company Report #A0063989

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS		ORAL
150MG;	TWICE	Tinnitus					
PER DAY;							
ORAL				Atenolol	C		
				Naproxen	C		
				Aspirin	C		

Date:05/29/98ISR Number: 3166213-4Report Type:Periodic  
Age:17 YR Gender:Male I/FU:I

Company Report #A0064001

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Consumer	Zyban	PS		ORAL
150 MG;	ORAL						

Date:05/29/98ISR Number: 3166215-8Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0064002

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Immobile	Consumer	Zyban	PS		ORAL
150MG	PER	Myalgia					
DAY	ORAL	Pain					

Date:05/29/98ISR Number: 3166217-1Report Type:Periodic  
Age:64 YR Gender:Male I/FU:I

Company Report #A0064004

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Consumer	Zyban	PS		ORAL
150 MG;							

TWICE PER  
 DAY; ORAL  
 Dizziness  
 Flushing  
 Headache  
 Nervousness  
 Diltiazem  
 Hydrochloride C  
 Atenolol C  
 Pravastatin Sodium C  
 Omeprazole C  
 Aspirin C  
 Isosorbide  
 Mononitrate C

Date:05/29/98ISR Number: 3166218-3Report Type:Periodic Company Report #A0064005  
 Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG	TWICE	Abdominal Pain Upper	Consumer	Zyban	PS		ORAL
PER DAY;		Anorexia					
ORAL		Asthenia					
		Diarrhoea		Nifedipine	C		
				Alprazolam	C		
				Cimetidine	C		
				Lorcet	C		
				Cisapride	C		
				Clonidine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3166221-3Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0064011

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pain	Consumer	Zyban	PS		ORAL
150 MG;	PER						
DAY;	ORAL						
				Ketoprofen	C		
				Indomethacin	C		
				Morphine	C		
				Baclofen	C		

Date:05/29/98ISR Number: 3166222-5Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0064012

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS		ORAL
150MG;	TWICE						
PER DAY;		Hypoaesthesia					
ORAL		Pruritus					

Date:05/29/98ISR Number: 3166225-0Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0064013

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity	Consumer	Zyban	PS		ORAL
150 MG							
TWICE PER DAY		Urticaria					
ORAL							

Date:05/29/98ISR Number: 3166226-2Report Type:Periodic  
Age:37 YR Gender:Male I/FU:I

Company Report #A0064014

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mood Altered	Consumer	Zyban	PS		ORAL
150MG	TWICE	Urticaria					
PER DAY							
ORAL							

Date:05/29/98ISR Number: 3166227-4Report Type:Periodic Company Report #A0064015  
 Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS		ORAL
150MG;							
TWICE PER							
DAY;	ORAL			Hormones	C		

Date:05/29/98ISR Number: 3166230-4Report Type:Periodic Company Report #A0064022  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Zyban	PS		ORAL
150 MG;							
TWICE PER							
DAY;	ORAL			Haloperidol	C		
				Paroxetine	C		
				Hydrochloride	C		
				Vitamin	C		
				Gabapentin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3166232-8Report Type:Periodic  
Age:72 YR Gender:Male I/FU:I

Company Report #A0063803

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY / ORAL		Depression Disorientation Insomnia Tremor	Consumer	Zyban	PS		ORAL
				Triamcinolone Acetonide	C		
				Zafirlukast	C		
				Insulin	C		
				Salbutamol Sulphate	C		
				Salmeterol Xinafoate	C		

Date:05/29/98ISR Number: 3166235-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0063810

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Abdominal Pain Upper Anxiety Dyspnoea Insomnia Nervousness Paranoia Tachycardia Tremor	Consumer	Zyban	PS		ORAL

Date:05/29/98ISR Number: 3166236-5Report Type:Periodic  
Age:67 YR Gender:Male I/FU:I

Company Report #A0063815

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Benign Prostatic	Consumer	Zyban	PS		ORAL

TWICE PER DAY  
 / ORAL  
 Hyperplasia  
 Confusional State  
 Dysuria  
 Headache  
 Micturition Urgency  
 Pollakiuria  
 Tremor  
 Hydrochlorothiazide C  
 Piroxicam C  
 Diazepam C

Date:05/29/98ISR Number: 3166237-7Report Type:Periodic Company Report #A0063827  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL				Salbutamol Sulphate	C		
				Azithromycin	C		

Date:05/29/98ISR Number: 3166238-9Report Type:Periodic Company Report #A0063828  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS		ORAL
150 MG /		Chest Discomfort					
TWICE PER DAY		Drug Ineffective					
/ ORAL		Flushing		Herbal Medication	C		



Freedom Of Information (FOI) Report

Thyroid Medication C

Date:05/29/98ISR Number: 3166240-7Report Type:Periodic Company Report #A0063829  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS		ORAL
150 MG /		Irritability					
TWICE PER DAY		Mood Altered					
/ ORAL		Sedation					

Date:05/29/98ISR Number: 3166241-9Report Type:Periodic Company Report #A0063830  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS		ORAL
150 MG/	ORAL						

Date:05/29/98ISR Number: 3166242-0Report Type:Periodic Company Report #A0063832  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Consumer	Zyban	PS		ORAL
150 MG /		Vision Blurred					
TWICE PER DAY							
/ ORAL							

Date:05/29/98ISR Number: 3166244-4Report Type:Periodic Company Report #A0063833  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Consumer	Zyban	PS		ORAL
150 MG/	ORAL						

Date:05/29/98ISR Number: 3166245-6Report Type:Periodic Company Report #A0063834  
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Consumer	Zyban	PS		ORAL
150 MG/	ORAL	Irritability					

Date:05/29/98ISR Number: 3166246-8Report Type:Periodic Company Report #A0063844  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Akinesia	Consumer	Zyban	PS		ORAL
150 MG /		Chest Discomfort					
ORAL		Depression		Diltiazem			
		Dermatitis		Hydrochloride	C		
		Hypoaesthesia		Aspirin	C		
		Movement Disorder					
		Musculoskeletal Pain					
		Musculoskeletal Stiffness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3166247-XReport Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0063845

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL	4 DAY	Dyspnoea Urticaria	Health Professional	Zyban	PS		ORAL

Date:05/29/98ISR Number: 3166248-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0063847

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ ORAL		Drug Ineffective Dysgeusia	Health Professional	Zyban	PS		ORAL

Date:05/29/98ISR Number: 3166250-XReport Type:Periodic  
 Age:36 YR Gender:Male I/FU:I

Company Report #A0063865

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Back Pain Dermatitis Fatigue Hypersensitivity Pharyngeal Oedema Swelling Urticaria	Consumer	Zyban   Omeprazole Cortisone	PS   C C		ORAL

Date:05/29/98ISR Number: 3166252-3Report Type:Periodic  
 Age:37 YR Gender:Female I/FU:I

Company Report #A0063867

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation	Consumer	Zyban	PS		ORAL
150 MG/ TWICE		Fatigue					
PERDAY/ ORAL		Feeling Drunk		Imipramine Hydrochloride	C		

Date:05/29/98ISR Number: 3166253-5Report Type:Periodic Company Report #A0063869  
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS		ORAL
150 MG/ TWICE		Pruritus					
PER DAY /		Urticaria					
ORAL				Conjugated Estrogens	C		
				Triamterene	C		
				Hydrochlorothiazide	C		

Date:05/29/98ISR Number: 3166254-7Report Type:Periodic Company Report #A0063870  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Consumer	Zyban	PS		ORAL
150 MG /		Headache					
TWICE PER DAY		Nausea					
/ ORAL		Pruritus		Oxaprozin	C		
		Vomiting		Paracetamol	C		
				Cimetidine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Salbutamol Sulphate C  
 Humibid Dm C  
 Duratuss C  
 Amitriptyline Hcl C  
 Levofloxacin C  
 Fluticasone  
 Propionate C  
 Salmeterol Xinafoate C  
 Beclomethasone  
 Dipropion. C  
 Nicotine C

Date:05/29/98ISR Number: 3166255-9Report Type:Periodic Company Report #A0063871  
 Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Diarrhoea	Consumer	Zyban	PS		ORAL
TWICE PER DAY		Headache					
/ ORAL							
				Cimetidine	C		
				Aspirin	C		
				Alka-Seltzer	C		

Date:05/29/98ISR Number: 3166274-2Report Type:Periodic Company Report #A0060065  
 Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bronchospasm	Health	Zyban	PS		ORAL
150 MG ORAL		Dermatitis	Professional	Ranitidine			
		Dyspnoea		Hydrochloride	C		
		Hypersensitivity		Amoxicillin	C		
		Pharyngeal Oedema		Entex La	C		
		Rash Macular					
		Tachypnoea					
		Urticaria					

Date:05/29/98ISR Number: 3166275-4Report Type:Periodic Company Report #A0059950  
Age:61 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / PER Initial or Prolonged DAY / ORAL	Alcohol Withdrawal Syndrome Convulsion Dry Mouth Nervousness Pancreatitis	Consumer	Zyban  Estrogen Naproxen Omeprazole Nadolol	PS  C C C C		ORAL

Date:05/29/98ISR Number: 3166276-6Report Type:Periodic Company Report #A0059853  
Age:59 YR Gender:Female I/FU:F

Outcome	PT
Other	Disorientation Dizziness Dysgeusia Headache Hypersensitivity Nausea

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sedation

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG /		Health	Zyban	PS		ORAL
TWICE PER DAY		Professional				
/ ORAL			Verapamil			
			Hydrochloride	C		
			Conjugated Estrogens	C		
			Frusemide	C		
			Famotidine	C		
			Vitamin	C		
			Paracetamol	C		

Date:05/29/98ISR Number: 3166277-8Report Type:Periodic Company Report #A0059794  
 Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Bronchospasm	Health	Zyban	PS		ORAL
150 MG			Professional				
Initial or Prolonged		Dyspnoea					
/SINGLE DOSE							
/ ORAL				Salbutamol Sulphate	C		
				Combivent	C		
				Theophylline	C		

Date:05/29/98ISR Number: 3166278-XReport Type:Periodic Company Report #A0059474  
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dermatitis	Health	Zyban	PS		ORAL
150 MG /			Professional				
TWICE PER DAY		Dyspnoea					

/ ORAL

Pruritus

Swelling  
Urticaria

Date:05/29/98ISR Number: 3167109-4Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0064432

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression	Health	Zyban	PS	Zyban	ORAL
150 MG /		Blister	Professional				
TWICE PER DAY		Cellulitis					

/ ORAL

Haemorrhage	Vitamin	C
Hypersensitivity	Herbal Medication	C
Insomnia	Mineral Supplement	C
Irritability	Hypericum	C
Oedema	Garlic	C
Pain		
Pruritus		
Skin Disorder		
Skin Lesion		
Urticaria		
Weight Increased		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3167111-2Report Type:Periodic Company Report #A0064470  
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG ORAL	Amnesia	Consumer	Zyban	PS	Zyban	ORAL
Initial or Prolonged	Convulsion Sedation		Asmacort Adrenaline	C C		

Date:05/29/98ISR Number: 3167536-5Report Type:Periodic Company Report #A0060739  
 Age:24 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL	Convulsion	Health	Zyban	PS		ORAL
Initial or Prolonged		Professional				

Date:05/29/98ISR Number: 3167538-9Report Type:Periodic Company Report #A0060648  
 Age:27 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG ORAL 17 DAY	Erythema Multiforme Pain Pruritus	Health Professional Company Representative	Zyban Levlan	PS C		ORAL

Date:05/29/98ISR Number: 3167602-4Report Type:Periodic Company Report #A0062366  
 Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE	Abdominal Pain	Consumer	Zyban	PS		ORAL
Initial or Prolonged PER DAY/ ORAL 14 DAY	Chest Pain Diarrhoea Herpes Simplex					

Nausea  
Pyrexia  
Swelling  
Urticaria  
Vomiting  
White Blood Cell Count  
Increased

Date:05/29/98ISR Number: 3167603-6Report Type:Periodic Company Report #A0062352  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Health	Zyban	PS		ORAL
150 MG/TWICE		Joint Swelling	Professional				
PER DAY/ ORAL		Oedema		Oral Contraceptive	C		
		Pruritus					
		Urticaria					

Date:05/29/98ISR Number: 3167604-8Report Type:Periodic Company Report #A0062254  
Age: Gender: I/FU:I

Outcome	PT	Report Source
Hospitalization -	Confusional State	Health
Initial or Prolonged	Liver Function Test	Professional
	Abnormal	Company

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Representative

Dose	Duration			Product	Role	Manufacturer	Route
150 MG/ORAL				Zyban	PS		ORAL

Date:05/29/98ISR Number: 3167605-XReport Type:Periodic Company Report #A0062246  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pruritus	Health	Zyban	PS		ORAL
ORAL			Professional				

Date:05/29/98ISR Number: 3167606-1Report Type:Periodic Company Report #A0062174  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dermatitis	Health	Zyban	PS		ORAL
150 MG/TWICE			Professional				
Initial or Prolonged		Urticaria					
PER DAY/ ORAL							

Date:05/29/98ISR Number: 3167607-3Report Type:Periodic Company Report #A0063416  
 Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Chest Discomfort	Health	Zyban	PS		ORAL
150 MG/TWICE			Professional				
Initial or Prolonged		Odynophagia					
PER DAY/ ORAL							
		Pruritus					
		Urticaria					

Date:05/29/98ISR Number: 3167608-5Report Type:Periodic Company Report #A0063490  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY	Nausea Tremor	Consumer	Zyban	PS		ORAL
/ ORAL						

Date:05/29/98ISR Number: 3167800-XReport Type:Periodic Company Report #A0060628  
Age:49 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150MG TWICE Initial or Prolonged PER DAY ORAL	Anorexia Chest Pain	Consumer	Zyban	PS	Zyban	ORAL
	Drug Ineffective		Isosorbide			
	Dry Mouth		Mononitrate	C		
	Dyspepsia		Metoprolol Succinate	C		
	Dyspnoea		Lansoprazole	C		
	Headache		Simvastatin	C		
	Insomnia		Aspirin	C		

Date:05/29/98ISR Number: 3167802-3Report Type:Periodic Company Report #A0060437  
Age:56 YR Gender:Male I/FU:I

Outcome	PT	Report Source
Other	Convulsion	Health Professional

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Company Representative	Product	Role	Manufacturer	Route
150MG TWICE			Zyban	PS	Zyban	ORAL
PER DAY ORAL	DAY					

Date:05/29/98ISR Number: 3167803-5Report Type:Periodic Company Report #A0060409  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG TWICE		Angioneurotic Oedema	Health	Zyban	PS	Zyban	ORAL
Initial or Prolonged PER DAY ORAL		Dysphagia	Professional				
		Face Oedema		Propranolol			
		Hypersensitivity		Hydrochloride	C		
		Oedema		Conjugated Estrogens	C		
		Tongue Oedema					
		Urticaria					

Date:05/29/98ISR Number: 3167805-9Report Type:Periodic Company Report #A0062559  
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG TWICE		Abdominal Pain Upper	Consumer	Zyban	PS	Zyban	ORAL
Initial or Prolonged PER DAY ORAL		Constipation					
Disability		Disturbance In Attention					
		Dyspnoea					
		Feeling Drunk					
		Hypersensitivity					
		Motor Dysfunction					
		Oedema Peripheral					
		Pruritus					
		Rash Papular					

Date:05/29/98ISR Number: 3167807-2Report Type:Periodic Company Report #A0062432  
Age:48 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150MG TWICE	Hypersensitivity	Health	Zyban	PS	Zyban	ORAL
Initial or Prolonged PER DAY ORAL	Oedema Peripheral	Professional				
	Pruritus Urticaria					

Date:05/29/98ISR Number: 3167809-6Report Type:Periodic Company Report #A0062403  
Age:35 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150MG ORAL	Cerebrovascular Accident	Health	Zyban	PS	Zyban	ORAL
Initial or Prolonged		Professional Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3168101-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0064023

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet-Zyban	PS		ORAL
ORAL		Pruritus					

Date:05/29/98ISR Number: 3168102-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0064024

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE		Hypersensitivity					
PER DAY/ORAL		Pharyngeal Oedema					
		Pruritus					
		Tongue Oedema					
		Urticaria					

Date:05/29/98ISR Number: 3168105-3Report Type:Periodic  
 Age:63 YR Gender:Male I/FU:I

Company Report #A0064026

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE		Rash Papular					
PER DAY/ORAL				Atenolol	C		

Date:05/29/98ISR Number: 3168107-7Report Type:Periodic  
 Age:67 YR Gender:Female I/FU:I

Company Report #A0064028

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ORAL							

Urticaria

Atorvastatin Calcium C  
Ibuprofen C  
Vitamin E C  
Oestradiol C

Date:05/29/98ISR Number: 3168109-0Report Type:Periodic Company Report #A0064030  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL							

Olanzapine C  
Antihypertensive C

Date:05/29/98ISR Number: 3168110-7Report Type:Periodic Company Report #A0064032  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disturbance In Attention	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ORAL		Dizziness					
		Dry Mouth					
		Feeling Abnormal					
		Nervousness					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/98ISR Number: 3168154-5Report Type:Periodic Company Report #A0062106  
 Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State	Health	Zyban	PS		ORAL
150 MG /		Crying	Professional				
TWICE PER DAY		Depression					
/ ORAL		Disorientation		Lansorprazole	C		
		Dyspnoea		Ipratropium Bromide	C		
		Emotional Disorder		Pain Medication	C		
		Furuncle					
		Hyperventilation					
		Nightmare					
		Palpitations					
		Tremor					

Date:05/29/98ISR Number: 3168156-9Report Type:Periodic Company Report #A0061979  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Discomfort	Health	Zyban	PS		ORAL
150 MG/ ORAL		Dermographism	Professional				
		Pruritus	Company				
		Unevaluable Event	Representative				

Date:05/29/98ISR Number: 3168158-2Report Type:Periodic Company Report #A0061938  
 Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Angioneurotic Oedema	Health	Zyban	PS		ORAL
150 MG /		Dyspnoea	Professional				
Hospitalization -							
TWICE PER DAY		Eyelid Oedema					
Initial or Prolonged							
/ ORAL							
Other		Face Oedema		Glimepiride	C		

Pharyngeal Oedema  
Speech Disorder  
Swelling

Troglitazone C  
Diltiazem  
Hydrochloride C  
Compazine C

Date:05/29/98ISR Number: 3168160-0Report Type:Periodic Company Report #A0061929  
Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Dermatitis	Consumer	Zyban	PS		ORAL
150 MG /		Dyspnoea					
TWICE PER DAY		Hypersensitivity					
/ ORAL		Pharyngeal Oedema		Diazepam	C		
		Urticaria		Ortho Cyclen	C		

Date:05/29/98ISR Number: 3188518-3Report Type:Periodic Company Report #A0063868  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Constipation	Consumer	Zyban	PS		ORAL
150 MG /		Dyspepsia					
TWICE PER DAY		Oligomenorrhoea					
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/98ISR Number: 3083794-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0055167

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hepatitis	Health	Wellbutrin Tablet	PS		ORAL
ORAL			Professional				

Date:06/01/98ISR Number: 3090040-XReport Type:Direct  
Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG PO BID		Condition Aggravated		Zyban	PS		ORAL
Initial or Prolonged		Post-Traumatic Stress Disorder Suicidal Ideation		Placebo	SS		
				Zoloft	C		
				Trazodone	C		
				Buspirone	C		
				Antabuse	C		
				Albuterol	C		
				Atrovent	C		
				Cimetidine	C		

Date:06/04/98ISR Number: 3090225-2Report Type:Expedited (15-DaCompany Report #A0062175  
Age:61 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE		Chest Pain	Health	Zyban	PS		ORAL
Initial or Prolonged PER DAY/ORAL/ TAB		Conversion Disorder	Professional				
		Coordination Abnormal					
		Crying		Digoxin	C		
		Depression		Atenolol	C		
		Fear					
		Pneumonia Mycoplasmal					
		Pulmonary Tuberculosis					

Date:06/04/98ISR Number: 3090226-4Report Type:Expedited (15-DaCompany Report #A0061539  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Nasal Congestion	Health	Zyban	PS		ORAL
150 MG/ PER		Pruritus	Professional				
DAY/ ORAL/		Serum Sickness					
TAB							

Date:06/04/98ISR Number: 3133320-1Report Type:Direct Company Report #  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Disorientation		Zyban	PS		ORAL
150MG BID PO		Face Oedema					
Other		Heart Rate Increased					
		Urticaria					

Date:06/05/98ISR Number: 3090962-XReport Type:Expedited (15-DaCompany Report #A0065877  
Age:65 YR Gender:Male I/FU:I

Outcome  
Hospitalization -  
Initial or Prolonged  
  
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Freedom Of Information (FOI) Report

Other

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150 MG/PER DAY/ORAL		Anaphylactic Shock	Consumer	Wellbutrin	PS		ORAL
	150 MG/THREE TIMES DAY/ORAL		Angioneurotic Oedema					
			Drug Ineffective Face Oedema		Wellbutrin-Controlled Release	SS		ORAL
			Paraesthesia					
			Respiratory Failure					
			Urticaria		Aspirin	C		

Date:06/05/98ISR Number: 3091334-4Report Type:Direct  
Age:35 YR Gender:Female I/FU:I

Company Report #

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG BID		Nightmare	Health Professional	Wellbutrin	PS		

Date:06/08/98ISR Number: 3091260-0Report Type:Expedited (15-Day)Company Report #A0064987  
Age:69 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG/TWICE	Initial or Prolonged PER DAY/ ORAL	Abdominal Pain	Consumer	Zyban	PS		ORAL
			Colitis					
			Diarrhoea		Herbal Dietary Supplement	C		
			Faecal Incontinence		Lorazepam	C		
			Fatigue					
			Nervousness					
			Tremor					
			Weight Decreased					
			White Blood Cell Count Increased					

Date:06/09/98ISR Number: 3091779-2Report Type:Direct  
Age:58 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Crying		Wellbutrin Sr	PS		
150 MG, BID		Depression		Zestoretic	C		
				Trental	C		
				Hytrin	C		
				Tylenol	C		
				Asa	C		

Date:06/09/98ISR Number: 3091815-3Report Type:Direct  
Age:62 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema		Zyban	PS	Glaxo	ORAL
150MG 1T PO		Urticaria					
QD				Hytrin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/11/98ISR Number: 3092899-9Report Type:Direct  
Age:42 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erythema Multiforme		Zyban	PS		
QD X 3 DAYS		Urticaria					
THEN BID							
				Tylenol Sinus	C		
				St. John'S Wart	C		

Date:06/12/98ISR Number: 3093932-0Report Type:Expedited (15-DaCompany Report #A0065116  
Age:50 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Convulsion	Health	Zyban	PS		ORAL
150 MG/ORAL		Drug Level Above Therapeutic	Professional Company Representative				

Date:06/15/98ISR Number: 3094408-7Report Type:Expedited (15-DaCompany Report #A0065665  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Consumer	Wellbutrin	PS		ORAL
75 MG/TWICE		Confusional State					
PER DAY/ORAL		Convulsion		Clonazepam	C		
		Dermatitis		Carisoprodol	C		
		Dry Mouth		Botulinum Toxin	C		
		Jaundice					
		Weight Decreased					

Date:06/15/98ISR Number: 3094825-5Report Type:Expedited (15-DaCompany Report #A0066342  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Convulsion	Health	Wellbutrin	PS		ORAL
Initial or Prolonged ORAL		Drug Interaction	Professional	Fluvoxamine Maleate	SS		ORAL
		Hypertension Intentional Misuse Tachycardia	Company Representative				

Date:06/17/98ISR Number: 3095280-1Report Type:Expedited (15-DaCompany Report #A0066297  
Age:14 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ORAL		Convulsion	Health	Wellbutrin	PS		ORAL
Initial or Prolonged ORAL		Depressed Level Of Consciousness	Professional	Diphenhydramine Hcl	SS		ORAL

Date:06/18/98ISR Number: 3095740-3Report Type:Direct Company Report #  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 BID		Grand Mal Convulsion		Wellbutrin	PS		



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Freedom Of Information (FOI) Report

Date:06/22/98ISR Number: 3097062-3Report Type:Expedited (15-DaCompany Report #A0063323

Age:67 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation	Health	Wellbutrin	PS		ORAL
150 MG PER Other DAY ORAL		Anxiety	Professional				
		Completed Suicide		Semisodium Valproate	C		
		Condition Aggravated		Lansoprazole	C		
		Diarrhoea		Ascorbic Acid	C		
		Irritability		Vitamin B Complex	C		
		Mood Altered		Tranylcypromine			
		Paranoia		Sulphate	C		

Date:06/22/98ISR Number: 3097400-1Report Type:Direct

Age:45 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cerebrovascular Accident		Zyban	PS	Glaxo-Wellcome	

Date:06/22/98ISR Number: 3097433-5Report Type:Direct

Age:42 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG 2 X'S Initial or Prolonged DAILY		Asthenia		Zyban	PS	Glaxo Wellcome	ORAL
		Chest Discomfort					
		Chest Pain					
		Dizziness					
		Muscle Tightness					
		Vomiting					

Date:06/23/98ISR Number: 3097760-1Report Type:Expedited (15-DaCompany Report #8-98076-001K

Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aortic Valve Incompetence	Health	Redux	PS		ORAL
15 MG TWICE		Basal Cell Carcinoma	Professional				
DAILY ORAL		Blister		Wellbutrin	SS		
75 TO 150 MG		Blood Pressure Increased					
TWICE DAILY		Bronchitis Acute		Klonpin	C		
		Bronchospasm		Multivitamin	C		
		Cough		Vitamin C	C		
		Crying		Vitamin E	C		
		Diarrhoea		Wellbutrin	C		
		Eczema					
		Erythema Nodosum					
		Fluid Retention					
		Nausea					
		Pyrexia					
		Sedation					

Date:06/24/98ISR Number: 3098682-2Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Irritability		Zyban	PS		ORAL
150 MG QD X 3		Mental Disorder					
D PO		Suicidal Ideation		Diabeta	C		

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Freedom Of Information (FOI) Report

Asa C  
 Menocor C  
 Lisinopril C

Date:06/24/98ISR Number: 3098713-XReport Type:Direct  
 Age:38 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1T PO BID Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage		Dermatitis		Zyban	PS	Glaxowellcome	ORAL
		Lymphadenopathy Pain Serum Sickness					

Date:06/29/98ISR Number: 3099734-3Report Type:Expedited (15-DaCompany Report #A0066903  
 Age:14 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening SINGLE DOSE / Hospitalization - ORAL Initial or Prolonged Disability		Convulsion	Health	Wellbutrin	PS		ORAL
		Intentional Misuse	Professional				
		Respiratory Failure	Company Representative				

Date:07/01/98ISR Number: 3100470-5Report Type:Expedited (15-DaCompany Report #A0066987  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG/ TWICE PER DAY/ ORAL		Csf Pressure Increased	Consumer	Zyban	PS		ORAL
		Hepatic Function Abnormal					
		Hepatitis Viral Optic Nerve Disorder Papilloedema					

Platelet Count Decreased  
Pyrexia  
Visual Field Defect  
Vomiting

Date:07/01/98ISR Number: 3100514-0Report Type:Expedited (15-DaCompany Report #A0065877  
Age:65 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG/PER	Anaphylactic Reaction	Health	Wellbutrin	PS		ORAL
Initial or Prolonged DAY/ORAL; Other 150MG/THREE	Angioneurotic Oedema	Professional				
TIMES PER DAY/ORAL	Face Oedema					
	Hypersensitivity					
	Paraesthesia					
	Respiratory Failure		Aspirin	C		
	Syncope Vasovagal		Ethanol	C		
	Urticaria		Amoxicillin	C		
			Ciprofloxacin Hcl	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/02/98ISR Number: 3101743-2Report Type:Expedited (15-DaCompany Report #A0066676  
Age:15 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Amnesia	Consumer	Wellbutrin	PS		ORAL
ORAL							
Other		Coordination Abnormal Disturbance In Attention Fall Joint Contracture Migraine Nervous System Disorder Peripheral Nerve Injury Petit Mal Epilepsy Wrist Fracture	Company Representative				

Date:07/06/98ISR Number: 3102317-XReport Type:Expedited (15-DaCompany Report #A0065505  
Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / PER Initial or Prolonged DAY / ORAL		Condition Aggravated	Consumer	Wellbutrin	PS		ORAL
50 MG / TWICE PER DAY / ORAL /TABLET		Mood Swings					
		Platelet Count Decreased		Lamictal	SS		ORAL
		Premenstrual Syndrome					
		Pyrexia					
ORAL/TABLET		White Blood Cell Count		Quetiapine Fumarate	SS		ORAL
ORAL/TABLET		Decreased		Carbamazepine	SS		ORAL
ORAL/TABLET				Paroxetine Hydrochloride	C		
				Lorazepam	C		
				Methysergide Maleate	C		

Date:07/06/98ISR Number: 3103894-5Report Type:Direct  
Age:40 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis		Wellbutrin	PS		ORAL
150 SR PO							

Date:07/07/98ISR Number: 3103927-6Report Type:Direct Company Report #  
 Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin	PS		
75-150 MG Q							
		Tremor					
DAY				Zoloft	C		
				Melatonin	C		

Date:07/07/98ISR Number: 3104072-6Report Type:Direct Company Report #  
 Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Face Oedema		Olanzapine	PS		
HS							
Intervention to		Rash Maculo-Papular		Bupropin	SS		
HS							
Prevent Permanent							
Impairment/Damage							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/07/98ISR Number: 3104149-5Report Type:Direct  
 Age:37 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG PO		Blepharospasm	Health	Zyban	PS	Glaxo Wellcome	ORAL
DAILY	3 DAY	Palpitations	Professional				
				Vitamins	C		

Date:07/08/98ISR Number: 3103114-1Report Type:Expedited (15-DaCompany Report #MPI 97612  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 15 OR 30 MG		Aortic Valve Incompetence	Health	Ionamin	PS		ORAL
DAILY-ORAL		Mitral Valve Incompetence	Professional				
VARIES		Palpitations					
20 MG TID		Tricuspid Valve Incompetence		Pondimin	SS		ORAL
ORAL				Redux	SS		ORAL
15 MG, BID				Fastin	SS		ORAL
ORAL				Wellbutrin	SS		ORAL
100 MG BID							
ORAL 75 MG,							
2, BID ORAL							

Date:07/08/98ISR Number: 3103407-8Report Type:Expedited (15-DaCompany Report #A0066944  
 Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG/TWICE		Chest Discomfort	Consumer	Wellbutrin	PS		ORAL
Initial or Prolonged PER DAY/ORAL		Chest Pain		Wellbutrin	SS		ORAL
150 MG	ORAL			Clonazepam	C		
				Aspirin	C		
				Sertraline			
				Hydrochloride	C		

Date:07/14/98ISR Number: 3104056-8Report Type:Direct Company Report #  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 PO QD X 7 DAYS THEN BID.		Agitation Anxiety Disorder Hostility		Zyban	PS		ORAL
		Screaming Speech Disorder Tearfulness Thinking Abnormal		Norvasc	C		
				Lozol	C		

Date:07/16/98ISR Number: 3105828-6Report Type:Expedited (15-DaCompany Report #A0067512  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG/ TWICE PER DAY/ ORAL		Death	Health Professional	Zyban	PS		ORAL
				Indinavir Sulfate	C		
				Lamivudine	C		
				Protease Inhibitor	C		



Freedom Of Information (FOI) Report

Sedative C  
Lorazepam C

Date:07/17/98ISR Number: 3106332-1Report Type:Expedited (15-DaCompany Report #9819711  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150.00 MG Initial or Prolonged TOTAL; DAILY; ORAL	Angina Pectoris Chest Discomfort Chest Pain	Consumer	Zoloft	PS		ORAL
150.00 MG TOTAL; DAILY; ORAL	Coronary Artery Occlusion Drug Interaction Erectile Dysfunction		Wellbutrin	SS		ORAL
	Eye Irritation Hyperhidrosis Keratoconjunctivitis Sicca Libido Decreased Mood Swings Pollakiuria Speech Disorder		Clonazepam	C		

Date:07/17/98ISR Number: 3188398-6Report Type:Periodic Company Report #A0063064  
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Arrhythmia	Health Professional Company Representative	Wellbutrin Tablet- Controlled Release	PS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Anaphylactic Reaction Dermatitis Drug Interaction	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ TWICE		Kidney Infection					
PER DAY/ ORAL		Pain Pyrexia Urticaria		Ciprofloxacin (Formulation Unknown)	SS		ORAL
ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Extrapyramidal Disorder Fall Parkinson'S Disease	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ TWICE			Representative				
PER DAY/ ORAL				Terazodin Hydrochloride Imipramine Famotidine Steroid ....	C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

... C  
 ..... C

Date:07/17/98ISR Number: 3188409-8Report Type:Periodic Company Report #A0064841  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abnormal Behaviour Agitation Cerebral Artery Embolism	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ PER DAY/ ORAL		Speech Disorder	Representative				
				Clozapine Semisodium Valproate Paroxetine Hydrochloride	C C C C		

Date:07/17/98ISR Number: 3188413-XReport Type:Periodic Company Report #A0064907  
 Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other		Deafness	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ ORAL			Representative	Trazodone	C		

Date:07/17/98ISR Number: 3188417-7Report Type:Periodic Company Report #A0065150  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other		Anaphylactic Reaction Blood Pressure Decreased Dermatitis	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL		Dyspnoea					

Pharyngeal Oedema  
Pruritus

Fluoxetine  
Hydrochloride C

Date:07/17/98ISR Number: 3188421-9Report Type:Periodic  
Age:22 YR Gender:Female I/FU:I

Company Report #A0065505

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Drug Interaction Mood Altered Platelet Count Decreased	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ PER DAY/ ORAL	Pyrexia					
50 MG/TWICE PER DAY/ ORAL	White Blood Cell Count Decreased		Lamictal Tablet	SS		ORAL
ORAL			Quetiapine Fumarate Tablet	SS		ORAL
ORAL			Carbamazepine Tablet	SS		ORAL
			Paroxetine Hydrochloride	C		
			Lorazepam	C		
			Methysergide Maleate	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/17/98ISR Number: 3188424-4Report Type:Periodic  
 Age:44 YR Gender:Female I/FU:I

Company Report #A0065573

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Aggression	Consumer	Wellbutrin			
		Crying		Tablet-Controlled			
		Delusion		Release	PS		ORAL
150 MG/TWICE							
PER DAY/ ORAL		Depression					
		Drug Ineffective		Zyban Tablet-Zyban	SS		ORAL
150 MG/ ORAL		Hallucination		Prednisone	C		
		Mood Altered		Salbutamol Sulphate	C		
		Speech Disorder		Beclomethasone			
		Tremor		Dipropion.	C		
				Ipratropium Bromide	C		

Date:07/17/98ISR Number: 3188428-1Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #A0065668

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Anxiety	Consumer	Wellbutrin			
Initial or Prolonged		Confusional State		Tablet-Controlled			
		Insomnia		Release	PS		ORAL
150 MG/ ORAL							

Date:07/17/98ISR Number: 3188431-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0065767

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Delirium	Health	Wellbutrin			
Initial or Prolonged			Professional	Tablet-Controlled			
			Company	Release	PS		ORAL
ORAL			Representative				

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dermatitis Disorientation Dizziness	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL		Haematemesis  Hallucination Pyrexia Stevens-Johnson Syndrome Urticaria	Representative				

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Asthenia Tremor	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
1 TABLET/ TWICE PER DAY/ ORAL			Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/17/98ISR Number: 3188435-9Report Type:Periodic  
Age:22 YR Gender:Female I/FU:I

Company Report #A0065875

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Disturbance In Attention Joint Dislocation	Health Professional	Wellbutrin Tablet-Controlled Release			ORAL
400 MG/ PER DAY/ ORAL	1	YR		Diuretic	C		

Date:07/17/98ISR Number: 3188436-0Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0067121

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Amnesia Convulsion	Consumer	Wellbutrin Tablet-Controlled Release			ORAL
150 MG/ TWICE PER DAY/ ORAL				Bromfenac Sodium Tramadol Hydrochloride Mirtazapine	C C C		

Date:07/17/98ISR Number: 3188438-4Report Type:Periodic  
Age:10 YR Gender:Male I/FU:I

Company Report #A0067233

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Condition Aggravated Convulsion Overdose	Health Professional Company	Wellbutrin Tablet-Controlled Release			ORAL
1500 MG/ ORAL			Representative				

Date:07/17/98ISR Number: 3188440-2Report Type:Periodic Company Report #A0067257  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Dermatitis	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:07/17/98ISR Number: 3188442-6Report Type:Periodic Company Report #A0067324  
Age:28 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Convulsion	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL			Asthma Medication Theophylline Haloperidol	C C C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/17/98ISR Number: 3188445-1Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #A0067426

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Urticaria	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL			Representative				

Date:07/20/98ISR Number: 3109633-6Report Type:Direct  
Age:48 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PO BID		Abdominal Pain Agitation Crying Feeling Jittery Paranoia		Zyban	PS		ORAL

Date:07/22/98ISR Number: 3108658-4Report Type:Expedited (15-DaCompany Report #MPI-981429  
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 15 MG DAILY Life-Threatening ORAL		Dysuria	Health	Ionamin	PS		ORAL
20-60 MG DAILY ORAL		Fall Insomnia	Professional	Pondimin	SS		ORAL
400 MG DAILY ORAL MANY YEARS		Ventricular Hypertrophy		Wellbutrin	SS		ORAL

Loestrin C  
Naprosyn C  
Ultram C

Date:07/22/98ISR Number: 3108831-5Report Type:Direct  
Age:65 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Life-Threatening	Anxiety		Zyban	PS		
TBCR 150 MG						
Hospitalization -	Chest Discomfort					
AS DIRECTED						
Initial or Prolonged	Chest Pain					
ON LABEL						
	Hemiparesis					
	Sensation Of Heaviness					
	Vision Blurred					

Date:07/22/98ISR Number: 3109036-4Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
3 WK	Dysphagia		Zyban	PS		
	Face Oedema					
	Periorbital Oedema					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/22/98ISR Number: 3109046-7Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS		
150 MG BID		Loss Of Consciousness		Effexor	C		
				Depakate	C		
				Klonopin	C		
				Zyprexa	C		

Date:07/23/98ISR Number: 3107804-6Report Type:Expedited (15-DaCompany Report #A0067257  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Auricular Swelling	Health	Wellbutrin	PS		ORAL
Initial or Prolonged ORAL		Blood Pressure Increased	Professional	Dexamphetamine	SS		ORAL
ORAL		Blood Thyroid Stimulating	Company	Olanzapine	SS		ORAL
		Hormone Decreased	Representative				
		Hyperglycaemia					
		Hypersensitivity					
		Leukocytosis					
		Pruritus					
		Rash Maculo-Papular					
		Tachycardia					

Date:07/27/98ISR Number: 3110819-5Report Type:Expedited (15-DaCompany Report #A0067945  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / TWICE PER DAY		Abdominal Pain Lower	Health	Zyban	PS		ORAL
Initial or Prolonged TWICE PER DAY		Asthenia	Professional				
/ ORAL		Bronchospasm	Company				

INTRAMUSCULAR	UNK /	Difficulty In Walking	Representative	Ceftriaxone	SS	
DOSE /		SINGLE				
		Dizziness				
		Muscle Spasms				
INTRAMUSCULAR		Neck Pain		Doxycycline	SS	ORAL
100 MG /		Pyrexia				
TWICE PER DAY		Serum Sickness				
/ ORAL		White Blood Cell Count				
		Increased				

Date:07/27/98ISR Number: 3110822-5Report Type:Expedited (15-DaCompany Report #A0068002  
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bronchitis	Health	Zyban	PS		ORAL
150 MG /		Bronchospasm	Professional				
TWICE PER DAY		Cardiac Failure					
/ ORAL		Congestive		Nicotinic Acid	C		
		Cough		Aspirin	C		
		Dyspnoea					
		Haemoptysis					
		Rales					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/27/98ISR Number: 3157553-3Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #M0157-98

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alcoholic Hangover	Health	Remeron	PS		
15MG/DAY PO							
		Drug Interaction	Professional	Wellbutrin Sr	SS		
150MG BID				Claritin	C		

Date:07/28/98ISR Number: 3112923-4Report Type:Direct  
Age:43 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Hepatitis		Wellbutrin Sr	PS		ORAL
150 MG PO BID							
Initial or Prolonged		Myalgia					
		Pyrexia					

Date:07/29/98ISR Number: 3123280-1Report Type:Direct  
Age:40 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased		Bupropion	PS		ORAL
75 MG PO Q AM							
		Chest Discomfort		Fluoxetine	SS		ORAL
20 MG PO Q AM							
		Dysphagia					
		Dyspnoea					
		Tachycardia					

Date:07/30/98ISR Number: 3111481-8Report Type:Direct  
Age:71 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Bupropion Sa	PS		ORAL
150 MG QD,							

ORAL

Pentoxifylline	C
Albuterol	C
Aspirin	C
Buspirone Hcl	C
Zolpidem	C
Nifedipime (Adalat Cc)	C
Ferrous Sulfate	C
Docusate	C
Levothyroxine	C
Lovastatin	C

Date:07/31/98ISR Number: 3111244-3Report Type:Expedited (15-DaCompany Report #A0067874

Age:78 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/IN THE Initial or Prolonged MORNING/ORAL	Hepatic Failure  Renal Failure	Consumer	Zyban	PS		ORAL

Paracetamol	C
Lorazepam	C
Aspirin	C
Cold Medication	C
Propranolol	
Hydrochloride	C
Methyldopa	C
Cimetidine	C

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Freedom Of Information (FOI) Report

Date:07/31/98ISR Number: 3111245-5Report Type:Expedited (15-DaCompany Report #A0068184

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Health	Zyban	PS		ORAL
150		Diabetic Retinopathy	Professional				
MG/UNK/ORAL			Other	Raloxifene Hydrochloride	SS		ORAL
60 MG/PER							
DAY/ORAL							

Date:07/31/98ISR Number: 3111780-XReport Type:Direct

Company Report #

Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Bupropion	PS		ORAL
75 MG, TWO		Drug Ineffective					
BID, ORAL				Valproic Acid	C		
				Propranolol Hcl	C		

Date:07/31/98ISR Number: 3200049-0Report Type:Periodic

Company Report #8-97351-028B

Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea	Consumer	Pondimin	PS		ORAL
20 MG THREE		Dizziness					
TIMES DAILY		Drug Withdrawal Syndrome					
ORAL		Dyspnoea		Phentermine	SS		
ABOUT 4 YEARS 4 YR				Serzone	SS		
				Wellbutrin	SS		
				Premarin	C		

Date:08/03/98ISR Number: 3112548-0Report Type:Expedited (15-DaCompany Report #A0068320  
Age:36 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL	Dermatitis  Dyspnoea  Feeling Abnormal Laboratory Test Abnormal Pyrexia Urticaria	Consumer	Zyban	PS		ORAL

Date:08/03/98ISR Number: 3112551-0Report Type:Expedited (15-DaCompany Report #A0066676  
Age:14 YR Gender:Female I/FU:F

Outcome	PT
Disability	Amnesia
Other	Bone Disorder Coordination Abnormal Disturbance In Attention Eye Movement Disorder Fall Impaired Healing Migraine Movement Disorder Nervous System Disorder Pain In Extremity Peripheral Nerve Injury

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG/ TWICE PER DAY/ORAL		Petit Mal Epilepsy Photophobia Staring Vomiting Wrist Deformity Wrist Fracture	Health Professional Company Representative	Wellbutrin Trazodone	PS C		ORAL

Date:08/03/98ISR Number: 3201442-2Report Type:Periodic Company Report #8-98008-037B  
Age:29 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 20 MG THREE TIMED DAILY ORAL		3 WKN	Asthenia Drug Withdrawal Syndrome	Health Professional	Pondimin Ionamin Wellbutrin	PS SS SS		ORAL
ORAL					Ionamin Valium	C C		

Date:08/04/98ISR Number: 3112697-7Report Type:Direct Company Report #  
Age:16 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG SQ PO BID			Anorexia Decreased Appetite		Wellbutrin	PS		ORAL

Date:08/04/98ISR Number: 3113209-4Report Type:Direct  
Age:9 YR Gender:Male I/FU:I

Company Report #

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG QD		Abnormal Behaviour Obsessive-Compulsive Disorder		Wellbutrin	PS		

Date:08/04/98ISR Number: 3113216-1Report Type:Direct  
Age:42 YR Gender:Female I/FU:I

Company Report #

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 100MG, TID, Intervention to BY MOUTH Prevent Permanent Impairment/Damage		Mouth Ulceration		Bupropion	PS		
				Insulin Syringe Easy Glucose Reagen Strip Insulin Nph Human Lisinopril Acetaminophen	C C C C C		

Date:08/05/98ISR Number: 3113392-0Report Type:Expedited (15-DaCompany Report #8-98016-002S  
Age:43 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 20-60 MG		Insomnia Ventricular Hypertrophy	Health Professional	Pondimin	PS		ORAL
DAILY; ORAL 400 MG DAILY;				Wellbutrin	SS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL  
 15 MG DIALY; Ionamin SS ORAL

ORAL  
 Naproxyn Ec C  
 Ultram C  
 Wellbutrin C  
 Ionamin C  
 Loestrin C

Date:08/05/98ISR Number: 3113632-8Report Type:Direct Company Report #  
 Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Hypotension		Zyban	PS		
150 MG BID	1 MON						
Intervention to		Orthostatic Hypotension		Hctz	C		
Prevent Permanent		Renal Impairment		Atenolol	C		
Impairment/Damage				Zestril	C		
				Zantac	C		
				Zyban	C		

Date:08/05/98ISR Number: 3202196-6Report Type:Periodic Company Report #97-12-0778  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Health	Intron A (Interferon			
Other		Drug Interaction	Professional	Alfa-2b Recombinant)			
		Hypertension		Injectable	PS		
3 MU TIW							
		Hypertriglyceridaemia		Bupropion			
		Influenza Like Illness		Hydrochloride			
				Tablets	SS		ORAL

Date:08/06/98ISR Number: 3113778-4Report Type:Direct Company Report #  
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Feeling Jittery		Wellbutrin Sr	PS		ORAL
150 MG PO		Nervousness					
DAILY				Paxil	C		

Date:08/10/98ISR Number: 3115264-4Report Type:Expedited (15-DaCompany Report #A0068687  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Arthralgia	Health	Wellbutrin	PS		ORAL
ORAL							
Initial or Prolonged		Dermatitis Exfoliative Pyrexia Urticaria	Professional Company Representative				

Date:08/12/98ISR Number: 3116962-9Report Type:Direct Company Report #  
Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia		Wellbutrin	PS		
100 MG BID		Chills Myalgia Pyrexia Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/12/98ISR Number: 3117261-1Report Type:Direct  
Age:35 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG X 2		Drug Withdrawal Syndrome					

Date:08/13/98ISR Number: 3116733-3Report Type:Direct  
Age:24 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Fall		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged		Grand Mal Convulsion Laceration					

Date:08/14/98ISR Number: 3117355-0Report Type:Expedited (15-DaCompany Report #A0068408  
Age:26 YR Gender:Female I/FU:I

Company Report #A0068408

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Platelet Count Decreased	Health	Wellbutrin	PS		ORAL
150 MG /			Professional				
TWICE PER							
DAY/ ORAL							

Lithium Salt C  
Naproxen C

Date:08/14/98ISR Number: 3117417-8Report Type:Expedited (15-DaCompany Report #A0068968  
Age:61 YR Gender:Female I/FU:I

Company Report #A0068968

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Chest Pain	Consumer	Zyban	PS		ORAL
150 MG /		Rash Pruritic					
Initial or Prolonged							
TWICE PER DAY							

/ ORAL

Aspirin C  
Morphine C

Date:08/14/98ISR Number: 3117449-XReport Type:Expedited (15-DaCompany Report #A0065568  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Health	Zyban	PS		ORAL
150 MG /		Connective Tissue	Professional				
TWICE PER DAY		Disorder					
/ ORAL		Genital Ulceration					
		Mouth Ulceration					
		Skin Disorder					

Date:08/18/98ISR Number: 3118512-XReport Type:Direct Company Report #  
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Urticaria		Welbutrin	PS		
Intervention to				Lasix	C		
Prevent Permanent				Serzone	C		
Impairment/Damage				Digoxin	C		
				Ambien	C		
				Ativan	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/20/98ISR Number: 3119552-7Report Type:Expedited (15-DaCompany Report #A0069321

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin	PS		ORAL
Other		Anxiety					
UNK/ UNK/			Professional				
ORAL		Coordination Abnormal					
		Movement Disorder	Company	Carisoprodol	SS		ORAL
UNK/ UNK/			Representative				
ORAL							

Date:08/20/98ISR Number: 3119555-2Report Type:Expedited (15-DaCompany Report #A0069088

Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS		ORAL
Other		Chest Pain					
150 MG/TWICE							
PER DAY/ORAL		Choking					
		Convulsion		Simvastatin	C		
		Thirst					
		Tremor					

Date:08/20/98ISR Number: 3119556-4Report Type:Expedited (15-DaCompany Report #A0067422

Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin	PS		ORAL
Other		Cognitive Deterioration					
Hospitalization -			Professional				
150 MG/TWICE		Hallucination					
Initial or Prolonged							
PER DAY/ORAL		Muscle Rigidity		Alpraxolam	C		
Other		Parkinson'S Disease		Estrogen	C		
		Tremor		Famotidine	C		
				Vitamin D	C		
				Percocet	C		
				Calcium Carbonate	C		

Date:08/21/98ISR Number: 3120797-0Report Type:Expedited (15-DaCompany Report #A0069307  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL		Joint Swelling	Health	Zyban	PS		ORAL
		Rash Erythematous	Professional				
		Serum Sickness	Company Representative				

Date:08/24/98ISR Number: 3120918-XReport Type:Expedited (15-DaCompany Report #A0069385  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Abortion Spontaneous	Consumer	Zyban	PS		ORAL

Date:08/24/98ISR Number: 3121056-2Report Type:Expedited (15-DaCompany Report #A0067746  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG; TWICE PER DAY; ORAL		Brain Neoplasm	Health	Zyban	PS		ORAL
		Grand Mal Convulsion	Professional Company Representative				



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/24/98ISR Number: 3121161-0Report Type:Direct  
Age:41 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG PO BID	Abdominal Pain		Wellbutrin Sr	PS		ORAL
				Paxil	C		

Date:08/25/98ISR Number: 3120747-7Report Type:Expedited (15-DaCompany Report #A0066676  
Age:14 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG / TWICE		Amnesia	Health	Wellbutrin	PS		ORAL
Initial or Prolonged PER DAY / Disability		Bone Disorder	Professional				
ORAL Other		Coordination Abnormal	Company				
		Disturbance In Attention Eyelid Function Disorder Fall Impaired Healing Migraine Movement Disorder Nervous System Disorder Peripheral Nerve Injury Petit Mal Epilepsy Photophobia Staring Vomiting Wrist Deformity Wrist Fracture	Representative	Trazodone	C		

Date:08/25/98ISR Number: 3122492-0Report Type:Direct  
Age:37 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150 MG BID PO		Agitation		Wellbutrin	PS		ORAL
Hospitalization -		Aspiration					

Initial or Prolonged      Blood Pressure Systolic  
Increased  
Confusional State  
Hallucination  
Hyperreflexia  
Overdose  
Respiratory Rate  
Increased  
Tachycardia

Date:08/27/98ISR Number: 3122730-4Report Type:Expedited (15-DaCompany Report #A0068687

Age:                      Gender:Male                      I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose                      Duration Hospitalization - 150 MG ORAL	Arthralgia	Health	Wellbutrin	PS		ORAL
Initial or Prolonged	Burning Sensation Dermatitis Exfoliative Malaise Pyrexia Rash Erythematous Rash Pruritic	Professional Company Representative				

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Freedom Of Information (FOI) Report

Date:08/27/98ISR Number: 3122733-XReport Type:Expedited (15-DaCompany Report #A0069140

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion Drug Level Above Therapeutic	Literature	Wellbutrin Unknown Ssri (Formulation Unknown)	PS  SS		

Date:08/27/98ISR Number: 3122735-3Report Type:Expedited (15-DaCompany Report #A0069142

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion Drug Level Above Therapeutic	Literature	Wellbutrin Unknown Ssri (Formulation Unknown)	PS  SS		

Date:08/27/98ISR Number: 3122737-7Report Type:Expedited (15-DaCompany Report #A0069143

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion Drug Level Above Therapeutic	Literature	Wellbutrin Unknown Ssri (Formulation Unknown)	PS  SS		

Date:08/27/98ISR Number: 3123040-1Report Type:Expedited (15-DaCompany Report #A0069623

Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL		Dehydration  Dizziness  Heart Rate Increased Insomnia	Consumer	Wellbutrin	PS		ORAL

Date:08/27/98ISR Number: 3123050-4Report Type:Expedited (15-DaCompany Report #A0069631

Age:55 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - TWICE PER DAY	Diarrhoea	Consumer	Zyban	PS		ORAL
Initial or Prolonged ORAL	Nausea					
	Pancreatitis		Lisinopril	C		
			Chlorpropamide	C		
			Fluoxetine			
			Hydrochloride	C		

Date:08/27/98ISR Number: 3123067-XReport Type:Direct

Company Report #

Age:39 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150 MG BID PO 1 WK	Blood Creatine		Zyban	PS		ORAL
Initial or Prolonged	Phosphokinase Increased					
	Pain In Extremity					
	Rhabdomyolysis					

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Freedom Of Information (FOI) Report

Date:08/28/98ISR Number: 3123552-0Report Type:Expedited (15-DaCompany Report #A0068941  
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG		Conversion Disorder		Zyban	PS		ORAL
Initial or Prolonged /UNK/ORAL		Feeling Abnormal					
		Hallucination Tremor					

Date:08/31/98ISR Number: 3123324-7Report Type:Expedited (15-DaCompany Report #A0069746  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Health Professional Company Representative	Wellbutrin Diphenhydramine Hcl	PS SS		ORAL

Date:08/31/98ISR Number: 3124018-4Report Type:Direct Company Report #  
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Health	Zyban	PS		ORAL
150 BID PO	2 MON						
Other TRANSDERMAL	14 MG		Professional	Habitrol	SS		
TRANSDERMAL	2 MON						

Date:09/01/98ISR Number: 3124703-4Report Type:Expedited (15-DaCompany Report #A0069000  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150 MG/TWICE		Apnoea	Health	Zyban	PS		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nabumetone C  
 Famotidine C  
 Vicodin C

Date:09/01/98ISR Number: 3204199-4Report Type:Periodic Company Report #A0064923  
 Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL		Delirium	Health	Zyban Tablet - Zyban	PS		ORAL
		Fall	Professional				
		Laceration		Iron Salt	C		
		Vision Blurred		Potassium Salt	C		
				Aspirin	C		
				Nephrocaps	C		

Date:09/01/98ISR Number: 3204201-XReport Type:Periodic Company Report #A0064852  
 Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ ORAL Initial or Prolonged		Myocardial Infarction	Consumer	Zyban Tablet-Zyban	PS		ORAL
		Thrombosis		Estratest	C		
				Multivitamin	C		

Date:09/01/98ISR Number: 3204203-3Report Type:Periodic Company Report #A0064613  
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG ORAL		Choking Sensation	Health	Zyban Tablet - Zyban	PS		ORAL
		Hypersensitivity	Professional				
		Pruritus					
		Urticaria					

Date:09/01/98ISR Number: 3204206-9Report Type:Periodic  
Age:38 YR Gender:Female I/FU:F

Company Report #A0064432

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG /TWICE PER DAY/ ORAL	Aggression Blister	Health Professional	Zyban Tablet - Zyban	PS		ORAL
	Cellulitis		Multivitamin	C		
	Haemorrhage		Herbal Medication	C		
	Hypersensitivity		Mineral Supplements	C		
	Insomnia		Hypericum	C		
	Irritability		Garlic	C		
	Oedema					
	Pain					
	Pruritus					
	Skin Disorder					
	Urticaria					
	Weight Increased					

Date:09/01/98ISR Number: 3204209-4Report Type:Periodic  
Age:39 YR Gender:Female I/FU:F

Company Report #A0064347

Outcome  
Hospitalization -  
Initial or Prolonged  
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Freedom Of Information (FOI) Report

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Angioneurotic Oedema Arthralgia	Health Professional	Zyban Tablets - Zyban	PS		ORAL
PER DAY/ ORAL		Serum Sickness	Company				
		Urticaria	Representative	Ibuprofen Cetirizine Hydrochloride	C C		

Date:09/01/98ISR Number: 3204212-4Report Type:Periodic Company Report #A0064061  
Age:64 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Tinnitus	Health Professional	Zyban Tablets - Zyban	PS		ORAL
150 MG /							
TWICE PER							
DAY/ ORAL							

Date:09/01/98ISR Number: 3204214-8Report Type:Periodic Company Report #A0064018  
Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Amnesia	Consumer	Zyban Tablets- Zyban	PS		ORAL
150 MG/ TWICE		Balance Disorder					
Initial or Prolonged		Confusional State					
PER DAY/ ORAL		Dissociation					
		Dizziness					
		Ecchymosis					
		Grand Mal Convulsion					
		Haemorrhage					
		Laceration					
		Nystagmus					

Date:09/01/98ISR Number: 3204218-5Report Type:Periodic  
Age:44 YR Gender:Female I/FU:F

Company Report #A0063779

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Agitation	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /TWICE						
Initial or Prolonged	Memory Impairment	Professional				
PER DAY/ ORAL	Tremor					

Date:09/01/98ISR Number: 3204219-7Report Type:Periodic  
Age:31 YR Gender:Male I/FU:F

Company Report #A0063416

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Chest Discomfort	Health	Zyban Tablets -			
Initial or Prolonged	Odynophagia	Professional	Zyban	PS		ORAL
150 MG/ TWICE						
	Pruritus					
PER DAY/ ORAL	Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/01/98ISR Number: 3204222-7Report Type:Periodic  
Age:44 YR Gender:Female I/FU:F

Company Report #A0063490

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ TWICE		Agitation	Health	Zyban Tablets- Zyban	PS		ORAL
Initial or Prolonged PER DAY/ ORAL		Conversion Disorder	Professional				
Other		Joint Stiffness Mental Disorder Movement Disorder Muscle Twitching Nausea Pain Tremor					

Date:09/01/98ISR Number: 3204223-9Report Type:Periodic  
Age:53 YR Gender:Female I/FU:F

Company Report #A0063213

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG/ PER		Dyspnoea	Health	Zyban Tablet - Zyban	PS		ORAL
DAY/ ORAL		Hypersensitivity	Professional				
		Insomnia		Etodolac	C		
		Pruritus		Oestradiol	C		
		Swelling		Claritin-D	C		
		Urticaria					

Date:09/01/98ISR Number: 3204226-4Report Type:Periodic  
Age:60 YR Gender:Female I/FU:F

Company Report #A0063157

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG ORAL		Cough	Health	Zyban Tablet - Zyban	PS		ORAL
		Hypoxia	Professional	Fluticasone			
		Leukocytosis		Propionate	C		
		Pneumonitis		Ipratropium Bromide	C		

Date:09/01/98ISR Number: 3204230-6Report Type:Periodic Company Report #A0062940  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Back Pain	Health	Zyban Tablet-Zyban	PS		ORAL
150 MG ORAL		Convulsion	Professional	Trazodone	C		

Date:09/01/98ISR Number: 3204232-XReport Type:Periodic Company Report #A0062673  
Age:27 YR Gender:Male I/FU:F

Outcome	PT
Other	Anaphylactic Reaction
	Angioneurotic Oedema
	Chills
	Condition Aggravated
	Face Oedema
	Flushing
	Heart Rate Increased
	Hypersensitivity
	Hypotension
	Pruritus
	Rash Erythematous
	Swelling
	Tongue Oedema



Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL	Dermatitis Dysphagia Dyspnoea Insomnia Pruritus Speech Disorder Swelling Urticaria	Consumer	Zyban Tablet- Zyban	PS	ORAL
			Glipizide	C	
			Theophylline	C	
			Dyazide	C	
			Salbutamol Sulphate	C	

Date:09/01/98ISR Number: 3204242-2Report Type:Periodic Company Report #A0057814  
Age:59 YR Gender:Female I/FU:F

Outcome	PT
Disability	Agitation
	Amnesia
	Arthralgia
	Bronchitis
	Cough
	Depression
	Difficulty In Walking
	Dry Mouth
	Dyspraxia
	Grand Mal Convulsion
	Hallucination
	Headache
	Nervousness
	Neurological Symptom
	Speech Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY / ORAL		Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:09/01/98ISR Number: 3204248-3Report Type:Periodic Company Report #A0066526  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL Other		Convulsion	Health Professional Company Representative	Zyban Tablet - Zyban	PS		ORAL

Date:09/01/98ISR Number: 3204251-3Report Type:Periodic Company Report #A0066427  
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG / TWICE PER DAY / ORAL		Arthralgia Chills Dyspnoea Hypersensitivity Myalgia Oedema Pain Pruritus Rash Maculo-Papular Serum Sickness Urticaria	Health Professional Company Representative	Zyban Tablet - Zyban	PS		ORAL

Date:09/01/98ISR Number: 3204255-0Report Type:Periodic Company Report #A0066361  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dermatitis	Professional				
TWICE PER DAY		Pruritus					
/ ORAL		Urticaria					

Date:09/01/98ISR Number: 3204259-8Report Type:Periodic Company Report #A0066333  
Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG ORAL							
Initial or Prolonged		Myocardial Infarction	Professional Company Representative				



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/01/98ISR Number: 3204260-4Report Type:Periodic  
Age:35 YR Gender:Male I/FU:I

Company Report #A0066226

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Urticaria	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /			Professional				
TWICE PER DAY							
/ ORAL							

Date:09/01/98ISR Number: 3204263-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0066137

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL				Wellbutrin Tablet - Controlled Release	SS		ORAL
ORAL							

Date:09/01/98ISR Number: 3204265-3Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0066078

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Pyrexia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
Initial or Prolonged		Urticaria					
TWICE PER DAY							
/ ORAL 2 WK							

Date:09/01/98ISR Number: 3204283-5Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #A0066028

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Disability	Confusional State	Health	Zyban Tablet - Zyban	PS	ORAL
150 MG /					
TWICE PER DAY	Deafness	Professional			
/ ORAL	Dizziness				
	Labyrinthitis		Hydrochlorothiazide	C	
	Meniere'S Disease		Fosinopril Sodium	C	
	Nausea				
	Vertigo				
	Vomiting				

Date:09/01/98ISR Number: 3204284-7Report Type:Periodic Company Report #A0066000  
Age:40 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Abdominal Pain Upper	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /						
Initial or Prolonged	Asthenia					
TWICE PER DAY						
/ ORAL	Hypersensitivity					
	Pain In Extremity					
	Urticaria					

Date:09/01/98ISR Number: 3204287-2Report Type:Periodic Company Report #A0065873  
Age:39 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Arterial Rupture
Initial or Prolonged	Face Oedema
Disability	Grand Mal Convulsion
	Hypersensitivity
	Nervous System Disorder
	Oedema Peripheral

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Urticaria				
		Visual Field Defect				
Dose	Duration		Report Source	Product	Role	Manufacturer
			Consumer	Zyban Tablet - Zyban	PS	
150 MG /						
TWICE PER DAY						
/ ORAL						ORAL

Date:09/01/98ISR Number: 3204288-4Report Type:Periodic Company Report #A0065854  
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet-Zyban	PS		ORAL
Hospitalization -		Convulsion					
150 MG /							
Initial or Prolonged		Depressed Level Of					
TWICE PER DAY							
/ ORAL		Consciousness					
		Dyskinesia		Aspirin	C		
		Eye Rolling					

Date:09/01/98ISR Number: 3204290-2Report Type:Periodic Company Report #A0065816  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban Tablet - Zyban	PS		ORAL
Other		Aggression					
150 MG / ORAL							
		Agitation	Professional	Medroxyprogesterone			
		Amnesia		Ace.	C		
		Anxiety					
		Disorientation					
		Disturbance In Attention					
		Excitability					
		Fluid Retention					
		Insomnia					
		Paranoia					
		Suicidal Ideation					
		Tinnitus					

Tremor

Date:09/01/98ISR Number: 3204291-4Report Type:Periodic Company Report #A0065790  
Age:73 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY  / ORAL	Decreased Appetite  Dizziness  Fall  Feeling Abnormal Hip Fracture Weight Decreased	Consumer	Zyban Tablet - Zyban	PS		ORAL
			Blood Pressure Medication	C		

Date:09/01/98ISR Number: 3204294-XReport Type:Periodic Company Report #A0065789  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG / ORAL	Dermatitis	Health  Professional	Zyban Tablet - Zyban	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/01/98ISR Number: 3204296-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0065788

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban Tablet - Zyban	PS		ORAL
Other		Dermatitis	Professional				
150 MG / ORAL							

Date:09/01/98ISR Number: 3204297-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0065786

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban Tablet - Zyban	PS		ORAL
Other		Dermatitis	Professional				
150 MG / ORAL							

Date:09/01/98ISR Number: 3204299-9Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0065762

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Respiratory Distress	Professional				
Initial or Prolonged		Urticaria	Company				
TWICE PER DAY							
Other			Representative				
/ ORAL							

Date:09/01/98ISR Number: 3204301-4Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0065736

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Health	Zyban Tablet - Zyban	PS		ORAL
150 MG ORAL		Convulsion	Professional				
Initial or Prolonged		Depressed Level Of Consciousness	Company	Nortriptyline	C		
Loss Of Consciousness							
Representative							

Date:09/01/98ISR Number: 3204304-XReport Type:Periodic  
Age:35 YR Gender:Male I/FU:I

Company Report #A0065658

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Asthenia	Professional				
TWICE PER DAY		Dermatitis					
/ ORAL		Hypersensitivity		Centrum	C		
		Myalgia					
		Oedema					
		Pain					
		Periorbital Oedema					
		Pruritus					
		Rash Erythematous					
		Swelling					
		Urticaria					

Date:09/01/98ISR Number: 3204307-5Report Type:Periodic  
Age:69 YR Gender:Female I/FU:I

Company Report #A0065655

Outcome	PT
Hospitalization -	Asthenia
Initial or Prolonged	Balance Disorder
	Dizziness
	Fall

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Headache Hypoaesthesia Vomiting	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER DAY / ORAL				Enalapril Maleate Alprazolam Verapamil	C C C		

Date:09/01/98ISR Number: 3205205-3Report Type:Periodic Company Report #A0056995  
Age:58 YR Gender:Female I/FU:F

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration		Health	Zyban Tablet - Zyban	PS		ORAL
Dose		Dysgeusia	Professional				
Disability		Nausea					
150 MG /UNK / ORAL		Oral Intake Reduced		Conjugated Estrogens Thyroxine Sodium	C C		

Date:09/01/98ISR Number: 3205207-7Report Type:Periodic Company Report #A0068175  
Age:40 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration		Health	Zyban Tablet- Zyban	PS		ORAL
Dose		Cerebral Infarction	Professional				
Hospitalization - 150 / MG / Initial or Prolonged ORAL		Cerebrovascular Accident	Company Representative	Simvastatin	C		

Date:09/01/98ISR Number: 3205209-0Report Type:Periodic Company Report #A0068019  
Age:49 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose							

Other                    Depression                    Health                    Zyban Tablet - Zyban   PS                    ORAL  
 150 / MG /  
 TWICE PER DAY                    Professional  
 / ORAL

...                    C  
 Diltiazem  
 Hydrochloride                    C  
 Estrogen                    C

Date:09/01/98ISR Number: 3205212-0Report Type:Periodic                    Company Report #A0067908  
 Age:55 YR    Gender:Female                    I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY / ORAL		Cerebrovascular Accident	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Chest Pain					
		Hypoaesthesia					
		Insomnia		Progesterone Calcium Salt Estrogen Multivitamin Alendronate	C C C C C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/01/98ISR Number: 3205214-4Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0067848

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG /	Abdominal Pain	Health	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Agitation	Professional				
/ ORAL		Anxiety					
		Crying					
		Paranoia					

Date:09/01/98ISR Number: 3205217-XReport Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #A0067448

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG /	Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
Initial or Prolonged	TWICE PER DAY	Chest Pain					
/ ORAL				Conjugated Estrogens	C		
				Medroxyprogesterone			
				Ace.	C		
				Multivitamin	C		
				Vitamin E	C		
				Ascorbic Acid	C		
				Aspirin	C		

Date:09/01/98ISR Number: 3205219-3Report Type:Periodic  
Age:28 YR Gender:Male I/FU:I

Company Report #A0067417

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	ORAL	Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
Initial or Prolonged							

Date:09/01/98ISR Number: 3205220-XReport Type:Periodic Company Report #A0067336  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Back Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Herbal Medication	C		

Date:09/01/98ISR Number: 3205224-7Report Type:Periodic Company Report #A0067218  
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Face Oedema	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pharyngeal Oedema					
TWICE PER DAY							
/ ORAL							
		Tongue Oedema					
		Urticaria					

Date:09/01/98ISR Number: 3205226-0Report Type:Periodic Company Report #A0067205  
Age:27 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Chest Pain
Initial or Prolonged	Hypoaesthesia
	Joint Swelling

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pharyngolaryngeal Pain  
Urticaria

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:09/01/98ISR Number: 3205229-6Report Type:Periodic Company Report #A0067189  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / PER Initial or Prolonged DAY / ORAL		Hallucination Mania	Health Professional Company Representative	Zyban Tablet - Zyban Lithium Salt	PS C		ORAL

Date:09/01/98ISR Number: 3205231-4Report Type:Periodic Company Report #A0067126  
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY / ORAL		Agitation Amnesia Anxiety Confusional State Disorientation Nervous System Disorder Speech Disorder	Health Professional	Zyban Tablet - Zyban Clarithromycin Cheratussin	PS C C		ORAL

Date:09/01/98ISR Number: 3205234-XReport Type:Periodic Company Report #A0066913  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dyspnoea					
TWICE PER DAY		Urticaria					
/ ORAL							

Date:09/01/98ISR Number: 3205236-3Report Type:Periodic Company Report #A0066889  
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Grand Mal Convulsion	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /			Professional				
Initial or Prolonged							
THREE TIMES							
PER DAY /							
ORAL							

Nabumetone C

Date:09/01/98ISR Number: 3205237-5Report Type:Periodic Company Report #A0066793  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Hypotension	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Urticaria					
Initial or Prolonged							
DAY / ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/01/98ISR Number: 3205240-5Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0066753

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Urticaria	Professional				
TWICE PER DAY							
/ ORAL							
				Multivitamin	C		
				Estrogen	C		
				Diphenhydramine Hcl	C		

Date:09/01/98ISR Number: 3205242-9Report Type:Periodic  
Age:55 YR Gender:Male I/FU:I

Company Report #A0066711

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Mania	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Suicidal Ideation	Professional				
TWICE PER DAY		Thinking Abnormal	Company				
/ ORAL			Representative				

Date:09/01/98ISR Number: 3205243-0Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #A0066646

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cerebral Infarction	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Grand Mal Convulsion	Professional				
Initial or Prolonged			Company				
TWICE PER DAY			Representative				
/ ORAL ;							

WEEKS

Date:09/01/98ISR Number: 3205245-4Report Type:Periodic Company Report #A0066587  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY  / ORAL		Dermatitis  Hypoaesthesia  Pharyngeal Oedema  Tachycardia	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:09/01/98ISR Number: 3205247-8Report Type:Periodic Company Report #A0066532  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG /  TWICE PER DAY  / ORAL		Amnesia  Convulsion  Dizziness  Eye Rolling Loss Of Consciousness Tremor	Health  Professional	Zyban Tablet -Zyban    Tramadol Hydrochloride Cyclobenzaprine Hcl	PS    C C		ORAL

Date:09/01/98ISR Number: 3205887-6Report Type:Periodic Company Report #A0065561  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG ORAL 6 DAY Initial or Prolonged		Convulsion	Consumer	Zyban Tablet-Zyban  Alprazolam Multivitamin Orqal Contraceptive	PS  C C C	Zyban	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/01/98ISR Number: 3205888-8Report Type:Periodic  
Age:28 YR Gender:Male I/FU:I

Company Report #A0065495

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG TWICE	Angioneurotic Oedema	Health	Zyban Tablet-Zyban	PS	Zyban	ORAL
Initial or Prolonged PER DAY ORAL	Dermatitis	Professional				
	Respiratory Distress Swelling Urticaria	Company Representative				

Date:09/01/98ISR Number: 3205889-XReport Type:Periodic  
Age:33 YR Gender:Male I/FU:I

Company Report #A0065339

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150MG PER DAY	Bladder Disorder	Health	Zyban Tablet-Zyban	PS	Zyban	ORAL
ORAL	Convulsion	Professional				
	Grand Mal Convulsion Hypoaesthesia Oral Loss Of Consciousness Urinary Incontinence		Zidovudine Lamivudine	C C		

Date:09/01/98ISR Number: 3205890-6Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0065334

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150MG TWICE	Convulsion	Health	Zyban Tablet-Zyban	PS	Zyban	ORAL
Other PER DAY ORAL	Syncope	Professional				
		Company Representative	Propranolol Hydrochloride	C		

Date:09/01/98ISR Number: 3205891-8Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0065298

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain Upper	Consumer	Zyban Tablet-Zyban	PS	Zyban	ORAL
150MG TWICE		Ageusia					
PER DAY ORAL		Anorexia					
		Back Pain					
		Chills					
		Dermatitis					
		Diarrhoea					
		Dysphagia					
		Hyperhidrosis					
		Pyrexia					
		Stevens-Johnson Syndrome					
		Thermal Burn					
		Throat Tightness					
		Tongue Oedema					

Date:09/01/98ISR Number: 3205892-XReport Type:Periodic Company Report #A0065206  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis	Consumer	Zyban Tablet-Zyban	PS	Zyban	ORAL
150MG TWICE		Hypersensitivity					
PER DAY ORAL		Swelling					
		Urticaria					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/01/98ISR Number: 3205893-1Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #A0064999

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG TWICE	Anaphylactic Reaction	Health	Zyban Tablet-Zyban	PS	Zyban	ORAL
Initial or Prolonged PER ORAL	Bronchospasm	Professional				
	Cyanosis Dermatitis Dyspnoea Pruritus Swelling Urticaria					

Date:09/01/98ISR Number: 3205894-3Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #A0064880

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG TWICE	Fall	Health	Zyban Tablet-Zyban	PS	Zyban	ORAL
Initial or Prolonged PER DAY ORAL	Grand Mal Convulsion	Professional				
Other	Hip Fracture	Company Representative	Flurbiprofen Gabapentin Nortriptyline Hcl Baclofen	C C C C		

Date:09/01/98ISR Number: 3205895-5Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #A0057792

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG PER DAY	Confusional State	Health	Zyban Tablet-Zyban	PS	Zyban	ORAL
Initial or Prolonged ORAL	Dehydration	Professional				
	Depressed Level Of Consciousness Hyponatraemia Polyuria		Oxpentifylline Nabumetone Claritin-D Lansoprazole	C C C C		

Vomiting  
Weight Decreased

Conjugated Estrogens C  
Dyazide C  
Alprazolam C  
Amitriptyline Hcl C  
Fluticasone  
Propionate C  
Aspirin C

Date:09/01/98ISR Number: 3205896-7Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #A0061791

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG TWICE Initial or Prolonged PER DAY ORAL	Blood Pressure Increased Cerebrovascular Accident	Health Professional	Zyban Tablet-Zyban	PS	Zyban	ORAL
	Dizziness Dyspnoea Fear Joint Stiffness Loss Of Consciousness Pain In Extremity Tachycardia		Conjugated Estrogens Calcium Salt + Vitamin	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/02/98ISR Number: 3125110-0Report Type:Expedited (15-DaCompany Report #A0069743  
Age:54 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - UNK/ORAL	Disorientation	Consumer	Zyban	PS		ORAL
Initial or Prolonged	Dizziness Palpitations		Nicotine	C		

Date:09/03/98ISR Number: 3125723-6Report Type:Expedited (15-DaCompany Report #A0065418  
Age:53 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening 150MG / PER	Angina Pectoris	Health	Zyban	PS		ORAL
Hospitalization - DAY / ORAL	Chest Discomfort	Professional				
Initial or Prolonged	Flushing	Company	Atenolol	C		
Other	Myocardial Infarction	Representative	Famotidine Amlodipine Oxaprozin	C C C		

Date:09/03/98ISR Number: 3125724-8Report Type:Expedited (15-DaCompany Report #A0069407  
Age:50 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150MG / TWICE	Dermatitis	Consumer	Zyban	PS		ORAL
Initial or Prolonged PER DAY /	Eyelid Oedema					
ORAL	Face Oedema					
	Syncope Urticaria					

Date:09/08/98ISR Number: 3126674-3Report Type:Expedited (15-DaCompany Report #A0070019  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Grand Mal Convulsion	Health	Zyban	PS		ORAL
150 MG ORAL			Professional	Amitriptyline Hcl	SS		ORAL
Hospitalization - ORAL			Company Representative				
Initial or Prolonged							

Date:09/09/98ISR Number: 3126593-2Report Type:Direct Company Report #  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Agitation		Wellbutrin	PS		
150 MG				Remeron	SS		
Intervention to		Akathisia		Effexor	SS		
30 MG HS							
Prevent Permanent		Muscle Rigidity					
150 BID							
Impairment/Damage		Serotonin Syndrome					

Date:09/09/98ISR Number: 3126616-0Report Type:Expedited (15-DaCompany Report #M086021  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Coma	Health	Serzone	PS		ORAL
Initial or Prolonged ORAL		Muscle Rigidity	Professional	Depakote	SS		ORAL
		Rash Generalised		Wellbutrin	SS		
				Triamterene	C		
				Timoptic	C		
				Potassium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Estrogen C  
 Synthroid C  
 Calcium C  
 Ativan C

Date:09/11/98ISR Number: 3128448-6Report Type:Expedited (15-DaCompany Report #A0069712  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Amylase Increased	Health	Wellbutrin	PS		ORAL
UNK, UNK,		Drug Interaction	Professional				
ORAL		Gout					
		Liver Function Test					
		Abnormal					
		Pancreatitis					
		Vomiting					

Date:09/14/98ISR Number: 3128407-3Report Type:Expedited (15-DaCompany Report #98090013  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Carisoprodol	PS		ORAL
DATA NA, PO		Coordination Abnormal		Wellbutrin	SS		ORAL
DATA NA. PO		Movement Disorder					

Date:09/14/98ISR Number: 3129147-7Report Type:Expedited (15-DaCompany Report #A0069197  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blister	Health	Wellbutrin	PS		ORAL
ORAL TAB		Epidermolysis Bullosa	Professional	Paroxetine			
		Porphyria Non-Acute		Hydrochloride	C		
		Pseudoporphyria		Trazodone	C		

Date:09/16/98ISR Number: 3130705-4Report Type:Direct  
Age:42 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS		ORAL
150 SR BID							

Date:09/16/98ISR Number: 3131117-XReport Type:Direct  
Age:77 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Mental Disorder		Famotidine	PS		ORAL
T1T PO B							
Initial or Prolonged				Bupropion Hcl	SS		ORAL
T2T PO Q							
				Acetaminophen	C		
				Aspirin	C		
				Digoxin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/17/98ISR Number: 3131637-8Report Type:Expedited (15-DaCompany Report #A0064074  
Age:27 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150MG/TWICE	Anaphylactic Shock	Health	Zyban	PS		ORAL
Hospitalization - PER DAY/ORAL	Chest Pain	Professional				
Initial or Prolonged	Dyspepsia Dyspnoea Food Interaction Hypersensitivity Pain In Extremity		Clove	SS		

Date:09/18/98ISR Number: 3132553-8Report Type:Expedited (15-DaCompany Report #A0070451  
Age:46 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability ORAL	Abdominal Distension	Foreign	Wellbutrin	PS		ORAL
	Oedema Peripheral	Consumer	Clonazepam Estropipate Medroxyprogesterone	C C C		

Date:09/21/98ISR Number: 3133432-2Report Type:Expedited (15-DaCompany Report #A0070019  
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG ORAL	Grand Mal Convulsion	Health	Zyban	PS		ORAL
Initial or Prolonged ORAL		Professional	Amitriptyline Hcl	SS		ORAL
		Company Representative				

Date:09/21/98ISR Number: 3133562-5Report Type:Expedited (15-DaCompany Report #A0070795  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Chest Pain	Health	Zyban	PS		ORAL
150 MG PER Hospitalization - ORAL		Dizziness	Professional				
Initial or Prolonged		Pulmonary Oedema	Company Representative	Fluoxetine Hydrochloride Dalmane Alprazolam	C C C		

Date:09/24/98ISR Number: 3134873-XReport Type:Expedited (15-DaCompany Report #98596.01  
Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Aggression Crying	Health Professional	Clonidine Hydrochloride	PS	Mylan	ORAL
0.2 MG Other QHS,ORAL		Delirium					
		Emotional Disorder Hallucination, Auditory Hypertonia		Wellbutrin Claritin-D	SS SS	Glaxo Schering	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/25/98ISR Number: 3135134-5Report Type:Expedited (15-DaCompany Report #A0071029  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Choking	Consumer	Zyban	PS		ORAL
150 MG/	ORAL	Foreign Body Trauma		Librax	C		
		Nausea					
		Oesophageal Obstruction					
		Pharyngolaryngeal Pain					

Date:09/25/98ISR Number: 3135138-2Report Type:Expedited (15-DaCompany Report #A0070661  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cardiac Arrest	Health	Wellbutrin	PS		ORAL
ORAL							
Initial or Prolonged		Ventricular Fibrillation	Professional	Lamictal	SS		ORAL
ORAL							
				Fluoxetine			
				Hydrochloride	SS		
				Clonazepam	SS		
				Enflurane	SS		

Date:09/30/98ISR Number: 3136930-0Report Type:Direct Company Report #  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased		Wellbutrin	PS		
75MG PM	150						
		Headache					

MG "SR" AM

(NDC#00173-01

35-00)

Date:10/01/98ISR Number: 3136982-8Report Type:Direct  
Age:50 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150MG/X2-4HR	Amnesia		Zyban	PS		
Hospitalization - Initial or Prolonged	Bipolar I Disorder Blast Injury Burns Second Degree Burns Third Degree Complex Partial Seizures Injury Insomnia Suicidal Ideation Suicide Attempt					

Date:10/01/98ISR Number: 3137126-9Report Type:Expedited (15-DaCompany Report #A0069363  
Age:86 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 100 MG/ ORAL	Grand Mal Convulsion	Foreign	Wellbutrin	PS		ORAL
Initial or Prolonged		Health	Frusemide	C		
Other		Professional	Omeprazole	C		
			Sinemet	C		
			Simethicone	C		
			Motilium	C		
			Olanzapine	C		
			Digoxin	C		
			Enalapril	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nitroglycerin C  
 Caffeine C  
 Sinemet 10/100 C  
 Benztropine C

Date:10/01/98ISR Number: 3137127-0Report Type:Expedited (15-DaCompany Report #A0070930  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiovascular Disorder	Foreign	Wellbutrin	PS		ORAL
ORAL			Health Professional Company Representative				

Date:10/02/98ISR Number: 3137372-4Report Type:Expedited (15-DaCompany Report #A0055625  
 Age:10 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Literature	Wellbutrin	PS		ORAL
Other		Drug Interaction	Health				
100 MG/THREE		Grand Mal Convulsion	Professional				
TIMES PER		Medication Error		Guanfacine Hydrochloride Tablet	SS		ORAL
DA/ORAL							
.5 MG/THREE							
TIMES PER							
DAY/ORAL							

Date:10/06/98ISR Number: 3138918-2Report Type:Expedited (15-DaCompany Report #A0070930  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

ORAL	Myocardial Infarction	Foreign	Wellbutrin	PS	ORAL
		Health Professional Company Representative			

Date:10/07/98ISR Number: 3139032-2Report Type:Direct Company Report #  
 Age:12 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 150 MG QAM, Intervention to 100 MG Q4 P Prevent Permanent 10 MG TAB QD Impairment/Damage 4 GTTS TID		Hypersensitivity  Urticaria		Wellbutrin Sr  Claritin  Cortisporin	PS  C  C		ORAL  ORAL

Date:10/07/98ISR Number: 3139488-5Report Type:Expedited (15-DaCompany Report #A0071135  
 Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 60 Hospitalization - TABLET/SINGLE Initial or Prolonged DOSE/ORAL		Convulsion  Overdose	Health  Professional Company Representative	Wellbutrin  Ethanol	PS  C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/07/98ISR Number: 3139498-8Report Type:Expedited (15-DaCompany Report #A0062864

Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Health	Zyban	PS		ORAL
150 MG/TWICE			Professional				
PER DAY/ORAL							

Date:10/09/98ISR Number: 3142162-2Report Type:Expedited (15-DaCompany Report #A0071731

Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Abnormal Behaviour	Health	Wellbutrin	PS		ORAL
ORAL							
Initial or Prolonged		Crying	Professional	Clonidine	C		ORAL
ORAL							
		Delirium	Other				
		Hallucination, Auditory					
		Muscle Disorder					

Date:10/09/98ISR Number: 3142164-6Report Type:Expedited (15-DaCompany Report #A0068282

Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis	Health	Wellbutrin	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL		Proteinuria	Professional				
			Company				
			Representative				

Date:10/09/98ISR Number: 3142498-5Report Type:Expedited (15-DaCompany Report #A0070674

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other ORAL	Constipation	Health	Wellbutrin	PS	ORAL
	Difficulty In Walking Fatigue Nausea	Professional			

Date:10/09/98ISR Number: 3142500-0Report Type:Expedited (15-DaCompany Report #A0067886  
Age:44 YR Gender:Female I/FU:I

Outcome Dose Hospitalization - TWICE PER DAY Initial or Prolonged / ORAL	PT	Report Source	Product	Role	Manufacturer	Route
	Depression	Consumer	Wellbutrin	PS		ORAL
	Diarrhoea					
	Influenza Like Illness		Phenelaine Sulphate	C		
	Myalgia		Clonazepam	C		
	Nausea		Antihistamine	C		
	Sleep Disorder					

Date:10/09/98ISR Number: 3142502-4Report Type:Expedited (15-DaCompany Report #A0071650  
Age:54 YR Gender:Female I/FU:I

Outcome Dose Hospitalization - 100 MG / Initial or Prolonged ORAL	Duration 6 WK	PT	Report Source	Product	Role	Manufacturer	Route
		Anaemia	Health	Wellbutrin	PS		ORAL
		Ecchymosis	Professional				
		Gingival Bleeding	Company				
		Neutropenia	Representative				
		Thrombocytopenia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/13/98ISR Number: 3141014-1Report Type:Expedited (15-DaCompany Report #A0070661

Age:61 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening ORAL	Cardiac Arrest	Health	Wellbutrin	PS		ORAL
Hospitalization - Initial or Prolonged Other	Ventricular Fibrillation	Professional	Lamictal Fluoxetine Hydrochloride	SS		
20 MG/PER DAY/ORAL			Clonazepam Enflurane Fentanyl	SS SS SS		
50 MCG / UNK / INTRAVENOUS			Propofol	SS		
150 MG / UNK / INTRAVENOUS			Mivacron	SS		
14 MG / UNK / INTRAVENOUS			Glycopyrronium Bromide	SS		
2 MG / UNK / INTRAVENOUS			Cephazolin	SS		
1 G / UNK / INTRAVENOUS			Oxygen Gas Nitrous Oxide Gas	SS SS		

Date:10/13/98ISR Number: 3141796-9Report Type:Expedited (15-DaCompany Report #A0072020

Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Hospitalization -  
ORAL  
Initial or Prolonged

Vasculitis

Health

Wellbutrin

PS

ORAL

Professional  
Company  
Representative

Date:10/13/98ISR Number: 3141798-2Report Type:Expedited (15-DaCompany Report #A0071991  
Age:40 YR Gender:Female I/FU:I

Outcome  
Hospitalization -  
Initial or Prolonged

- PT
- Abdominal Pain
- Abdominal Pain Upper
- Anaemia
- Chest Discomfort
- Chest Pain
- Chills
- Cholelithiasis
- Cough
- Dehydration
- Diarrhoea
- Drug Interaction
- Eye Irritation
- Gallbladder Disorder
- Gingival Bleeding
- Hypersensitivity
- Ileus Paralytic
- Insomnia
- Loss Of Consciousness
- Muscle Twitching
- Nasal Congestion
- Pollakiuria
- Pyrexia



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG/THREE TIMES PER DAY/ORAL		Rash Erythematous Red Blood Cell Sedimentation Rate Decreased Road Traffic Accident Throat Irritation	Consumer	Wellbutrin	PS		ORAL
		Weight Decreased White Blood Cell Count Increased		Prednisone Ortho-Cept Lansoprazole Pseudoephedrine Hcl Semisodium Valproate	C C C C C		

Date:10/14/98ISR Number: 3141239-5Report Type:Expedited (15-DaCompany Report #A0071455  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG ORAL		Dizziness Hypertension Loss Of Consciousness	Foreign Consumer	Zyban Blood Pressure Medication	PS C		ORAL

Date:10/19/98ISR Number: 3142501-2Report Type:Expedited (15-DaCompany Report #A0071152  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death ORAL		Death	Foreign Health Professional Company Representative	Wellbutrin	PS		ORAL

Date:10/19/98ISR Number: 3142870-3Report Type:Expedited (15-DaCompany Report #A0071650  
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	200 MG /	Anaemia	Health	Wellbutrin	PS		ORAL
Hospitalization -		Ecchymosis	Professional				
TWICE PER DAY							
Initial or Prolonged	6 WK	Gingival Bleeding	Company				
/ ORAL		Neutropenia	Representative	Conjugated Estrogens	C		
		Thrombocytopenia		Calcium Salt	C		
				Clonazepam	C		

Date:10/19/98ISR Number: 3142871-5Report Type:Expedited (15-DaCompany Report #A0072073  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG /TWICE	Chills	Consumer	Wellbutrin	PS		ORAL
Initial or Prolonged	PER DAY	Hyperhidrosis					
		Hypoaesthesia		Ranitidine			
		Memory Impairment		Hydrochloride	C		
		Pyrexia		Lisinopril	C		
		Vasculitis Cerebral					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/19/98ISR Number: 3142873-9Report Type:Expedited (15-DaCompany Report #A0072191  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Dizziness	Consumer	Zyban	PS		ORAL
Hospitalization - TWICE PER DAY		Haemorrhagic Stroke	Company				
Initial or Prolonged / ORAL		Headache	Representative				
TOPICAL	SEE TEXT /	Hypertension		Nicotine	SS		
TOPICAL		Intracranial Aneurysm					
		Loss Of Consciousness		Antibiotics	C		
		Malaise		Multivitamin	C		
		Pneumonia					
		Sinusitis					
		Tinnitus					

Date:10/19/98ISR Number: 3142874-0Report Type:Expedited (15-DaCompany Report #A0072222  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG /		Blood Creatinine	Health	Zyban	PS		ORAL
Initial or Prolonged TWICE PER DAY		Increased	Professional				
/ ORAL		Disorientation	Company				
		Dizziness	Representative	Calcium Salt	C		
		Dysarthria					
		Facial Palsy					
		Gait Disturbance					
		Headache					
		Hypercalcaemia					
		Hypertension					
		Nausea					
		Renal Failure Acute					
		Vomiting					

Date:10/19/98ISR Number: 3144277-1Report Type:Direct Company Report #  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Back Pain		Zyban	PS		ORAL
150 MG	ORAL						
Other		Chest Pain Palpitations Pyrexia Restlessness					

Date:10/20/98ISR Number: 3143685-2Report Type:Expedited (15-DaCompany Report #A0071912  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Confusional State	Foreign	Zyban	PS	Zyban	ORAL
150 MG/PER							
		Disturbance In Attention	Health				
DAY/ORAL							
		Hypercapnia Malaise	Professional				

Date:10/20/98ISR Number: 3143715-8Report Type:Expedited (15-DaCompany Report #1998-10-0429  
Age:15 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Abnormal Behaviour Anxiety

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
0.2 MG QHS		Crying Delirium Hallucination, Auditory	Report Source				
ORAL		Hypertonia	Other	Clonidine	PS		ORAL
1 TAB QD		Post-Traumatic Stress Disorder		Claritin-D 24 Hour	SS		ORAL
ORAL,							
EXTENDED							
RELEASE							
ORAL				Wellbutrin	SS		ORAL

Date:10/21/98ISR Number: 3144232-1Report Type:Direct  
Age:30 YR Gender:Female I/FU:I

Company Report #

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required COUMADIN DAW		International Normalised Ratio Increased		Wellbutrin Sr	PS		
Intervention to 10 MG		Prothrombin Time Prolonged		Procardia Lx	C		
Prevent Permanent Impairment/Damage							

Date:10/21/98ISR Number: 3145350-4Report Type:Expedited (15-DaCompany Report #A0072552  
Age:51 YR Gender:Male I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ TWICE		Feeling Abnormal	Consumer	Wellbutrin	PS		ORAL
Initial or Prolonged PER DAY /		Mydriasis Pallor					
ORAL		Sensation Of Pressure		Atorvastatin Calcium	C		

Date:10/21/98ISR Number: 3145351-6Report Type:Expedited (15-DaCompany Report #A0072624  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspnoea	Consumer	Zyban	PS		ORAL
150 MG /		Hypersensitivity					
TWICE PER DAY		Syncope					
/ ORAL		Urticaria					

Date:10/22/98ISR Number: 3145378-4Report Type:Expedited (15-DaCompany Report #A0072111  
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Drug Toxicity	Health	Wellbutrin	PS		ORAL
ORAL		Overdose	Professional	Lithium Salt	C		
				Valproic Acid	C		
				Risperidone	C		

Date:10/22/98ISR Number: 3145380-2Report Type:Expedited (15-DaCompany Report #A0064936  
Age:58 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Angina Pectoris	Health	Zyban	PS		ORAL
150 MG TWICE							
Initial or Prolonged		Heart Rate Increased	Professional				
PER DAY ORAL		Supraventricular Tachycardia		Lotrel	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/22/98ISR Number: 3145387-5Report Type:Expedited (15-DaCompany Report #A0068968  
Age:61 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG TWICE	Chest Pain	Health	Zyban	PS		ORAL
Initial or Prolonged PER DAY ORAL	Dyspnoea	Professional				
Other	Hypersensitivity Oesophagitis Pruritus Rash Pruritic		Aspirin	C		

Date:10/22/98ISR Number: 3145395-4Report Type:Expedited (15-DaCompany Report #A0072365  
Age:81 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 600 MG ORAL	Loss Of Consciousness	Foreign	Zyban	PS		ORAL
Initial or Prolonged		Consumer				

Date:10/22/98ISR Number: 3145397-8Report Type:Expedited (15-DaCompany Report #A0072775  
Age:46 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG TWICE	Asthenia	Foreign	Zyban	PS		ORAL
Initial or Prolonged PER DAY ORAL	Dizziness	Health				
	Myalgia Nausea Orthostatic Hypotension Pyrexia Vomiting	Professional				

Date:10/22/98ISR Number: 3145399-1Report Type:Expedited (15-DaCompany Report #A0072800  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG ORAL		Anxiety	Foreign	Zyban	PS		ORAL
Initial or Prolonged		Palpitations Peripheral Coldness	Health Professional				

Date:10/22/98ISR Number: 3145660-0Report Type:Expedited (15-DaCompany Report #A0072801  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG / PER DAY / ORAL		Chest Pain  Convulsion  Dizziness Dyspnoea Headache Tremor	Foreign  Consumer	Zyban	PS		ORAL

Date:10/26/98ISR Number: 3146418-9Report Type:Direct Company Report #  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged		Dermatitis Toxic Epidermal Necrolysis		Atenolol Wellbutrin Pepcid Solu-Medrol	PS SS C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/26/98ISR Number: 3146429-3Report Type:Direct  
Age:39 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 200 MG BID	Drug Toxicity		Tegretol	PS		ORAL
Initial or Prolonged ORAL	Intentional Misuse					
75 MG TID	Lethargy		Bupropion	SS		ORAL
ORAL	Sinus Tachycardia					
			Clonazepam	C		
			Fluphenazine	C		
			Risperidone	C		

Date:10/26/98ISR Number: 3147232-0Report Type:Expedited (15-DaCompany Report #A0073034  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG TWICE	Facial Palsy	Foreign	Zyban	PS		ORAL
Initial or Prolonged PER DAY ORAL	Hypoaesthesia	Consumer				

Date:10/26/98ISR Number: 3147234-4Report Type:Expedited (15-DaCompany Report #A0073094  
Age:15 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG ORAL	Convulsion	Health	Wellbutrin	PS		ORAL
Initial or Prolonged	Fall	Professional				
	Musculoskeletal Stiffness	Company				
	Overdose	Representative				
	Pyrexia					

Date:10/26/98ISR Number: 3147237-XReport Type:Expedited (15-DaCompany Report #A0072470  
Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypothyroidism	Health	Wellbutrin	PS		ORAL
150 MG TWICE		Petit Mal Epilepsy	Professional				
PER DAY ORAL			Company Representative				

Date:10/26/98ISR Number: 3150742-3Report Type:Expedited (15-DaCompany Report #A0072311  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion	Health	Zyban	PS		ORAL
150MG /		Dysarthria	Professional				
Initial or Prolonged		Fall					
SINGLE DOSE /		Head Injury					
ORAL TABLET		Tardive Dyskinesia					

Date:10/26/98ISR Number: 3150743-5Report Type:Expedited (15-DaCompany Report #A0073099  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Glucose Decreased	Foreign	Zyban	PS		ORAL
150MG / ORAL		Malaise	Consumer				
Initial or Prolonged		Palpitations		Glibenclamide	C		
TABLET				Metformin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/27/98ISR Number: 3244022-5Report Type:Periodic  
Age:60 YR Gender:Male I/FU:I

Company Report #9826591

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Viagra Tablets	PS		ORAL
50.00 MG		Hyperaesthesia					
TOTAL: PRN:							
ORAL							
				Zyban	SS		
NOT SPECIFIED							
				Nefazodone	SS		
NOT SPECIFIED							
				Buspirone	SS		
NOT SPECIFIED							

Date:10/27/98ISR Number: 3244523-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #9831535

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Laboratory Test Abnormal	Health	Viagra Tablets	PS		ORAL
100.00 MG			Professional				
TOTAL: PRN:							
ORAL							
				Zyban	SS		
				Ibuprofen	C		
				Flonase	C		
				Multi-Vitamin	C		
				Glucophage	C		

Date:10/27/98ISR Number: 3244937-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #9825988

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Health	Viagra Tablets	PS		ORAL
ORAL							

ORAL Drug Interaction Professional Wellbutrin SS ORAL

Date:10/29/98ISR Number: 3149880-0Report Type:Direct Company Report #  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Hepatocellular Damage		Wellbutrin	PS	Glaxo Wellcome	
450 MG QD				Tylenol	C		

Date:10/29/98ISR Number: 3163205-6Report Type:Periodic Company Report #A0060952  
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Convulsion	Health	Zyban Tablet-Zyban	PS		ORAL
150 MG /			Professional				
Other			Company				
TWICE PER DAY			Representative				
/ ORAL							

Date:10/30/98ISR Number: 3150122-0Report Type:Expedited (15-DaCompany Report #A0069746  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health	Wellbutrin	PS		ORAL
300 MG/ORAL		Intentional Misuse	Professional	Diphenhydramine Hcl	SS		
			Company				
			Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/30/98ISR Number: 3150123-2Report Type:Expedited (15-DaCompany Report #A0073387

Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG/TWICE	Dermatitis	Foreign	Zyban	PS		ORAL
PER DAY/ORAL		Difficulty In Walking	Consumer				
		Pain In Extremity					

Date:10/30/98ISR Number: 3150161-XReport Type:Expedited (15-DaCompany Report #A0087650

Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other	200 MG/TWICE	Anaemia Ecchymosis Gingival Bleeding	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
PER PAY/ORAL		Haemorrhage	Representative				
		Neutropenia Thrombocytopenia		Conjugated Estrogens Calcium Salt Clonazepam	C C C		

Date:10/30/98ISR Number: 3150372-3Report Type:Expedited (15-DaCompany Report #A0073559

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG/TWICE	Dehydration	Consumer	Zyban	PS	Zyban	ORAL
PER DAY/ORAL		Drug Ineffective					
		Nausea Pyrexia Sleep Disorder Weight Increased		Conjugated Estrogens Propranolol Hydrochloride	C C C		

Date:10/30/98ISR Number: 3150373-5Report Type:Expedited (15-DaCompany Report #A0073660

Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE		Angina Pectoris	Foreign	Zyban Tablet-Zyban	PS	Zyban	ORAL
Initial or Prolonged PER DAY/ORAL		Chest Pain	Consumer				
		Tremor					

Date:11/02/98ISR Number: 3257399-1Report Type:Periodic Company Report #8-98161-046A

Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 0.625 MG		Dermatitis	Consumer	Premarin Tablets	PS		
DAILY				Wellbutrin (Bupropion Hcl) One-A-Day Vitamins	SS C		

Date:11/02/98ISR Number: 3258317-2Report Type:Periodic Company Report #A0067845

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged 150 MG/TWICE		Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
PER DAY/ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tramadol  
Hydrochloride C

Date:11/02/98ISR Number: 3258325-1Report Type:Periodic Company Report #A0067984  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dermatitis Vomiting	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL							

Date:11/02/98ISR Number: 3258330-5Report Type:Periodic Company Report #A0068012  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other TWICE PER DAY/ ORAL		Eyelid Oedema Oedema Peripheral Pain In Extremity Urticaria	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:11/02/98ISR Number: 3258335-4Report Type:Periodic Company Report #A0068064  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG ORAL		Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release Sertraline Hydrochloride	PS C		ORAL

Date:11/02/98ISR Number: 3258339-1Report Type:Periodic Company Report #A0068180  
Age:61 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Dyspnoea Hypersensitivity Throat Tightness	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/TWICE						
PER DAY/ORAL			Propranolol Hydrochloride Ranitidine Hydrochloride Simvastatin	C C C		

Date:11/02/98ISR Number: 3258346-9Report Type:Periodic Company Report #A0068242  
Age:48 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Circulatory Collapse Convulsion	Health Professional Company  Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/TWICE						
PER DAY/ORAL						



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/02/98ISR Number: 3258447-5Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0068766

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
200 MG/TWICE			Representative				
PER DAY/ ORAL				Unknown Risperidone	C		

Date:11/02/98ISR Number: 3258455-4Report Type:Periodic  
Age:29 YR Gender:Male I/FU:I

Company Report #A0069672

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Consciousness Sleep Disorder	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/TWICE				Ethanol (Formulation Unknown)	SS		
PER DAY/ ORAL				Hyoscyamine Sulphate	C		

Date:11/02/98ISR Number: 3258470-0Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #A0070086

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
200 MG/TWICE				Fluoxetine Hydrochloride	C		
PER DAY/ ORAL				Clonazepam	C		

Olanzapine C  
Lithium Carbonate C

Date:11/02/98ISR Number: 3258479-7Report Type:Periodic Company Report #A0070432  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression Suicidal Ideation	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/TWICE							
PER DAY/ ORAL							
Mesalazine C							

Date:11/02/98ISR Number: 3258486-4Report Type:Periodic Company Report #A0070942  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/TWICE							
PER DAY/ ORAL							
Insulin C							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/02/98ISR Number: 3258490-6Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #A0071326

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	150 MG/TWICE PER DAY/ ORAL	Alcohol Interaction Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
				Ethanol (Formulation Unknown)	SS		ORAL

Date:11/02/98ISR Number: 3258494-3Report Type:Periodic  
Age:27 YR Gender:Female I/FU:F

Company Report #A0065796

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other	150 MG/TWICE PER DAY/ ORAL	Dermatitis Disorientation Dizziness Haematemesis Photopsia Pyrexia Stevens-Johnson Syndrome Urticaria	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:11/02/98ISR Number: 3258499-2Report Type:Periodic  
Age:39 YR Gender:Female I/FU:F

Company Report #A0065951

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Anorexia Convulsion	Health Professional	Wellbutrin Tablet-Controlled			

150 MG ORAL

Release

PS

ORAL

Date:11/02/98ISR Number: 3258505-5Report Type:Periodic  
Age:40 YR Gender:Female I/FU:F

Company Report #A0066958

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Urticaria	Health	Wellbutrin			
Other			Professional	Tablet-Controlled			
				Release	PS		ORAL

150 MG/TWICE

PER DAY/

ORAL

Date:11/02/98ISR Number: 3258509-2Report Type:Periodic  
Age:32 YR Gender:Female I/FU:F

Company Report #A0067131

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Condition Aggravated	Health	Wellbutrin			
		Depression	Professional	Tablet-Controlled			
		Suicidal Ideation		Release	PS		ORAL

150 MG/TWICE

PER DAY/ ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/02/98ISR Number: 3258519-5Report Type:Periodic  
 Age:46 YR Gender:Male I/FU:F

Company Report #A0067426

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Face Oedema Pruritus Urticaria	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/TWICE PER DAY/ ORAL							

Date:11/03/98ISR Number: 3151103-3Report Type:Direct  
 Age:45 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG PO BID		Flatulence		Wellbutrin Zoloft	PS C		ORAL

Date:11/03/98ISR Number: 3151105-7Report Type:Direct  
 Age:44 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG PO DAILY/150MG PO BID		Headache		Wellbutrin Prozac Prempro	PS C C		ORAL

Date:11/03/98ISR Number: 3151108-2Report Type:Direct  
 Age:44 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspnoea		Wellbutrin	PS		ORAL
150MG PO BID				Prozac	C		
				Prempro	C		

Date:11/03/98ISR Number: 3151545-6Report Type:Expedited (15-DaCompany Report #A0070549  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased	Health	Zyban	PS		ORAL
150 MG /TWICE							
PER DAY /ORAL		Difficulty In Walking	Professional				
		Hallucination		Allopurinol	C		
				Potassium Chloride	C		
				Atenolol	C		
				Indapamide	C		
				Nadolol	C		

Date:11/03/98ISR Number: 3152649-4Report Type:Expedited (15-DaCompany Report #107134  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Pressure Systolic	Health	Klonopin	PS		ORAL
ORAL							
Initial or Prolonged		Decreased	Professional	Fluoxetine	SS		ORAL
ORAL							
		Blood Pressure Systolic	Company	Wellbutrin	SS		ORAL
ORAL							
		Increased	Representative	Lamictal	SS		ORAL
ORAL							
		Cardiac Arrest		Anesthesia	SS		ORAL
ORAL							
		Ventricular Fibrillation		Anesthesia	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Epinephrine C

Date:11/04/98ISR Number: 3151914-4Report Type:Direct  
Age:30 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complications Of Maternal		Wellbutrin	PS		ORAL
1TAB BID PO		Exposure To Therapeutic Drugs Intra-Uterine Death					

Date:11/04/98ISR Number: 3152211-3Report Type:Expedited (15-DaCompany Report #A0071137  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Crying	Foreign	Wellbutrin	PS		ORAL
100 MG PER							
Hospitalization -		Dissociation	Health				
DAY ORAL							
Initial or Prolonged		Suicidal Ideation	Professional	Clonazepm	C		
Disability				Indapamide	C		
Other				Metformin	C		
				Sertraline			
				Hydrochloride	C		
				Zopiclone	C		
				Lorazepam	C		

Date:11/04/98ISR Number: 3152212-5Report Type:Expedited (15-DaCompany Report #A0073784  
Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin	PS		ORAL
150 MG TWICE							
PER DAY ORAL		Feeling Abnormal	Professional				
		Headache	Company	Venlafaxine			
		Loss Of Consciousness	Representative	Hydrochloride	C		

Sedation  
Speech Disorder  
Urinary Incontinence

Date:11/05/98ISR Number: 3152899-7Report Type:Expedited (15-DaCompany Report #A0072801  
Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain	Foreign	Zyban	PS		ORAL
150 MG PER							
DAY ORAL		Convulsion	Consumer				
		Dizziness					
		Dyspnoea					
		Headache					
		Tremor					

Date:11/05/98ISR Number: 3152977-2Report Type:Expedited (15-DaCompany Report #A0073934  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Muscle Spasms	Foreign	Zyban	PS		ORAL
150 MG/ PER							
DAY/ ORAL			Consumer				

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Freedom Of Information (FOI) Report

Date:11/05/98ISR Number: 3153105-XReport Type:Direct  
Age:43 YR Gender: I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening PO BID	Dysarthria		Zyban	PS		ORAL
Hospitalization - Initial or Prolonged	Fatigue Hypercalcaemia Hypertension Lethargy Renal Failure Acute Vomiting		Tenormin Cardizem Lasix	C C C		

Date:11/06/98ISR Number: 3153443-0Report Type:Expedited (15-DaCompany Report #A0064936  
Age:58 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY	Angina Pectoris Supraventricular Tachycardia	Health Professional	Zyban	PS		ORAL
/ ORAL			Lotrel	C		

Date:11/06/98ISR Number: 3153459-4Report Type:Expedited (15-DaCompany Report #A0073938  
Age:37 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY	Amnesia Asthenia Convulsion	Consumer	Zyban	PS		ORAL
/ ORAL	Dehydration Hypoaesthesia Nausea Viral Infection		Tramadol Hydrochloride (Formulation Unknown)	SS		ORAL

ORAL

Vision Blurred

Amoxicillin	C
Fluticasone	
Propionate	C
Fexofenadine	
Hydrochlorid	C
Amitriptyline Hcl	C
Lorcet	C
Cyclobenzaprine Hcl	C
Multivitamin	C

Date:11/06/98ISR Number: 3153463-6Report Type:Expedited (15-DaCompany Report #A0073662

Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY		Convulsion	Health  Professional	Wellbutrin	PS		ORAL
/ ORAL			Other	Paroxetine Hydrochloride (Formulation Unknown)	SS		ORAL
20 MG / PER DAY / ORAL				Omeprazoleq	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/10/98ISR Number: 3155811-XReport Type:Expedited (15-DaCompany Report #A0073917  
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Health	Wellbutrin	PS		ORAL
150 MG /		Injury	Professional				
TWICE PER		Injury Asphyxiation	Company				
DAY/ ORAL			Representative				

Date:11/10/98ISR Number: 3155814-5Report Type:Expedited (15-DaCompany Report #A0074104  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Health	Zyban	PS		
Other			Professional				
ORAL							

Date:11/10/98ISR Number: 3155816-9Report Type:Expedited (15-DaCompany Report #A0074143  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Facial Palsy	Foreign	Zyban	PS		ORAL
150 MG /			Consumer				
TWICE PER DAY							
/ ORAL							

Date:11/13/98ISR Number: 3156666-XReport Type:Expedited (15-DaCompany Report #A0074228  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Headache	Consumer	Zyban Tablet-Zyban	PS	Zyban	ORAL
150MG TWICE							

Initial or Prolonged Hypersensitivity  
PER DAY ORAL

Nervousness  
Tremor

Influenza Vaccine  
Injection SS

INTRAMUSCULAR INTRAMUSCULAR  
Weight Decreased  
Weight Increased

Presnisone C  
Salbutamol Sulphate C  
Ipratropium Bromide C  
Salmeterol Xinafoate C  
Salbutamol Sulphate C  
Oxygen C

Date:11/13/98ISR Number: 3157186-9Report Type:Expedited (15-DaCompany Report #A0074224  
Age:64 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ PER	Asthenia	Consumer	Wellbutrin	PS		ORAL
Initial or Prolonged DAY/ ORAL	Blood Pressure Increased					
	Dizziness Nausea		Atenolol Prempro	C C		

Date:11/13/98ISR Number: 3157189-4Report Type:Expedited (15-DaCompany Report #A0073385  
Age:45 YR Gender:Male I/FU:I

Outcome PT  
Hospitalization - Arthralgia  
Initial or Prolonged Delirium  
Dermatitis  
Malaise  
Myalgia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Pyrexia Serum Sickness	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Health	Wellbutrin	PS		ORAL
150 MG/	ORAL		Professional				

Date:11/13/98ISR Number: 3157480-1Report Type:Expedited (15-DaCompany Report #A0071499  
Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Creatine	Health	Zyban	PS		ORAL
Other		Phosphokinase Increased	Professional				
150MG/TWICE		Difficulty In Walking	Company				
PER DAY/ORAL		Liver Function Test	Representative				
		Abnormal					
		Monoparesis					
		Muscle Spasms					
		Rhabdomyolysis					

Date:11/13/98ISR Number: 3157491-6Report Type:Expedited (15-DaCompany Report #A0073503  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyskinesia	Foreign	Zyban	PS		ORAL
Disability		Dysphagia	Health				
150MG/TWICE		Dysphasia	Professional	Paracetamol	C		
PER DAY/ORAL		Hallucination					
		Hypoaesthesia					
		Tremor					

Date:11/13/98ISR Number: 3157495-3Report Type:Expedited (15-DaCompany Report #A0064936  
Age:58 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG/TWICE		Heart Rate Increased	Health	Zyban	PS		ORAL
Initial or Prolonged PER DAY/ORAL		Supraventricular Tachycardia	Professional	Lotrel	C		

Date:11/13/98ISR Number: 3157501-6Report Type:Expedited (15-DaCompany Report #A0069743  
Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Disorientation	Health	Zyban	PS		ORAL
Initial or Prolonged		Dizziness Palpitations	Professional	Nicotine	C		

Date:11/13/98ISR Number: 3157766-0Report Type:Direct Company Report #  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100MG QD PO		Paraesthesia		Bupropion	PS		ORAL
		Paraesthesia Oral		Ritalin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/16/98ISR Number: 3157769-6Report Type:Expedited (15-DaCompany Report #A0072801

Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain	Foreign	Zyban	PS		ORAL
150 MG/PER		Convulsion	Health				
DAY/ORAL		Dizziness	Professional				
		Dyspnoea					
		Headache					
		Panic Attack					
		Tremor					

Date:11/16/98ISR Number: 3157772-6Report Type:Expedited (15-DaCompany Report #A0072222

Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Creatine Increased	Health	Zyban	PS		ORAL
150 MG/TWICE		Disorientation	Professional				
Initial or Prolonged		Dizziness	Company	Calcium Salt	C		
PER DAY/ORAL		Dysarthria	Representative	Atenolol	C		
		Facial Palsy		Diltiazem			
		Gait Disturbance		Hydrochloride	C		
		Headache		Frusemide	C		
		Hypercalcaemia					
		Nausea					
		Renal Failure Acute					
		Secondary Hypertension					
		Vomiting					

Date:11/19/98ISR Number: 3160208-2Report Type:Expedited (15-DaCompany Report #A0074896

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspnoea	Foreign	Zyban	PS		ORAL
150 MG/TWICE							

Facial Palsy  
Health  
Professional

Date:11/19/98ISR Number: 3160234-3Report Type:Expedited (15-DaCompany Report #A0073975  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Toxic Epidermal	Health	Wellbutrin	PS		ORAL
ORAL	3 WK						
		Necrolysis	Professional	Lamictal	SS		ORAL
ORAL	5 MON						

Date:11/19/98ISR Number: 3160235-5Report Type:Expedited (15-DaCompany Report #A0074905  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cardiac Fibrillation	Foreign	Zyban	PS		ORAL
150 MG/ TWICE							
Initial or Prolonged			Consumer				
PER DAY/ ORAL							

Date:11/19/98ISR Number: 3160538-4Report Type:Expedited (15-DaCompany Report #A0068408  
Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Thrombocytopenia	Health	Wellbutrin	PS		ORAL
150 MG/TWICE							
			Professional				
PER DAY/ORAL							

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Freedom Of Information (FOI) Report

Lithium Salt C  
 Naproxen C

Date:11/20/98ISR Number: 3160780-2Report Type:Expedited (15-DaCompany Report #A0066944  
 Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG /TWICE Initial or Prolonged PER DAY /		Chest Discomfort	Consumer	Wellbutrin	PS		ORAL
ORAL 150 MG		Chest Pain		Wellbutrin	SS		ORAL
ORAL/CONTROLL ED RELEASE				Clonazepam Aspirin Sertraline Hydrochloride	C C C C		

Date:11/20/98ISR Number: 3160782-6Report Type:Expedited (15-DaCompany Report #A0072470  
 Age:19 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG TWICE PER DAY ORAL		Hypothyroidism	Health	Wellbutrin	PS		ORAL
		Petit Mal Epilepsy	Professional Company Representative				

Date:11/20/98ISR Number: 3160930-8Report Type:Expedited (15-DaCompany Report #A0073853  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Asthenia Balance Disorder Decreased Appetite	Foreign Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
UNK/UNK/ORAL		Difficulty In Walking Migraine Myalgia Nausea Speech Disorder		Clonazepam Oestradiol Misoprostol	C C C		

Date:11/23/98ISR Number: 3161793-7Report Type:Expedited (15-DaCompany Report #A0070019  
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG ORAL Initial or Prolonged		Grand Mal Convulsion	Health Professional Company Representative	Zyban Amitriptyline Hcl (Formulation Unknown)	PS SS		ORAL ORAL
ORAL							

Date:11/23/98ISR Number: 3161795-0Report Type:Expedited (15-DaCompany Report #A0074755  
Age:32 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Accidental Overdose Aggression Agitation

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Delirium Electrocardiogram Qrs Complex Prolonged	Report Source	Product	Role	Manufacturer	Route
9 G SINGLE DOSE ORAL		Electrocardiogram Qt Prolonged	Literature	Wellbutrin	PS		ORAL
		Feeling Jittery Grand Mal Convulsion Sinus Tachycardia Supraventricular Tachycardia					

Date:11/24/98ISR Number: 3281999-6Report Type:Periodic Company Report #0175957A  
Age:65 YR Gender:Female I/FU:I

Outcome Dose Other	Duration	PT Drug Effect Decreased	Report Source	Product	Role	Manufacturer	Route
1 PATCH/QD/TTS UNKNOWN/UNK/B ID			Consumer	Habitrol-Nicotine Transdermal 21mg-Nvch	PS		
				Zyban-Bupropion Hcl 150mg-Glaxo	SS	Glaxo	
				Lorazepam Bid	C		

Date:11/24/98ISR Number: 3283068-8Report Type:Periodic Company Report #0182444A  
Age: Gender:Male I/FU:I

Outcome Dose Other	Duration	PT Abdominal Pain Dyspepsia Tremor	Report Source	Product	Role	Manufacturer	Route
TRANSDERMAL TTS	1 PATCH QD		Consumer	Habitrol-Nicotine Transdermal 21mg-Nvch	PS	Nvch	

BID PO

Zyban-Bupropion Hcl	SS	Glaxo	ORAL
150mg-Glaxo			
Insulin	C		
Univasc	C		
Hctz	C		
Glyburide	C		
Lipitor	C		
Lanoxin	C		
Verapamil	C		
Quinaglute	C		
Zantac	C		
Catapres	C		

Date:11/25/98ISR Number: 3286987-1Report Type:Periodic Company Report #8-97286-019L  
 Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiovascular Disorder	Consumer	Redux	PS		ORAL
15 MG TWICE		Palpitations					
DAILY ORAL				Fastin (Phentermine)	SS		ORAL
30 MG ONCE							
DAILY ORAL				Ionamin (Phentermine)	SS		ORAL
15-30 MG							
DAILY ORAL				Ionamin (Phentermine)	SS		ORAL
8/31/96-10/11							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

/1996;

5/22/1997-

8/25/97 ORAL 5 MON

Pondimin  
(Fenfluramine)  
Tablet

SS

ORAL

INTERMITTENT:

03APR96-11OCT

96;

22MAY97-25AUG

97 ORAL

Wellbutrin  
(Bupropion)

SS

ORAL

100-150 MG

TWICE DAILY

ORAL

Date:11/27/98ISR Number: 3163468-7Report Type:Expedited (15-DaCompany Report #A0075616

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hypertensive Crisis	Foreign	Zyban	PS		ORAL
150 MG/TWICE			Health				
PER DAY/ORAL			Professional	Ramipril	C		

Date:11/27/98ISR Number: 3163469-9Report Type:Expedited (15-DaCompany Report #A0075934

Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Dermatitis	Foreign	Zyban	PS		ORAL
150 MG/TWICE							

Initial or Prolonged Respiratory Disorder  
PER DAY/ORAL

Health  
Professional

Date:11/27/98ISR Number: 3163513-9Report Type:Expedited (15-DaCompany Report #A0075174  
Age:59 YR Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY	Psychotic Disorder Suicidal Ideation	Health Professional Company Representative	Wellbutrin Gabapentin	PS C		ORAL

Date:11/27/98ISR Number: 3163544-9Report Type:Expedited (15-DaCompany Report #A0072311  
Age:4 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ Initial or Prolonged SINGLE DOSE/ ORAL	Convulsion Dysarthria Fall Head Injury Laboratory Test Abnormal Medication Error Tardive Dyskinesia	Health Professional	Zyban	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/27/98ISR Number: 3163848-XReport Type:Expedited (15-DaCompany Report #A0075570

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2 WK	Completed Suicide	Consumer	Wellbutrin	PS		ORAL
100 MG/ORAL							

Date:11/30/98ISR Number: 3164009-0Report Type:Direct

Age:31 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Convulsion		Wellbutrin	PS		
150MG BID;				Lamictal	SS		

Date:12/01/98ISR Number: 3164970-4Report Type:Direct

Age:15 YR Gender: I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Ineffective		Wellbutrin	PS		
150 MG SR BID							

Date:12/01/98ISR Number: 3290599-3Report Type:Periodic

Age:37 YR Gender:Female I/FU:F

Company Report #A0068116

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Arthralgia Hypersensitivity	Health Professional	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Pruritus					
PER DAY/ORAL		Rash Erythematous Swelling Urticaria					

Date:12/01/98ISR Number: 3290605-6Report Type:Periodic Company Report #A0068091  
Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema Arthralgia	Health Professional	Zyban Tablets - Zyban	PS		ORAL
150 MG/TWICE		Dysphagia					
PER DAY/ORAL		Oedema Peripheral Rash Erythematous Urticaria		Vicodin	C		

Date:12/01/98ISR Number: 3290611-1Report Type:Periodic Company Report #A0067771  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypersensitivity Pruritus	Health Professional	Zyban Tablets - Zyban	PS		ORAL
150 MG/TWICE		Pyrexia					
PER DAY/ORAL		Rash Erythematous Urticaria					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/01/98ISR Number: 3290617-2Report Type:Periodic  
Age:27 YR Gender:Female I/FU:F

Company Report #A0067616

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Arthralgia	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Chest Pain	Professional				
PER DAY ORAL		Dysphagia					
		Dyspnoea					
		Urticaria					

Date:12/01/98ISR Number: 3290621-4Report Type:Periodic  
Age:54 YR Gender:Female I/FU:F

Company Report #A0065854

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Convulsion	Health	Zyban Tablet- Zyban	PS		ORAL
150 MG/TWICE		Depressed Level Of	Professional				
Initial or Prolonged		Consciousness		Arpirin	C		
PER DAY/ORAL		Dyskinesia					
		Eye Movement Disorder					

Date:12/01/98ISR Number: 3290623-8Report Type:Periodic  
Age:50 YR Gender:Female I/FU:F

Company Report #A0066078

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Pyrexia	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Urticaria	Professional				
Initial or Prolonged							
PER DAY/ORAL	2 WK						
Other							

Date:12/01/98ISR Number: 3290626-3Report Type:Periodic  
Age:49 YR Gender:Female I/FU:F

Company Report #A0065298

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Abdominal Pain Upper	Health	Zyban Tablet - Zyban	PS		ORAL
Other		Ageusia	Professional				
150 MG/TWICE		Anorexia					
PER DAY/ORAL		Back Pain					
		Chills					
		Dermatitis					
		Diarrhoea					
		Dysphagia					
		Hyperhidrosis					
		Oedema Mouth					
		Pyrexia					
		Stevens-Johnson Syndrome					
		Throat Irritation					
		Throat Tightness					
		Tongue Oedema					

Date:12/01/98ISR Number: 3290628-7Report Type:Periodic Company Report #A0065206  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dermatitis	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Ecchymosis	Professional				
PER DAY/ORAL		Hypersensitivity		Nabumetone	C		
		Pruritus		Naproxen	C		
		Swelling		Multivitamin	C		
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/01/98ISR Number: 3290631-7Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0074805

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Zyban Tablet - Zyban	PS		ORAL
ORAL							

Date:12/01/98ISR Number: 3290653-6Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #A0073925

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL		Dysphagia					
		Hypersensitivity					
		Pruritus					
		Urticaria					

Date:12/01/98ISR Number: 3290656-1Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #A0073954

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Insomnia	Consumer	Zyban Tablets -			
Initial or Prolonged		Syncope		Zyban	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL		Vomiting					
				Temazepam	C		
				Fioricet	C		

Date:12/01/98ISR Number: 3290658-5Report Type:Periodic  
Age:41 YR Gender:Male I/FU:I

Company Report #A0073851

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Zyban Tablets -			

Decreased Activity Zyban PS ORAL  
150 MG/TWICE  
DAILY PER  
DAY/ORAL  
Glimepiride C

Date:12/01/98ISR Number: 3290661-5Report Type:Periodic Company Report #A0073618  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Study	Zyban Tablet - Zyban	PS		ORAL
ORAL		Convulsion Ear Infection Syncope	Consumer				

Date:12/01/98ISR Number: 3290664-0Report Type:Periodic Company Report #A0072686  
Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anorexia	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER							
Initial or Prolonged		Grand Mal Convulsion					
DAY/ORAL		Insomnia		Co-Trimoxazole Phenazopyridine Hcl	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/01/98ISR Number: 3290669-XReport Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0071863

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ORAL		Angioneurotic Oedema	Health	Zyban Tablet- Zyban	PS		ORAL
Initial or Prolonged Disability		Urticaria	Professional Company Representative				
Other							

Date:12/01/98ISR Number: 3290675-5Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #A0071611

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Health Professional Company Representative	Zyban Tablet - Zyban	PS		ORAL
150MG/PER DAY/ORAL							

Date:12/01/98ISR Number: 3290681-0Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0071580

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE PER DAY/ORAL		Tongue Oedema					
		Urticaria		Conjugated Estrogens	C		

Date:12/01/98ISR Number: 3290686-XReport Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0071549

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE		Confusional State	Consumer	Zyban Tablet - Zyban	PS		ORAL

Initial or Prolonged Convulsion Company  
PER DAY/ORAL  
Other Loss Of Consciousness Representative Oral Contraceptive C

Date:12/01/98ISR Number: 3290691-3Report Type:Periodic Company Report #A0071499  
Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG/TWICE	Alanine Aminotransferase	Health	Zyban Tablet - Zyban	PS		ORAL
PER DAY/ORAL		Increased	Professional				
		Aspartate Aminotransferase	Company Representative				
		Increased					
		Blood Creatine Phosphokinase					
		Increased					
		Blood Lactate Dehydrogenase					
		Increased					
		Difficulty In Walking Muscle Spasms					
		Rhabdomyolysis					

Date:12/01/98ISR Number: 3290695-0Report Type:Periodic Company Report #A0071281  
Age:38 YR Gender:Male I/FU:I

Outcome	PT
Other	Arthralgia Eye Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Face Oedema Oedema Peripheral Pruritus	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Rash Erythematous	Consumer	Zyban Tablet- Zyban	PS		ORAL

Claritin-D	C
Robitussin-Dm	C
Clarithromycin	C

Date:12/01/98ISR Number: 3291062-6Report Type:Periodic Company Report #A0071278  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG/ TWICE PER DAY/ ORAL		Dyspnoea Pruritus Rash Vesicular Urticaria	Consumer	Zyban Tablet - Zyban Adrenaline	PS C		ORAL

Date:12/01/98ISR Number: 3291063-8Report Type:Periodic Company Report #A0070995  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG/ TWICE PER DAY/ ORAL		Aphasia Balance Disorder Convulsion Dyspraxia Hypersomnia Sedation	Consumer	Zyban Tablet - Zyban Lorazepam Dyazide	PS C C		ORAL

Date:12/01/98ISR Number: 3291064-XReport Type:Periodic Company Report #A0070995  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphasia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Balance Disorder					
PER DAY/		Convulsion					
ORAL		Dyspraxia		Lorazepam	C		
		Hypersomnia		Dyazide	C		
		Sedation					

Date:12/01/98ISR Number: 3291065-1Report Type:Periodic Company Report #A0070651  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG			Professional				
ORAL			Company Representative				

Date:12/01/98ISR Number: 3291066-3Report Type:Periodic Company Report #A0070459  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Neck Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY							
ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Prempro

C

Date:12/01/98ISR Number: 3291067-5Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #A0070204

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Consciousness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG PER		Pruritus					
DAY	ORAL	Tremor					

Date:12/01/98ISR Number: 3291068-7Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0070194

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG PER		Convulsion					
DAY	ORAL	Drug Ineffective		Ibuprofen	C		

Date:12/01/98ISR Number: 3291069-9Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #A0070089

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Drug Ineffective					
Initial or Prolonged		Headache					
PER DAY							
ORAL				Ibuprofen	C		
				Cocaine	C		
				Ethanol	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG		Convulsion	Health	Zyban Tablet - Zyban	PS		ORAL
ORAL			Professional				
				Salbutamol Sulphate	C		
				Antibiotics	C		
				Beclomethasone			
				Dipropion	C		
				Theophylline	C		
				Benzonatate	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG/ TWICE		Anxiety	Health	Zyban Tablet - Zyban	PS		ORAL
PER DAY		Suicidal Ideation	Professional				
ORAL			Company				
			Representative	Thyroxine Sodium	C		
				Metoprolol Tartrate	C		
				Oestradiol	C		
				Triamcinolone			
				Acetonide	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/01/98ISR Number: 3291072-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0069735

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG/ TWICE	Hypersensitivity	Health	Zyban Tablet - Zyban	PS		ORAL
Hospitalization -	PER DAY	Hypertension	Professional				
Initial or Prolonged							
ORAL							
Other				Moexipril Hydrochloride	C		
				Omeprazole	C		

Date:12/01/98ISR Number: 3291073-0Report Type:Periodic  
 Age:24 YR Gender:Female I/FU:I

Company Report #A0069624

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG	Convulsion	Consumer	Zyban Tablet - Zyban	PS		ORAL
THREE TIMES							
PER DAY							
ORAL							

Date:12/01/98ISR Number: 3291074-2Report Type:Periodic  
 Age:39 YR Gender:Male I/FU:I

Company Report #A0069512

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG	Dysgraphia	Health	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Eyelid Oedema	Professional				
ORAL		Hypersensitivity					
		Pruritus					
		Swelling					
		Urticaria					

Date:12/01/98ISR Number: 3291075-4Report Type:Periodic Company Report #A0069465  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Claustrophobia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG	PER	Dizziness					
DAY	ORAL	Loss Of Consciousness					
		Panic Disorder Without					
		Agoraphobia					
		Paranoia					

Date:12/01/98ISR Number: 3291076-6Report Type:Periodic Company Report #A0069451  
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Asthenia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG		Dizziness					
Initial or Prolonged		Drug Effect Decreased					
TWICE PER DAY		Dry Mouth					
ORAL							

Date:12/01/98ISR Number: 3291077-8Report Type:Periodic Company Report #A0069149  
Age:63 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Chills
Initial or Prolonged	Erythema Multiforme

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Pruritus Rash Maculo-Papular	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Health	Zyban Tablet - Zyban	PS		ORAL
150 MG			Professional				
TWICE PER DAY							
ORAL							
				Methyldopa	C		
				Famotidine	C		
				Thyroxine Sodium	C		
				Diazepam	C		
				Conjugated Estrogens	C		
				Felodipine	C		
				Fluvastatin Sodium	C		
				Dextropropoxyphene	C		
				Hydrocodone	C		
				Morphine	C		
				Prednisone	C		

Date:12/02/98ISR Number: 3165485-XReport Type:Direct  
Age:41 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction		Bupropion Sr	PS		
150MG BID							
Required		Haemorrhage		Warfarin	SS		
7.5MG QD							
Intervention to		International Normalised					
Prevent Permanent		Ratio Increased					
Impairment/Damage		Pain					
		Prothrombin Time					
		Prolonged					

Date:12/02/98ISR Number: 3165971-2Report Type:Direct  
Age:15 YR Gender: I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other  
150MG SR BID

Drug Ineffective

Health

Wellbutrin

PS

Professional

Date:12/03/98ISR Number: 3166694-6Report Type:Expedited (15-DaCompany Report #A0075267

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Health	Wellbutrin	PS		ORAL
ORAL			Professional Company Representative	Fluoxetine Hydrochloride	C		

Date:12/03/98ISR Number: 3166722-8Report Type:Expedited (15-DaCompany Report #A0075941

Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Consumer	Zyban	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL				Propranolol	C		
				Montelukast Sodium	C		
				Omeprazole	C		
				Captopril	C		
				Cetirizine			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride C

Date:12/04/98ISR Number: 3167501-8Report Type:Direct  
 Age:29 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	3 WK	Angioneurotic Oedema		Zyban	PS	Glaxo Wellcome	
Hospitalization - Initial or Prolonged Required		Dermatitis Face Oedema Oedema Mouth Tongue Oedema					
Intervention to Prevent Permanent Impairment/Damage							

Date:12/04/98ISR Number: 3167537-7Report Type:Direct  
 Age:34 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Zyban	PS		
PER							

DIRECTIONS

Synthroid C  
 Antihypertensives C

Date:12/08/98ISR Number: 3168529-4Report Type:Expedited (15-DaCompany Report #A0075616  
 Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG /	Hypertensive Crisis	Foreign	Zyban	PS		ORAL
TWICE PER DAY			Health				
/ ORAL			Professional				
				Ramipril	C		
				Lorazepam	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Arthralgia	Health	Wellbutrin	PS		ORAL
150 MG ORAL						
Hospitalization -	Decreased Appetite	Professional				
Initial or Prolonged	Dyspnoea Exertional					
Disability	Erythema Multiforme					
Other	Eyelid Oedema					
	Face Oedema					
	Fatigue					
	Insomnia					
	Periorbital Oedema					
	Proteinuria					
	Pruritus					
	Stevens-Johnson Syndrome					
	Throat Tightness					
	Urticaria					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/08/98ISR Number: 3168536-1Report Type:Expedited (15-DaCompany Report #A0076073

Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - ORAL	Duration Alcohol Withdrawal Syndrome	Health	Wellbutrin	PS		ORAL
Initial or Prolonged ORAL	Electroencephalogram Abnormal	Professional Company	Ethanol (Formulation Unknown)	SS		ORAL
ORAL	Grand Mal Convulsion Overdose Suicide Attempt	Representative	Glipizide Tablet	SS		ORAL
			Glucophage	C		

Date:12/08/98ISR Number: 3168578-6Report Type:Expedited (15-DaCompany Report #A0076277

Age:36 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY	Duration Anxiety Conversion Disorder	Foreign	Zyban	PS		ORAL
	Diarrhoea Dizziness Nervousness Tremor Vomiting	Consumer				

Date:12/08/98ISR Number: 3168731-1Report Type:Expedited (15-DaCompany Report #A0076538

Age:39 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - ORAL	Duration 1 YR Dermatitis	Health	Wellbutrim	PS		ORAL
Initial or Prolonged ORAL	Drug Interaction	Professional	Diphenhydramine Hcl	SS		ORAL
Other	Pyrexia	Company Representative	Co-Trimoxazole	C		

Date:12/08/98ISR Number: 3168853-5Report Type:Expedited (15-DaCompany Report #A0076428  
Age:51 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY	Cellulitis Fall	Consumer	Zyban	PS		ORAL
/ORAL	Grand Mal Convulsion		Conjugated Estrogens	C		

Date:12/10/98ISR Number: 3168834-1Report Type:Expedited (15-DaCompany Report #8-98338-012A  
Age:10 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other 0.5 MG TWICE DAILY, ORAL 75 MG TO 300 MG DAILY, ORAL	Drug Interaction Grand Mal Convulsion	Literature	Tenex Bupropion	PS SS		ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/10/98ISR Number: 3169454-5Report Type:Direct  
 Age:51 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG BID PO	Cerebral Arteritis		Zyban	PS		ORAL
		Cognitive Disorder					
		Coronary Artery Occlusion					
		Flushing					
		Hallucination, Auditory					
		Neurosis					
		Temporal Arteritis					
		Vomiting					

Date:12/11/98ISR Number: 3169752-5Report Type:Expedited (15-DaCompany Report #A0076551  
 Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG /	Arthritis	Foreign	Zyban	PS		ORAL
	TWICE PER DAY	Dermatitis	Health				
		Pharyngolaryngeal Pain	Professional				

Date:12/11/98ISR Number: 3169755-0Report Type:Expedited (15-DaCompany Report #A0076359  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	PER DAY	Overdose	Health	Wellbutrin	PS		ORAL
			Professional				

Date:12/11/98ISR Number: 3170074-7Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Required	Dermatitis	Wellbutrin Sr	PS	ORAL
1 PO BID				
Intervention to		Habitol	C	
Prevent Permanent		Valium	C	
Impairment/Damage				

Date:12/11/98ISR Number: 3170075-9Report Type:Direct Company Report #  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Dyspnoea		Wellbutrin Sr	PS	Glaxo/Wellcome	ORAL
1 PO BID							
Intervention to		Eyelid Oedema					
Prevent Permanent		Urticaria					
Impairment/Damage							

Date:12/14/98ISR Number: 3170787-7Report Type:Expedited (15-DaCompany Report #A0076758  
 Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebrovascular Accident	Consumer	Zyban	PS		ORAL
150 MG /							
Hospitalization -		Chest Pain					
SINGLE DOSE /							
Initial or Prolonged		Insomnia					
ORAL							
		Myocardial Infarction		Paracetamol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/14/98ISR Number: 3170791-9Report Type:Expedited (15-DaCompany Report #A0066944  
 Age:68 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100 MG / Initial or Prolonged TWICE PER DAY Disability / ORAL Other 150 MG / UNK / ORAL	Chest Discomfort  Chest Pain  Coronary Artery Atherosclerosis  Coronary Artery Occlusion	Health  Professional	Wellbutrin	PS		ORAL
			Clonazepam	C		
			Aspirin	C		
			Sertraline			
			Hydrochloride	C		

Date:12/16/98ISR Number: 3172028-3Report Type:Direct Company Report #  
 Age:62 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required 100 MG BID Intervention to Prevent Permanent Impairment/Damage	Convulsion  Tremor		Wellbutrin  Vicodin Prn	PS  C		

Date:12/17/98ISR Number: 3171578-3Report Type:Expedited (15-DaCompany Report #A0076725  
 Age:57 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ ORAL	Arrhythmia  Convulsion	Consumer	Zyban	PS		ORAL
			Carbamazepine (Formulaton Unknown)	SS		
			Pravastatin Sodium	C		

Date:12/17/98ISR Number: 3171579-5Report Type:Expedited (15-DaCompany Report #A0077179  
Age:61 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - PER DAY ORAL Initial or Prolonged	Atrial Fibrillation	Consumer	Zyban	PS		ORAL

Date:12/17/98ISR Number: 3171581-3Report Type:Expedited (15-DaCompany Report #A0073598  
Age:42 YR Gender:Female I/FU:I

Outcome	PT
Other	Anaphylactic Reaction Blood Cholesterol Increased Blood Pressure Increased Chest Discomfort Dermatitis Disturbance In Attention Dizziness Electrocardiogram Abnormal Feeling Jittery Flushing Movement Disorder Palpitations

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Psychomotor Hyperactivity Tremor	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Health	Zyban	PS		ORAL
150 MG/TWICE			Professional				
PER DAY/ORAL				Oestradiol	C		
				Ascorbic Acid	C		
				Vitamin E	C		

Date:12/18/98ISR Number: 3171870-2Report Type:Expedited (15-DaCompany Report #A0074822  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Foreign	Zyban	PS		ORAL
Life-Threatening		Angina Pectoris	Health				
ORAL			Professional				
Hospitalization -		Chest Pain					
Initial or Prolonged		Myocardial Infarction					
Other							

Date:12/18/98ISR Number: 3171878-7Report Type:Expedited (15-DaCompany Report #A0073094  
Age:15 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin	PS		ORAL
Hospitalization -		Convulsion	Professional				
150 MG ORAL;			Company				
Initial or Prolonged		Fall					
CONTROLLED							
RELEASE		Musculoskeletal Stiffness					
		Overdose	Representative				
		Pyrexia					

Date:12/18/98ISR Number: 3171879-9Report Type:Expedited (15-DaCompany Report #A0076551  
Age:23 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Arthritis	Foreign	Zyban	PS		ORAL
150 MG/TWICE		Dermatitis	Health				
PER DAY/ORAL		Pharyngolaryngeal Pain	Professional				

Date:12/18/98ISR Number: 3171882-9Report Type:Expedited (15-DaCompany Report #A0073660  
Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Angina Pectoris	Foreign	Zyban	PS		ORAL
150 MG /TWICE		Chest Pain	Consumer				
Initial or Prolonged		Tremor		Nitroglycerin	C		
PER DAY/ORAL				Aspirin	C		
				Didrocal	C		
				Tamoxifen	C		
				Amlodipine	C		
				Metoprolol	C		

Date:12/18/98ISR Number: 3171886-6Report Type:Expedited (15-DaCompany Report #A0076883  
Age: Gender:Unknown I/FU:I

Outcome	PT
Hospitalization -	Dehydration
Initial or Prolonged	Headache
	Nausea



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Freedom Of Information (FOI) Report

Vomiting

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/SEE		Consumer	Zyban	PS		ORAL
TEXT/ORAL						

Date:12/18/98ISR Number: 3171888-XReport Type:Expedited (15-DaCompany Report #A0077049  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 250 MG/ORAL		Dyspnoea	Foreign	Wellbutrin	PS		ORAL
Initial or Prolonged Other		Hypersensitivity Lobar Pneumonia	Consumer	Lithium Salt Clomipramine Hcl Clonazepam	C C C		

Date:12/21/98ISR Number: 3173160-0Report Type:Direct Company Report #  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage		Status Epilepticus		Bupropion Divalproex Sodium	PS C		

Date:12/22/98ISR Number: 3173545-2Report Type:Direct Company Report #  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG PO BID		Erythema Multiforme Urticaria		Zyban	PS	Glaxo/Wellcome	ORAL

Date:12/23/98ISR Number: 3227168-7Report Type:Periodic Company Report #9816588  
Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams	Consumer	Zithromax Capsules	PS		ORAL
250.00 MG		Agitation					
TOTAL:DAILY:0		Anxiety					
RAL		Tremor		Zyban	SS		ORAL
ORAL				Alprazolam	SS		ORAL
ORAL				Albuterol	C		
				Atrovent	C		
				Azmacort	C		
				Indocet	C		

Date:12/24/98ISR Number: 3173587-7Report Type:Direct Company Report #  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eye Pain		Zyban	PS		ORAL
300MG PO QD		Face Oedema					
DIVIDED BID	2	Wk					
		Urticaria					

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Freedom Of Information (FOI) Report

Date:12/24/98ISR Number: 3173882-1Report Type:Expedited (15-DaCompany Report #A0077622  
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Foreign	Zyban	PS		ORAL
150 MG /		Cyanosis	Consumer				
TWICE PER DAY		Dermatitis					
/ ORAL		Dyspnoea					
		Eyelid Oedema					
		Hypersensitivity					
		Joint Swelling					
		Myalgia					

Date:12/28/98ISR Number: 3175799-5Report Type:Expedited (15-DaCompany Report #A0076883  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dehydration	Consumer	Zyban	PS		ORAL
150 MG/SEE		Headache					
Initial or Prolonged		Vomiting		Carisoprodol	C		
TEXT/ORAL				Conjugated Estrogens	C		

Date:12/28/98ISR Number: 3175827-7Report Type:Expedited (15-DaCompany Report #A0070795  
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Atrial Fibrillation	Health	Zyban	PS		ORAL
150 MG / PER		Chest Pain	Professional				
Hospitalization -		Dizziness	Company	Fluoxetine			
DAY / ORAL		Pulmonary Oedema	Representative	Hydrochloride	C		
Initial or Prolonged				Dalmane	C		
				Alprazolam	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150 MG/TWICE	Arthralgia	Consumer	Wellbutrin	PS		ORAL
Initial or Prolonged PER DAY/ORAL	Difficulty In Walking	Company				
	Face Oedema	Representative				
	Fall					
	Fatigue					
	Feeling Cold					
	Insomnia					
	Joint Stiffness					
	Migraine					
	Pain					
	Pain In Extremity					
	Pruritus					
	Rash Erythematous					
	Serum Sickness					
	Stomatitis					
	Urticaria					
	Vomiting					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/28/98 ISR Number: 3175913-1 Report Type:Expedited (15-DaCompany Report #A0077761  
 Age:49 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG /ORAL 5 DAY	Anxiety	Health	Wellbutrin	PS		ORAL
Initial or Prolonged	Depression Drug Ineffective Insomnia Jealous Delusion Paranoia Rebound Effect Suicidal Ideation	Professional	Antihypertensive	C		

Date:12/28/98 ISR Number: 3175914-3 Report Type:Expedited (15-DaCompany Report #A0075249  
 Age:43 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 300 MG / PER DAY / ORAL	Back Pain	Health	Wellbutrin	PS		ORAL
Other	Dysphonia	Professional				
	Fibromyalgia Pain In Extremity Weight Decreased	Company Representative	Lithium Salt	C		

Date:12/28/98 ISR Number: 3175915-5 Report Type:Expedited (15-DaCompany Report #A0072073  
 Age:48 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG /TWICE PER DAY	Chills Hyperhidrosis Hypoaesthesia	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
	Memory Impairment					
	Pyrexia Vasculitis Cerebral		Ranitidine Hydrochloride Lisinopril	C C		

Date:12/28/98ISR Number: 3175962-3Report Type:Expedited (15-DaCompany Report #A0077426

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Overdose	Health	Wellbutrin	PS		ORAL
ORAL			Professional	Fluoxetine Hydrochloride	SS		ORAL
ORAL							

Date:12/28/98ISR Number: 3175964-7Report Type:Expedited (15-DaCompany Report #A0077765

Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY		Anaphylactic Reaction	Health	Wellbutrin	PS		ORAL
ORAL		Dermatitis	Professional				
		Dyspnoea					
		Hypotension					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/28/98ISR Number: 3176034-4Report Type:Expedited (15-DaCompany Report #A0071029  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysphagia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG ORAL		Dyspnoea		Librax	C		
		Foreign Body Trauma					
		Nausea					
		Pharyngolaryngeal Pain					
		Retching					

Date:12/28/98ISR Number: 3176356-7Report Type:Expedited (15-DaCompany Report #A0077049  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cryptogenic Organizing	Foreign	Wellbutrin			
Initial or Prolonged		Pneumonia	Consumer	Tablet-Controlled			
Other		Drug Hypersensitivity		Release	PS		ORAL
250 MG/ORAL		Dyspnoea		Lithium Salt	C		
		Lobar Pneumonia		Clomipramine Hcl	C		
				Clonazepam	C		

Date:12/28/98ISR Number: 3176532-3Report Type:Expedited (15-DaCompany Report #A0077928  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Pancreatitis	Foreign	Zyban	PS		ORAL
150 MG TWICE							
Initial or Prolonged			Health				
PER DAY ORAL			Professional				

Date:12/28/98ISR Number: 3176533-5Report Type:Expedited (15-DaCompany Report #A0077971  
Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysarthria	Consumer	Zyban	PS		ORAL
150 MG TWICE		Dyspnoea					
PER DAY ORAL		Erythema		Lorazepam	C		
		Hypersensitivity		Temazepam	C		
		Hypertension		Amitriptyline Hcl	C		
		Tongue Oedema					

Date:12/28/98ISR Number: 3176534-7Report Type:Expedited (15-DaCompany Report #A0077674

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Health	Zyban	PS		ORAL
ORAL			Professional				

Date:12/31/98ISR Number: 3176184-2Report Type:Direct

Company Report #

Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Aggression		Zyban	PS		
1TABLET 2 X		Agitation					
Hospitalization -		Psychotic Disorder					
DAY 21 DAY		Suicidal Ideation					
Initial or Prolonged							
Other							

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Date:12/31/98ISR Number: 3176190-8Report Type:Direct  
Age:40 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300MG PO QD		Eye Pain		Zyban	PS		ORAL
DIVIDED BID	2 WK	Face Oedema Urticaria					

Date:12/31/98ISR Number: 3177016-9Report Type:Expedited (15-DaCompany Report #A0077426  
Age:22 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 100 MG TWICE PER DAY ORAL		Overdose	Health Professional	Wellbutrin Tablet- Controlled Release	PS		ORAL
PER DAY ORAL				Fluoxetine Hydrochloride (Formulation Unknown)	SS		ORAL

Date:12/31/98ISR Number: 3177019-4Report Type:Expedited (15-DaCompany Report #A0072073  
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG TWICE PER DAY ORAL		Back Pain Chills Hyperhidrosis Hypoaesthesia Pyrexia Vasculitis Cerebral	Health Professional	Wellburtin Tablet - Controlled Release	PS		ORAL
				Ranitidine Hydrochloride Lisinopril	C C		

Date:12/31/98ISR Number: 3177021-2Report Type:Expedited (15-DaCompany Report #A0076538  
Age:39 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Dermatitis	Health	Wellbutrin Tablet	PS		ORAL
ORAL 1 YR						
Initial or Prolonged	Drug Interaction	Professional	Diphenhydramine Hcl			
Other	Pyrexia	Company	Capsule	SS		ORAL
ORAL		Representative	Co-Trimoxazole	C		

Date:12/31/98ISR Number: 3177509-4Report Type:Expedited (15-DaCompany Report #A0078087  
Age:63 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Facial Bones Fracture	Foreign	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE						
Initial or Prolonged	Fall	Consumer				
PER DAY/ORAL						
	Haemorrhagic Stroke					
	Headache					

Date:01/05/99ISR Number: 3176747-4Report Type:Direct Company Report #  
Age:38 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Convulsion		Wellbutrin 300 Mg			
Initial or Prolonged			Bid	PS		
			Allergro D Qd	SS		

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Freedom Of Information (FOI) Report

Date:01/06/99ISR Number: 3177851-7Report Type:Expedited (15-DaCompany Report #A0077049  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 250MG ORAL		Cryptogenic Organizing	Foreign	Wellbutrin	PS		ORAL
Initial or Prolonged Other		Pneumonia Dyspnoea Hypersensitivity Pneumonia	Consumer	Lithium Salt Clomipramine Hcl Clonazepam	C C C		

Date:01/07/99ISR Number: 3177983-3Report Type:Direct Company Report #  
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other TWICE A DAY		Confusional State Fall Grand Mal Convulsion		Wellbutrin	PS		

Date:01/07/99ISR Number: 3178754-4Report Type:Direct Company Report #  
Age:50 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG TID & PO >		Hypoaesthesia Loss Of Consciousness Muscle Twitching Paraesthesia Syncope		Bupropion	PS		ORAL

Date:01/07/99ISR Number: 3376494-XReport Type:Periodic Company Report #SYM980001  
Age:10 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

PO	Convulsion	Health	Symmetrel	PS	ORAL
	Corneal Oedema	Professional	Wellbutrin	SS	

Date:01/08/99ISR Number: 3178923-3Report Type:Expedited (15-DaCompany Report #9844878  
 Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL	Alcohol Poisoning	Health	Glucotrol Tablets	PS		ORAL
Initial or Prolonged ORAL	Alcohol Withdrawal Syndrome	Professional	Bupropion Ethanol	SS SS		ORAL ORAL
ORAL	Grand Mal Convulsion Suicide Attempt		Glucophage	C		

Date:01/11/99ISR Number: 3179377-3Report Type:Expedited (15-DaCompany Report #A0078500  
 Age:60 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY / ORAL	Fall Syncope Upper Limb Fracture	Consumer	Zyban	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/11/99ISR Number: 3179532-2Report Type:Expedited (15-DaCompany Report #LBID002990002

Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 600 MG, PER	Agitation	Consumer	Lithobid	PS		ORAL
Initial or Prolonged DAY, PER ORAL	Drug Level Above Therapeutic		Zyprexa	SS		ORAL
PER ORAL	Feeling Drunk		Benzotropine	SS		ORAL
PER ORAL	Nausea Tremor		Wellbutrin Neurontin	SS SS		ORAL
PER ORAL			Klonopin	SS		ORAL
PER ORAL			Ambien	SS		ORAL
PER ORAL			Zoloft	SS		ORAL
600 MG, PER DAY, PER ORAL			Lithobid	SS		ORAL

Date:01/11/99ISR Number: 3179695-9Report Type:Expedited (15-DaCompany Report #A0074755

Age:32 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 9 G/ SINGLE	Aggression	Literature	Wellbutrin Tablet	PS		ORAL
Initial or Prolonged DOSE/ ORAL	Agitation Delirium Electrocardiogram Qrs Complex Prolonged Feeling Jittery Grand Mal Convulsion Overdose Sinus Tachycardia Supraventricular Tachycardia					

Date:01/11/99ISR Number: 3192589-8Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #1998006516-1

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Convulsion	Consumer	Nicoderm Cq 21 Mg Smithkline Beecham Consumer Healthcare	PS		
TRANSDERMAL	21 MILLIGRAMS						
1.0 DAILY							
TRANSDERMAL							
ORAL				Zyban (Cupropion) Glaxo Wellcome	SS		ORAL

Date:01/12/99ISR Number: 3370259-0Report Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #WAES 98111411

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation	Consumer	Tab Propecia 1 Mg	PS		ORAL
PO		Anxiety		Wellbutrin	SS		ORAL
PO		Mental Disorder		Ativan Seroquel Synthroid	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/12/99ISR Number: 3382240-6Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #1998015300-1

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased Depression	Consumer	Nicoderm Cq 21 Mg Smithkline Beecham Consumer Healthcare	PS	Smithkline Beecham Consumer Healthcare	
TRANSDERMAL	TRANSDERMAL			Wellbutrin (Bupropion) Glaxo Wellcome	SS	Glaxo Wellcome	
				Prozac (Fluoxetine) Dista	SS	Dista	
				Albuterol	C		
				Librium (Chlordiazepoxide)	C		
				Rhinocort (Budesonide)	C		
				Aerobid (Flunisolide)	C		

Date:01/14/99ISR Number: 3180952-0Report Type:Direct  
Age:50 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - TABLET		Medication Error	Health	Wellbutrin	PS	Bw Glaxo	ORAL
Initial or Prolonged TABLET			Professional	Buspar	SS	Mead Johnson	ORAL

Date:01/19/99ISR Number: 3181867-4Report Type:Direct  
Age:78 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Bupropion	PS		
Other				Diazepam	SS		

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Retinal Vein Thrombosis	Health	Zyban	PS		ORAL
ORAL			Professional				

Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Drowning	Health	Wellbutrin	PS		ORAL
150 MG /			Professional				
TWICE PER DAY			Company				
/ ORAL			Representative	Fluoxetine			
				Hydrochloridde	C		
				Perphenazine	C		
				Benztropine Mesylate	C		
				Propranolol			
				Hydrochloride	C		
				Lorazepam	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/19/99ISR Number: 3182156-4Report Type:Expedited (15-DaCompany Report #A0077928

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY  / ORAL	Pancreatitis	Foreign  Health  Professional	Zyban	PS		ORAL

Date:01/20/99ISR Number: 3182658-0Report Type:Expedited (15-DaCompany Report #A0073559

Age:58 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG TWICE  PER DAY ORAL	Asthenia  Blood Chloride Increased  Blood Sodium Increased Chills Cough Dizziness Drug Dependence Flushing Haematocrit Increased Haematuria Haemoglobin Increased Headache Mean Cell Volume Increased Nausea Platelet Count Decreased Protein Total Increased Pyrexia Rash Erythematous Sleep Disorder Urinary Tract Infection Urine Analysis Abnormal Vomiting Weight Increased	Health  Professional	Zyban    Conjugated Estrogens Propranolol Hydrochloride Multivitamin	PS    C  C C		ORAL

Date:01/20/99ISR Number: 3182664-6Report Type:Expedited (15-DaCompany Report #A0074622  
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis	Health	Wellbutrin Tablet -			
		Serum Sickness	Professional	Controlled Release	PS		ORAL
150 MG/ TWICE							
		Swelling	Company				
PER DAY/ ORAL 5 DAY		Tooth Abscess	Representative				

Date:01/21/99ISR Number: 3182993-6Report Type:Direct Company Report #  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspepsia	Health	Zyban	PS		
2 WK							
		Rash Pruritic	Professional	Prilosec	C		
		Throat Tightness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/21/99ISR Number: 3183279-6Report Type:Direct  
Age:61 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150MG PO BID	Disorientation		Zyban	PS		ORAL
Hospitalization - Initial or Prolonged	Grand Mal Convulsion					

Date:01/21/99ISR Number: 3183993-2Report Type:Expedited (15-DaCompany Report #9821939  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 25.00 MG	Abdominal Pain	Health	Zoloft	PS		ORAL
Hospitalization - TOTAL:DAILY:0 Initial or Prolonged RAL	Blood Amylase Increased	Professional				
Required 75.00 MG	Drug Ineffective		Wellbutrin	SS		ORAL
Intervention to TOTAL:DAILY:0 Prevent Permanent RAL	Gastroesophageal Reflux Disease					
Impairment/Damage 900.00 MG	Obstruction		Lithane	SS		
TOTAL:DAILY	Pancreatitis Acute		Motrin	C		

Date:01/22/99ISR Number: 3184598-XReport Type:Expedited (15-DaCompany Report #A0079791  
Age:64 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE	Aneurysm	Consumer	Zyban	PS		ORAL
Initial or Prolonged PER DAY/ORAL	Cerebrovascular Accident Hemiplegia		Digoxin	C		

Frusemide C  
Diclofenac Sodium C  
Warfarin Sodium C

Date:01/22/99ISR Number: 3184668-6Report Type:Expedited (15-DaCompany Report #A0077891  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Foreign	Zyban	PS		ORAL
150 MG TWICE		Depressed Level Of	Consumer				
PER DAY ORAL		Consciousness		Clozapine	C		
		Fall		Lorazepam	C		
		Speech Disorder		Ranitidine			
				Hydrochloride	C		

Date:01/22/99ISR Number: 3184756-4Report Type:Expedited (15-DaCompany Report #A0076551  
Age:23 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Arthritis	Foreign	Zyban	PS		ORAL
150 MG /		Dermatitis	Health				
TWICE PER DAY		Pharyngolaryngeal Pain	Professional				
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/22/99 ISR Number: 3184760-6 Report Type:Expedited (15-Da Company Report #A0078500  
 Age:60 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY / ORAL	Fall Forearm Fracture Rib Fracture Syncope	Consumer	Zyban	PS		ORAL

Date:01/22/99 ISR Number: 3193184-7 Report Type:Periodic Company Report #980325-LX278  
 Age:35 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 20.000 MG UNK Initial or Prolonged 350.000 MG QD PO	Coordination Abnormal	Health	Ambien	PS		
100.000 MG QD PO		Professional	Effexor	SS		ORAL
			Wellbutrin	SS		ORAL
			Lithium	C		
			Risperidone	C		
			Thyroid	C		
			Alprazolam	C		

Date:01/25/99 ISR Number: 3185444-0 Report Type:Direct Company Report #  
 Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG BID 3 WK Initial or Prolonged	Convulsion	Health	Bupropion	PS		ORAL
		Professional	Tylenol #4	C		
			Atenolol	C		
			Lipitor (Atrovastin)	C		

Aspirin C  
Cyclobenzamine C

Date:01/25/99ISR Number: 3185551-2Report Type:Expedited (15-DaCompany Report #A0079132  
Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Health	Wellbutrin			
Hospitalization -		Convulsion	Professional	Tablet-Controlled			
Initial or Prolonged		Intentional Misuse	Company	Release	PS		ORAL
150 MG/ SEE							
TEXT/ ORAL		Loss Of Consciousness	Representative				
				Paroxetine			
				Hydrochloride	C		

Date:01/25/99ISR Number: 3185553-6Report Type:Expedited (15-DaCompany Report #A0078231  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Cerebrovascular Accident	Foreign	Zyban	PS		ORAL
150 MG /							
Initial or Prolonged		Fatigue	Health				
TWICE PER DAY							
/ ORAL		Insomnia	Professional				
		Tremor		Human Int/Long			
				Insulin	C		
				Atorvastatin Calcium	C		
				Fenofibrate	C		
				Enalapril Maleate	C		
				Human Short-Act.			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Insulin C

Date:01/25/99ISR Number: 3185628-1Report Type:Expedited (15-DaCompany Report #A0076146  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG / TWICE PER DAY / ORAL 150 MG / TWICE PER DAY / ORAL		Aggression Agitation Anger Anxiety Chromaturia Confusional State Disorientation Emotional Disorder Feeling Abnormal Irritability Pollakiuria Urine Odour Abnormal	Health Professional	Wellbutrin          Zyban	PS          SS		ORAL          ORAL

Date:01/25/99ISR Number: 3185677-3Report Type:Expedited (15-DaCompany Report #A0080209  
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL Other ORAL		Agitation Alcohol Interaction Confusional State Delusion Hostility Insomnia Paranoia	Foreign Health Professional	Zyban   Ethanol Liquid	PS   SS		ORAL       ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Health	Viagra Tablets	PS		ORAL
50.00MG TOTAL		Drug Interaction	Professional				
ORAL				Wellbutrin	SS		ORAL
ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/TWICE		Aggression	Consumer	Zyban	PS		ORAL
PER DAY/ ORAL		Anxiety					
		Convulsion		Percocet	C		
		Dry Mouth		Diazepam	C		
		Haematemesis		Quinalbarbitone			
		Insomnia		Sodium	C		
		Tremor					
		Vision Blurred					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3189484-7Report Type:Expedited (15-DaCompany Report #A0073975  
 Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100 MG/ PER DAY / ORAL		Toxic Epidermal Necrolysis	Health Professional	Wellbutrin	PS		ORAL
200 MG / TWICE PER DAY / ORAL				Lamictal	SS		ORAL

Date:02/01/99ISR Number: 3189507-5Report Type:Expedited (15-DaCompany Report #A0080414  
 Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL		Serum Sickness Urticaria	Foreign Consumer	Zyban	PS		ORAL

Date:02/01/99ISR Number: 3189508-7Report Type:Expedited (15-DaCompany Report #A0071029  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG/ORAL		Choking Dysphagia Dyspnoea Foreign Body Trauma Nausea Oral Intake Reduced Pharyngolaryngeal Pain Retching	Health Professional	Zyban Librax	PS C		ORAL

Date:02/01/99ISR Number: 3189705-0Report Type:Expedited (15-DaCompany Report #A0071893  
Age:43 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150 MG/PER Other DAY/ORAL	Brain Oedema Cerebellar Infarction Cerebrovascular Accident Embolism	Health Professional	Wellbutrin Loratadine	PS C		ORAL

Date:02/01/99ISR Number: 3189884-5Report Type:Expedited (15-DaCompany Report #A0077761  
Age:49 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Other 150 MG / ORAL 5 DAY	Anxiety Depression Drug Ineffective Insomnia Jealous Delusion Paranoia Suicidal Ideation	Health Professional	Wellbutrin Tablet-Controlled Release Antihypertensive Zestoretic	PS C C		ORAL

Date:02/01/99ISR Number: 3196512-1Report Type:Periodic Company Report #A0067641  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source
Other	Convulsion	Health Professional

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Company Representative	Product	Role	Manufacturer	Route
			Wellbutrin Tablet	PS		
Date:02/01/99ISR Number: 3196513-3Report Type:Periodic Company Report #A0067421						
Age:12 YR Gender:Female I/FU:I						
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose			Health	Wellbutrin Tablet	PS	ORAL
Other		Convulsion				
75 MG/TWICE		Headache	Professional			
PER DAY/ ORAL		Loss Of Consciousness		Salbutamol Sulphate	C	
		Mental Impairment				
Date:02/01/99ISR Number: 3196514-5Report Type:Periodic Company Report #A0066609						
Age: Gender: I/FU:I						
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose			Health	Wellbutrin Tablet	PS	ORAL
Other		Convulsion				
ORAL			Professional			
			Company Representative			
Date:02/01/99ISR Number: 3196515-7Report Type:Periodic Company Report #A0066320						
Age:40 YR Gender:Female I/FU:I						
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose			Health	Wellbutrin Tablet -		
Other		Dyspnoea	Professional	Zyban	PS	ORAL
150 MG /		Rash Erythematous				
TWICE PER		Urticaria				
DAY/ ORAL						

Date:02/01/99ISR Number: 3196516-9Report Type:Periodic Company Report #A0066185  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Tablet	PS		ORAL
ORAL			Professional Company Representative				

Date:02/01/99ISR Number: 3196517-0Report Type:Periodic Company Report #A0065523  
Age:9 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia Joint Swelling	Health Professional	Wellbutrin Tablet - Controlled Release	PS		ORAL
100 MG /		Pruritus					
TWICE PER DAY		Serum Sickness					
/ ORAL		Skin Lesion		Desmopression	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3196518-2Report Type:Periodic  
Age:10 YR Gender:Male I/FU:I

Company Report #A0065371

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	100 MG/ THREE	Grand Mal Convulsion	Health	Wellbutrin Tablet	PS		ORAL
	TIMES PER		Professional				
	DAY/ ORAL						

Date:02/01/99ISR Number: 3196519-4Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #A0065219

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	75 MG / PER	Arrhythmia	Consumer	Wellbutrin Tablet	PS		ORAL
Initial or Prolonged	DAY / ORAL	Chest Discomfort					
		Flushing		Maxzide	C		
		Tachycardia		Multivitamin	C		
				Calcium Salt	C		
				Salbutamol Sulphate	C		
				Aspirin	C		
				Lorazepam	C		

Date:02/01/99ISR Number: 3196520-0Report Type:Periodic  
Age:41 YR Gender:Male I/FU:I

Company Report #A0064535

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG /	Pruritus	Health	Wellbutrin Tablet -			
Initial or Prolonged	Other	Urticaria	Professional	Zyban	PS		ORAL
	TWICE PER DAY		Company				
	/ ORAL		Representative				

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL				Hyoscine (Formulation Unknown)	SS		
TRANSDERMAL	TRANSDERMAL						

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Health Professional	Wellbutrin Tablet	PS		ORAL
100 MG /		Drug Hypersensitivity					
THREE TIMES		Dysphoria					
PER DAY/ ORAL		Insomnia		Lorazepam	C		
		Pruritus					
		Rash Maculo-Papular					
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3196523-6Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #A0063000

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG / PER DAY / ORAL		Dermatitis Face Oedema Rash Erythematous Throat Tightness	Health Professional	Wellbutrin Tablet - Zyban  Nicotine	PS		ORAL   C

Date:02/01/99ISR Number: 3196524-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0063037

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose  ORAL		Face Oedema Periorbital Oedema	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3196525-XReport Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0062802

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150 MG / TWICE PER DAY / ORAL		Asthenia Difficulty In Walking Dyskinesia Dysphagia Dyspnoea Fatigue Hyperventilation Jaw Disorder Muscle Rigidity Nervous System Disorder Speech Disorder Tremor	Health Professional	Wellbutrin Tablet - Zyban     Medroxyprogesterone Ace.	PS		ORAL     C

Date:02/01/99ISR Number: 3196527-3Report Type:Periodic Company Report #A0063358  
Age:36 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY	Auricular Swelling	Health	Wellbutrin	PS		ORAL
Other / ORAL	Erythema Multiforme	Professional				
	Gastric Dilatation					
	Malaise		Allergy Medication	C		
	Nausea					
	Oedema Peripheral					
	Pain					
	Pharyngeal Oedema					
	Pyrexia					
	Rash Erythematous					
	Rash Macular					
	Stevens-Johnson Syndrome					
	Urticaria					

Date:02/01/99ISR Number: 3196528-5Report Type:Periodic Company Report #A0063149  
Age:35 YR Gender:Male I/FU:I

Outcome	PT	Report Source
Hospitalization - Initial or Prolonged	Coordination Abnormal	Health
		Professional

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Freedom Of Information (FOI) Report

Other

Dose	Duration	Product	Role	Manufacturer	Route
VARIABLE DOSE		Wellbutrin	PS		ORAL
/ ORAL					
350 MG / PER		Venlafaxine Hydrochloride Tablet	SS		ORAL
DAY / ORAL					
20 MG		Zolpidem Tartrate (Formulation Unknown)	SS		
		Lithium Salt	C		
		Risperidone	C		
		Thyroid	C		
		Alprazolam	C		

Date:02/01/99ISR Number: 3196529-7Report Type:Periodic Company Report #A0061149  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Tablet	PS		ORAL
150 MG /			Professional				
TWICE PER DAY			Company				
/ ORAL			Representative				

Date:02/01/99ISR Number: 3196530-3Report Type:Periodic Company Report #A0059271  
 Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization -		Abdominal Pain Angioneurotic Oedema	Health Professional	Wellbutrin Tablet - Zyban	PS		ORAL
150 MG / ORAL							

Initial or Prolonged	Face Oedema	Company
Other	Headache	Representative
	Nausea	
	Pharyngeal Oedema	
	Urticaria	
	Vomiting	

Date:02/01/99ISR Number: 3196531-5Report Type:Periodic Company Report #A0059079  
 Age:6 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY  / ORAL	Grand Mal Convulsion	Health  Professional	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3196532-7Report Type:Periodic Company Report #A0072937  
 Age:52 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 100 MG /  THREE TIMES  PER DAY ORAL	Conversion Disorder  Convulsion  Crying  Depression Drug Withdrawal Syndrome Headache Nausea	Health  Professional	Wellbutrin Tablet    Aspirin	PS    C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3196533-9Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0072356

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Tablet	PS		ORAL
Other		Anxiety					
100 MG THREE		Clonic Convulsion	Professional				
TIMES PER DAY		Dystonia	Company				
ORAL		Hyperreflexia	Representative	Triamcinolone			
		Tardive Dyskinesia		Acetonide	C		
		Tremor		Melatonin	C		

Date:02/01/99ISR Number: 3196534-0Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #A0071371

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Tablet	PS		ORAL
Other		Pruritus					
ORAL	3 WK	Urticaria	Professional	Wellbutrin Tablet -			
			Company	Controlled Release	SS		ORAL
ORAL			Representative	Nitrofurantoin	C		

Date:02/01/99ISR Number: 3196535-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0071498

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Wellbutrin	PS		ORAL
Other		Convulsion					
ORAL							

Date:02/01/99ISR Number: 3196536-4Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #A0071924

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 150 MG (TWICE Initial or Prolonged PER DAY ORAL) Other	Angioneurotic Oedema Chest Pain Urticaria	Health Professional Company Representative	Wellbutrin	PS	ORAL
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Date:02/01/99ISR Number: 3196537-6Report Type:Periodic Company Report #A0070647  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG THREE TIMES PER DAY ORAL		Convulsion Loss Of Consciousness	Health Professional	Wellbutrin Tablet	PS		ORAL
				Fluoxetine Hydrochloride Thyroxine Sodium Nortriptyline	C C C		

Date:02/01/99ISR Number: 3196538-8Report Type:Periodic Company Report #A0070221  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG / TWICE PER DAY / ORAL		Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Tablet Narcotic	PS C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3196539-XReport Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0070222

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG TWICE PER DAY / ORAL		Convulsion	Health Professional Company Representative	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3196540-6Report Type:Periodic  
 Age:41 YR Gender:Male I/FU:I

Company Report #A0070053

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Convulsion	Health Professional Other	Wellbutrin Tablet Paroxetine Hydrochloride Tablet	PS SS		ORAL ORAL
				Colchicine Allopurinol Temazepam	C C C		

Date:02/01/99ISR Number: 3196541-8Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:I

Company Report #A0070058

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG / TWICE PER DAY / ORAL	1 WK	Convulsion	Health Professional Company Representative	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3196542-XReport Type:Periodic Company Report #A0069723  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Grand Mal Convulsion	Health	Wellbutrin Tablet	PS		ORAL
150 MG / PER			Professional				
Other			Company				
DAY / ORAL			Representative				

Date:02/01/99ISR Number: 3196543-1Report Type:Periodic Company Report #A0069889  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Health	Wellbutrin Tablet	PS		ORAL
150 MG ORAL	1 MON		Professional				
			Company				
			Representative				

Date:02/01/99ISR Number: 3196544-3Report Type:Periodic Company Report #A0069441  
Age:27 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Chest Pain
Initial or Prolonged	Convulsion
Other	Depressed Level Of Consciousness Disorientation Overdose



Other Grand Mal Convulsion Consumer Wellbutrin Tablet PS ORAL  
PER DAY ORAL  
Nortriptyline C

Date:02/01/99ISR Number: 3196548-0Report Type:Periodic Company Report #A0068887  
Age:35 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1200 MG / Initial or Prolonged SINGLE DOSE / ORAL	Bundle Branch Block Right Convulsion Electrocardiogram Qrs Complex Prolonged Sinus Tachycardia	Health Professional	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3196549-2Report Type:Periodic Company Report #A0068803  
Age:50 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG / TWICE PER DAY / ORAL	Convulsion Feeling Abnormal	Health Professional	Wellbutrin Tablet - Controlled Release	PS		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3196550-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0068334

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Tablet	PS		ORAL
150 MG /			Professional				
THREE TIMES							
PER DAY /							
ORAL							

Date:02/01/99ISR Number: 3196551-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0068155

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Health	Wellbutrin	PS		ORAL
ORAL			Professional				
			Company				
			Representative				

Date:02/01/99ISR Number: 3197253-7Report Type:Periodic  
Age:12 YR Gender:Female I/FU:I

Company Report #7397306

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaphylactic Reaction	Health	Abbott-Cylert	PS	Abbott	ORAL
75.000 MG PO			Professional				
QD							
		Face Oedema		Wellbutrin Sr	SS		ORAL
150.000 MG							
BID							

Date:02/01/99ISR Number: 3198313-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0069125

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Tablet	PS		ORAL
300 MG/ UNK/		Overdose	Professional				
ORAL			Company Representative				

Date:02/01/99ISR Number: 3198314-9Report Type:Periodic Company Report #A0077763  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspnoea	Consumer	Wellbutrin Tablet	PS		ORAL
150 MG /TWICE		Pharyngeal Oedema					
PER DAY /ORAL		Urticaria		Melatonin	C		
				Buspirone			
				Hydrochloride	C		

Date:02/01/99ISR Number: 3198315-0Report Type:Periodic Company Report #A0077656  
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Tablet	PS		ORAL
150 MG /TWICE			Professional				
PER DAY /			Company				
ORAL			Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3198316-2Report Type:Periodic  
 Age:25 YR Gender:Male I/FU:I

Company Report #A0077409

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY  / ORAL / 7  DAYS	Anaphylactic Reaction	Health  Professional  Company  Representative	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3198317-4Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:I

Company Report #A0078055

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY  / ORAL	Convulsion	Health  Professional  Company  Representative	Wellbutrin Tablet   Carisoprodol Temazepam Carisoprodol	PS   C C C		ORAL

Date:02/01/99ISR Number: 3198318-6Report Type:Periodic  
 Age:30 YR Gender:Male I/FU:I

Company Report #A0077917

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other UNK/ UNK /  ORAL	Grand Mal Convulsion	Health  Professional  Company  Representative	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3198319-8Report Type:Periodic Company Report #A0076229  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State Grand Mal Convulsion Loss Of Consciousness	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /							
TWICE PER DAY							
/ORAL							
				Thyroxine Sodium	C		

Date:02/01/99ISR Number: 3198320-4Report Type:Periodic Company Report #A0067837  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Health	Wellbutrin Tablet	PS		ORAL
Life-Threatening							
150 MG /							
Hospitalization -							
TWICE PER DAY							
Initial or Prolonged							
/ ORAL							
Other							

Date:02/01/99ISR Number: 3198321-6Report Type:Periodic Company Report #A0071371  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pruritus	Health	Wellbutrin Tablet	PS		ORAL
UNK / UNK /							
ORAL / 3							
WEEKS							
			Company				
			Representative	Wellbutrin			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

UNK / UNK /  
 ORAL  
 Tablet-Controlled Release SS ORAL  
 Nitrofurantoin C

Date:02/01/99ISR Number: 3198322-8Report Type:Periodic Company Report #A0071495  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Tablet	PS		ORAL
Other		Convulsion	Professional				
150 MG /		Loss Of Consciousness					
TWICE PER DAY		Urinary Incontinence					
/ ORAL				Ortho Tri-Cyclen	C		

Date:02/01/99ISR Number: 3198323-XReport Type:Periodic Company Report #A0071644  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Tablet	PS		ORAL
Other		Electroencephalogram	Professional				
200 MG /		Abnormal					
TWICE PER DAY		Grand Mal Convulsion	Company Representative	Venlafaxine Hydrochloride	C		
/ ORAL				Olanzapine	C		
				Sertraline			
				Hydrochloride	C		
				Carbamazepine	C		
				Semisodium Valproate	C		

Date:02/01/99ISR Number: 3198324-1Report Type:Periodic Company Report #A0072757  
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Intestinal Perforation	Health	Wellbutrin Tablet	PS		ORAL
100 MG / UNK			Professional				
/ ORAL				Conjugated Estrogens	C		
				Fiorinal	C		

Date:02/01/99ISR Number: 3198325-3Report Type:Periodic Company Report #A0073013  
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Tablet	PS		ORAL
UNK/ UNK /			Professional				
ORAL			Company Representative				

Date:02/01/99ISR Number: 3198326-5Report Type:Periodic Company Report #A0073098  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Urticaria	Health	Wellbutrin Tablet	PS		ORAL
150 MG / UNK			Professional				
Initial or Prolonged			Company Representative				
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3198327-7Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #A0073332

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Health	Wellbutrin Tablet	PS		ORAL
150 MG /		Insomnia	Professional				
TWICE PER DAY			Company				
/ ORAL			Representative	Amitriptyline Hcl Tablet	SS		ORAL
25 MG / UNK /							
ORAL							

Date:02/01/99ISR Number: 3198328-9Report Type:Periodic  
Age:10 YR Gender:Unknown I/FU:I

Company Report #A0073334

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Tablet	PS		ORAL
150 MG /		Staring	Professional				
TWICE PER DAY			Company				
/ ORAL			Representative				

Date:02/01/99ISR Number: 3198329-0Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0073418

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bite	Health	Wellbutrin Tablet	PS		ORAL
UNK / SEE		Depression	Professional				
TEXT / ORAL		Grand Mal Convulsion		Venlafaxine Hydrochloride Olanzapine	C C		

Date:02/01/99ISR Number: 3198330-7Report Type:Periodic Company Report #A0073628  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Tablet	PS		ORAL
150 MG /		Electroencephalogram	Professional				
TWICE PER DAY		Abnormal	Company				
/ORAL		Fall	Representative				
		Loss Of Consciousness					
		Muscle Spasms					
		Snoring					

Date:02/01/99ISR Number: 3198331-9Report Type:Periodic Company Report #A0073629  
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Tablet	PS		ORAL
150 MG / PER			Professional				
DAY / ORAL			Company	Meclofenamate Sodium	C		
			Representative	Entex	C		
				Unknown	C		



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Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3198332-0Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #A0074135

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY  / ORAL	Depressed Level Of Consciousness  Grand Mal Convulsion	Consumer	Wellbutrin Tablet	PS		ORAL

Paracetamol C

Date:02/01/99ISR Number: 3295673-3Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #A0076853

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration  150 MG ORAL	Nausea	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:02/01/99ISR Number: 3295674-5Report Type:Periodic  
Age:77 YR Gender:Female I/FU:I

Company Report #A0076854

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration  150 MG TWICE  PER DAY ORAL	Tremor	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

Ascorbid Acid C  
Nitroglycerin C  
Calcium Salt C  
Aspirin C  
Vitamin E C  
Chromium Picolinate C  
Insulin C

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER		Chest Discomfort Dry Mouth	Consumer	Wellbutrin Tablet -Controlled Release	PS		ORAL
DAY ORAL		Dyspnoea					
400 MG PER		Fatigue		Raxar Tablet	SS		ORAL
DAY ORAL		Insomnia					
150 MG PER		Tension Headache		Zyban Tablet-Zyban	SS		ORAL
DAY ORAL		Urinary Incontinence					
				Wellbutrin (Formulation Unknown) Loratadine	SS C		

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER		Agitation Insomnia	Consumer	Wellbutrin Tablet -Controlled Release	PS		ORAL
DAY ORAL		Tremor					
				Donnatal	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3295677-0Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #A0079236

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Accommodation Disorder Disturbance In Attention	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG TWICE							
DAILY ORAL				Nafazodone Hydrochloride Ginseng	C C		

Date:02/01/99ISR Number: 3295678-2Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0059262

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oedema Urticaria	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3295679-4Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0066438

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Myalgia	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3295680-0Report Type:Periodic  
Age:54 YR Gender:Male I/FU:I

Company Report #A0067314

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Epileptic Aura	Health Professional	Wellbutrin Tablet -Controlled Release	PS		ORAL
150 MG TWICE							
PER DAY ORAL							

Paroxetine Hydrochloride	C		
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Date:02/01/99ISR Number: 3295681-2Report Type:Periodic Company Report #A0067487  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health Professional	Wellbutrin Tablet-Controlled Release	PS		

Date:02/01/99ISR Number: 3295682-4Report Type:Periodic Company Report #A0069807  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Wellbutrin Tablet -Controlled Release	PS		ORAL
1 TABLET PER							
DAY ORAL							

Conjugated Estrogens	C		
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Hyperglycaemia  
Health Professional Wellbutrin  
Tablet-Controlled Release PS ORAL

Date:02/01/99ISR Number: 3296030-6Report Type:Periodic Company Report #A0078019  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia Nausea	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
TWICE PER DAY							
/ ORAL							
					Conjugated Estrogens	C	

Date:02/01/99ISR Number: 3296034-3Report Type:Periodic Company Report #A0078021  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accommodation Disorder Lethargy	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / PER							
DAY / ORAL							
					Amitriptyline	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3296036-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0078249

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Drug Interaction Memory Impairment Tinnitus	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
				Sertraline Hydrochloride (Formulation Unknown)	SS		

Date:02/01/99ISR Number: 3296038-0Report Type:Periodic  
 Age:51 YR Gender:Male I/FU:I

Company Report #A0078462

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Drug Interaction Formication	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
				Lithium Salt	C		
				Insulin	C		
				Salbutamol	C		

Date:02/01/99ISR Number: 3296041-0Report Type:Periodic  
 Age:76 YR Gender:Male I/FU:I

Company Report #A0078463

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Chest Pain Hallucination Vision Blurred	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
				Verapamil	C		

Allopurinol C  
Unknown C  
Potassium Salt C  
Unknown C

Date:02/01/99ISR Number: 3296043-4Report Type:Periodic Company Report #A0078471  
Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus Urticaria	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / TWICE PER DAY / ORAL							

Lithium C  
Zolpidem C  
Gabapentin C

Date:02/01/99ISR Number: 3296047-1Report Type:Periodic Company Report #A0078472  
Age:19 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Twitching	Consumer	Wellbutrin Tablet-Controlled			



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

150 MG /  
 TWICE PER DAY  
 / ORAL

Release PS ORAL

Pirbuterol Acetate C

Date:02/01/99ISR Number: 3296049-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0078473

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Depression	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

150 MG / PER  
 DAY / ORAL

Buspirone C  
 Diazepam C

Date:02/01/99ISR Number: 3296052-5Report Type:Periodic  
 Age:33 YR Gender:Female I/FU:I

Company Report #A0078474

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

150 MG /  
 TWICE PER DAY  
 / ORAL

Claritin-D C  
 Diphenhydramine Hcl C  
 Birth Control C  
 Paroxetine C  
 Ethanol C

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia Insomnia Irritability	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
TWICE PER DAY							
/ ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
100 MG /							
TWICE PER DAY							
/ ORAL							
				Semisodium	C		
				Olanzapine	C		
				Liothyronine	C		
				Lamotrigine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3296059-8Report Type:Periodic  
 Age: Gender:Unknown I/FU:F

Company Report #A0058749

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Acarodermatitis Pruritus	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL

ORAL

Date:02/01/99ISR Number: 3296067-7Report Type:Periodic  
 Age:41 YR Gender:Male I/FU:F

Company Report #A0070348

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG / THREE TIMES PER DAY / ORAL		Blister Chills Flushing  Hyperhidrosis  Rash Erythematous  Rash Pruritic  Rhinorrhoea Skin Lesion	Health Professional Company  Representative	Wellbutrin Tablet-Controlled Release      Diphenhydramine	PS      C		ORAL

Date:02/01/99ISR Number: 3296068-9Report Type:Periodic  
 Age:65 YR Gender:Female I/FU:F

Company Report #A0071172

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Abdominal Distension Diarrhoea Muscle Spasms	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:02/01/99ISR Number: 3296072-0Report Type:Periodic Company Report #A0071449  
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus Rash Erythematous	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / PER DAY / ORAL							
				Nicotine Ketoconazole	C C		

Date:02/01/99ISR Number: 3296076-8Report Type:Periodic Company Report #A0071646  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chills	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / PER DAY / ORAL							

Date:02/01/99ISR Number: 3296091-4Report Type:Periodic Company Report #A0076958  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
150 MG / TWICE PER DAY							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

/ ORAL

Methadone C  
 Buspirone  
 Hydrochloride C  
 Paroxetine  
 Hydrochloride C  
 Hydrochlorothiazide C  
 Oxaprozin C

Date:02/01/99ISR Number: 3296092-6Report Type:Periodic Company Report #A0076960  
 Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anorgasmia Insomnia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /							
TWICE PER DAY							

/ ORAL

Dalmane C  
 Hydrochlorothiazide C

Date:02/01/99ISR Number: 3296093-8Report Type:Periodic Company Report #A0076961  
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation Psychomotor Hyperactivity	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / ORAL							

Date:02/01/99ISR Number: 3296097-5Report Type:Periodic Company Report #A0077000  
 Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Feeling Abnormal	Consumer	Wellbutrin			

Hyperhidrosis  
Nervousness

Tablet-Controlled  
Release

PS

ORAL

150 MG / PER

DAY / ORAL

Date:02/01/99ISR Number: 3296098-7Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #A0077069

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State Depression	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
PER DAY /		Disturbance In Attention					
ORAL		Increased Appetite					

Date:02/01/99ISR Number: 3296102-6Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0077070

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Urinary Tract Infection	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
TWICE PER DAY							
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3296103-8Report Type:Periodic  
 Age:24 YR Gender:Male I/FU:I

Company Report #A0077080

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Paroxetine Hydrochloride	C
Temazepam	C

Date:02/01/99ISR Number: 3296106-3Report Type:Periodic  
 Age:15 YR Gender:Female I/FU:I

Company Report #A0077095

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eye Pain Eyelid Oedema Eyelid Ptosis	Health Professional	Wellbutrin Tablet - Controlled Release	PS		ORAL
100 MG /							
TWICE PER DAY							
/ ORAL							

Date:02/01/99ISR Number: 3296108-7Report Type:Periodic  
 Age:59 YR Gender:Male I/FU:I

Company Report #A0077096

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension Discomfort Flatulence	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / PER							
DAY / ORAL							

Date:02/01/99ISR Number: 3296110-5Report Type:Periodic Company Report #A0077101  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia Oedema Peripheral Pruritus	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /		Swelling					
TWICE PER DAY		Urticaria					
/ORAL							

Date:02/01/99ISR Number: 3296113-0Report Type:Periodic Company Report #A0080314  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
100 MG / ORAL							

Date:02/01/99ISR Number: 3296117-8Report Type:Periodic Company Report #A0077341  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Wellbutrin Tablet- Controlled Release	PS		ORAL
150 MG / PER				Lithium Salt	C		
DAY / ORAL				Prempro	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3296120-8Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0077427

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health Professional	Wellbutrin Tablet - Controlled Release	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3296123-3Report Type:Periodic  
 Age:34 YR Gender:Male I/FU:I

Company Report #A0077452

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Headache	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
150 MG / TWICE PER DAY / ORAL							
ORAL							
				Zyban Tablet - Zyban	SS		ORAL

Date:02/01/99ISR Number: 3296125-7Report Type:Periodic  
 Age:31 YR Gender:Male I/FU:I

Company Report #A0077587

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accommodation Disorder Balance Disorder Disorientation	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / TWICE PER DAY / ORAL							
		Muscle Spasms		Combivir	C		
				Ascriptin	C		
				Dronabinol	C		
				Clonazepam	C		
				Semisodium Valproate	C		
				Amoxicillin	C		

Vicoprofen C  
Venlafaxine  
Hydrochloride C

Date:02/01/99ISR Number: 3296127-0Report Type:Periodic Company Report #A0077670  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
TWICE PER DAY							

/ ORAL

Diuretic C  
Blood Pressure  
Medication C

Date:02/01/99ISR Number: 3296129-4Report Type:Periodic Company Report #A0077671  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Palpitations Tremor	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL

150 MG /

TWICE PER DAY

/ ORAL

Estrogen C  
Progesterone C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lisinopril C  
Folic Acid C

Date:02/01/99ISR Number: 3296131-2Report Type:Periodic Company Report #A0077699  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue Insomnia	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
100 MG / PER DAY / ORAL	4 WK			Clonazepam	C		

Date:02/01/99ISR Number: 3296132-4Report Type:Periodic Company Report #A0077701  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Hypotension	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
150 MG / TWICE PER DAY / ORAL				Crataegus Clonazepam	C C		

Date:02/01/99ISR Number: 3297093-4Report Type:Periodic Company Report #A0067140  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Angioneurotic Oedema	Health	Lamictal Tablet	PS		ORAL
ORAL 5 WK		Dermatitis	Professional	Wellbutrin Tablet	SS		
		Lichen Planus Periorbital Oedema					

Date:02/01/99ISR Number: 3301098-4Report Type:Periodic Company Report #A0067430  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Lamictal Tablet	PS		ORAL
ORAL		Headache		Wellbutrin Tablet	SS		ORAL
ORAL		Nausea Pruritus Rash Papular		Olanzapine	C		

Date:02/01/99ISR Number: 3304217-9Report Type:Periodic Company Report #A0071892  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Herpes Simplex	Consumer	Lamictal Tablet	PS		ORAL
300 MG / PER							
DAY / ORAL	2	MON		Wellbutrin Tablet-Controlled Release	SS		ORAL
ORAL	3	WK		Clomazepam Thyroxide Sodium	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3393841-3Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0074623

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Behaviour Anxiety Crying	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / IN		Depression					
THE MORNING /		Emotional Disorder					
ORAL		Screaming					

Date:02/01/99ISR Number: 3393842-5Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #A0074642

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /				Sertraline Hydrochloride	C		
TWICE PER				Loratadine	C		
DAY/ ORAL				Nedocromil Sodium	C		
				Budesonide	C		
				Vitamin	C		

Date:02/01/99ISR Number: 3393844-9Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0074643

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Nausea	Consumer	Wellbutrin Tablet-Controlled			

150 MG/ THREE

Release

PS

ORAL

TIMES PER DA

/ ORAL

Conjugated Estrogens C

Date:02/01/99ISR Number: 3393846-2Report Type:Periodic

Company Report #A0074644

Age:36 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Sedation	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

150 MG/

TWICER PER

DAY/ ORAL

Insulin C

Date:02/01/99ISR Number: 3393848-6Report Type:Periodic

Company Report #A0074645

Age:48 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Paraesthesia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

150 MG /

TWICE PER DAY

/ ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3393849-8Report Type:Periodic  
 Age:42 YR Gender:Female I/FU:I

Company Report #A0074646

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urticaria	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /							
TWICE PER							
DAY/ ORAL							

Date:02/01/99ISR Number: 3393851-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0074660

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / ORAL	12 DAY						

Date:02/01/99ISR Number: 3393853-XReport Type:Periodic  
 Age:39 YR Gender:Female I/FU:I

Company Report #A0074720

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Pain Agitation Dizziness	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / PER		Drug Interaction					
DAY / ORAL		Hypoaesthesia Mania Syncope		Procaine Hydrochloride (Formulation Unknown)	SS		
				Clonazepam	C		
				Nizatidine	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /			Representative				
TWICE PER DAY							
/ ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /			Representative				
TWICE PER DAY							
/ ORAL				Lorazepam	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3393860-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0074911

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Insomnia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:02/01/99ISR Number: 3393863-2Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #A0074912

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Anxiety Insomnia Oedema	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
1 MG / TWICE PER DAY		Rash Maculo-Papular Tension Urticaria		Clonazepam (Formulation Unknown)	SS		
ORAL				Wellbutrin Tablet-Controlled Release	SS		ORAL

Date:02/01/99ISR Number: 3393866-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0074957

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Galactorrhoea	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:02/01/99ISR Number: 3393868-1Report Type:Periodic  
Age:41 YR Gender:Male I/FU:I

Company Report #A0074981

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Arthralgia Drug Interaction Haematoma	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
7.5 MG /		International Normalised Ratio Increased Joint Swelling	Representative	Warfarin Sodium (Formulation Unknown) Warfarin Sodium (Formulation Unknown)	SS SS		

ALTERNATE  
DAYS

Date:02/01/99ISR Number: 3393905-4Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0073758

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dizziness Headache	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

TWICE PER DAY  
/ ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3393907-8Report Type:Periodic  
 Age:40 YR Gender:Male I/FU:I

Company Report #A0073760

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
TWICE PER							
DAY/ ORAL							

Insulin C

Date:02/01/99ISR Number: 3393908-XReport Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #A0073779

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hair Texture Abnormal Nausea	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / PER							
DAY / ORAL							

Alprazolam C

Date:02/01/99ISR Number: 3393911-XReport Type:Periodic  
 Age:18 YR Gender:Male I/FU:I

Company Report #A0073847

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
1 TABLET/							
TWICE PER							
DAY/ ORAL							

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY /ORAL		Abdominal Pain Back Pain Dermatitis Exfoliative Diarrhoea Disorientation Erythema Oedema Peripheral Onycholysis Vomiting	Consumer	Wellbutrin Tablet-Controlled Release  Estropipate Hydrochlorothiazide Fexofenadine Hydrochlorid Alprazolam Herbal Vitamin	PS    C C  C C C		ORAL

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Gingivitis	Consumer	Wellbutrin Tablet-Controlled Release  Trazodone	PS   C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3395316-4Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0072684

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Deafness	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ PER							
DAY/ ORAL							

Date:02/01/99ISR Number: 3395317-6Report Type:Periodic  
Age:66 YR Gender:Female I/FU:I

Company Report #A0072698

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Mental Impairment Nervousness	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ PER							
DAY/ ORAL				Multivitamin Estrogen	SS C		

Date:02/01/99ISR Number: 3395318-8Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0072750

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Venous Stasis	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL			Representative	Paroxetine Hydrochloride	C		

Date:02/01/99ISR Number: 3395319-XReport Type:Periodic  
Age:33 YR Gender:Male I/FU:I

Company Report #A0074990

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Chromaturia Dizziness Mood Altered	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
1 TABLET/ PER							
DAY/ ORAL				Clonazepam	C		

Date:02/01/99ISR Number: 3395320-6Report Type:Periodic Company Report #A0075002  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Interaction International Normalised Ratio Increased	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ ORAL				Warfarin Sodium (Formulation Unknown)	SS		
5 MG / PER							
DAY							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3395321-8Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0075003

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction International Normalised Ratio Increased	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ ORAL			Representative	Warfarin Sodium (Formulation Unknown)	SS		

Date:02/01/99ISR Number: 3395322-XReport Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0075004

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction International Normalised Ratio Increased	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ ORAL			Representative	Warfarin Sodium (Formulation Unknown)	SS		

Date:02/01/99ISR Number: 3395323-1Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0075006

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Irritability	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL			Representative				

Date:02/01/99ISR Number: 3395324-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0075053

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
300 MG/ PER							
DAY / ORAL							

Date:02/01/99ISR Number: 3395325-5Report Type:Periodic Company Report #A0075054  
 Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Insomnia Testicular Pain	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
100 MG/ THREE							
TIMES PER							
DAY/ ORAL							

Simvastatin C

Date:02/01/99ISR Number: 3395326-7Report Type:Periodic Company Report #A0075057  
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Effect Decreased Malaise	Consumer	Wellbutrin Tablet-Controlled			



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

150 MG/ TWICE				Release	PS		ORAL
PER DAY /							
ORAL				Nicotine	C		
				Alprazolam	C		

Date:02/01/99ISR Number: 3395327-9Report Type:Periodic Company Report #A0074209  
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL				Alprazolam	C		

Date:02/01/99ISR Number: 3395328-0Report Type:Periodic Company Report #A0074210  
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Insomnia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ PER							
DAY/ ORAL				Atorvastatin Calcium	C		
				Tenoretic	C		
				Estropipate	C		

Date:02/01/99ISR Number: 3395329-2Report Type:Periodic Company Report #A0074213  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

	Nausea	Consumer	Wellbutrin Tablet-Controlled Release	PS	ORAL
TWICE PER DAY/ ORAL			Paroxetine Hydrochloride Trazodone Estratest Medroxyprogesterone Ace.	C C C C	

Date:02/01/99ISR Number: 3395330-9Report Type:Periodic Company Report #A0074379  
 Age:21 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG/ PER DAY/ ORAL		Agitation Dizziness Feeling Abnormal  Tremor	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
				Salbutamol Sulphate Fluoxetine Hydrochloride	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3395331-0Report Type:Periodic  
 Age:72 YR Gender:Female I/FU:I

Company Report #A0074380

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia Hyperhidrosis	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / TWICE PER DAY/ ORAL				Guanethidine Monosulphate Glipizide Glucophage Multivitamin	C C C C		

Date:02/01/99ISR Number: 3395332-2Report Type:Periodic  
 Age:10 YR Gender:Female I/FU:I

Company Report #A0074450

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Enanthema Hypersensitivity	Health Professional Other	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ TWICE PER DAY / ORAL	2 WK	Oedema Mouth		Pemoline	C		

Date:02/01/99ISR Number: 3395333-4Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0074504

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Overdose	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3395334-6Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0074576

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea Influenza Like Illness Pain	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
100 MG/ PER DAY/ ORAL				Paroxetine Hydrochloride	SS		

Date:02/01/99ISR Number: 3395335-8Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0074578

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion Drug Interaction	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL	1 YR		Representative	Fluvoxamine Maleate Tablet	SS		
50 MG							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3395336-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0073880

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia Liver Function Test Abnormal	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / THREE TIMES PER DAY		Tinnitus		Wellbutrin Tablet Gabapentin	SS C		

Date:02/01/99ISR Number: 3395337-1Report Type:Periodic  
 Age:32 YR Gender:Male I/FU:I

Company Report #A0073881

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression Dermatitis Mood Swings	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / TWICE PER DAY / ORAL		Nightmare		Pseudoephedrine Hcl Paracetamol	C C		

Date:02/01/99ISR Number: 3395338-3Report Type:Periodic  
 Age:48 YR Gender:Female I/FU:I

Company Report #A0073882

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nausea	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
UNK/ PER DAY/ ORAL							

Semisodium Valproate C  
 Doxepin C  
 Estratest C  
 Potassium Chloride C  
 Atorvastatin Calcium C  
 Prednisone C  
 Thyroxine Sodium C  
 Lansoprazole C  
 Ciprofloxacin Hcl C  
 Compazine C

Date:02/01/99ISR Number: 3395339-5Report Type:Periodic Company Report #A0073883  
 Age:26 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Headache Tachycardia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /		Urticaria					
TWICE PER DAY							
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3395340-1Report Type:Periodic  
 Age:31 YR Gender:Female I/FU:I

Company Report #A0073884

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / PER DAY / ORAL				No Concurrent Medication	C		

Date:02/01/99ISR Number: 3395341-3Report Type:Periodic  
 Age:50 YR Gender:Female I/FU:I

Company Report #A0073932

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Nausea Tremor	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
200 MG / TWICE PER DAY / ORAL		Weight Decreased		Atenolol	C		

Date:02/01/99ISR Number: 3395342-5Report Type:Periodic  
 Age:50 YR Gender:Male I/FU:I

Company Report #A0073933

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Simple Partial Seizures Tremor	Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / TWICE PER DAY / ORAL				Tramadol			

Hydrochloride C  
Arthritis Medication C

Date:02/01/99ISR Number: 3395343-7Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0073937

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disturbance In Attention Headache Loss Of Consciousness Photophobia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Fluoxetine  
Hydrochloride C  
Zafirlukast C  
Sodium Cromoglycate C  
Omeprazole C  
Beclomethasone  
Dipropion C  
Salbutamol Sulphate C  
Fluticasone  
Propionate C  
Metronidazole C



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3395344-9Report Type:Periodic  
 Age:41 YR Gender:Female I/FU:I

Company Report #A0073939

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Wellbutrin Tablet- Controlled Release	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:02/01/99ISR Number: 3395345-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0073940

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
UNK / UNK /							
ORAL							

Date:02/01/99ISR Number: 3395346-2Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #A0073943

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Urticaria	Health Professional	Wellbutrin Tablet - Controlled Release	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Trazodone C

Date:02/01/99ISR Number: 3395347-4Report Type:Periodic  
 Age:41 YR Gender:Female I/FU:I

Company Report #A0073944

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dysgeusia Insomnia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
UNK/ TWICE							
PER DAY /							
ORAL							

Date:02/01/99ISR Number: 3395348-6Report Type:Periodic Company Report #A0073968  
 Age:39 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia Pruritus Rash Papular	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /							
THREE TIMES							
PER DAY ORAL							

Thyroxine Sodium	C
Ibuprofen	C
Lortab	C

Date:02/01/99ISR Number: 3395349-8Report Type:Periodic Company Report #A0073977  
 Age:33 YR Gender:Female I/FU:I

Outcome	PT
	Anxiety Depression

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Flushing Nausea Vomiting	Report Source	Product	Role	Manufacturer	Route
150 MG / IN	THE MORNING /		Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL				Ranitidine Hydrochloride	C		
				Piroxicam	C		
				Dicyclomine	C		
				Nicotine	C		

Date:02/01/99ISR Number: 3395350-4Report Type:Periodic Company Report #A0074032  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER		Dermatitis Erythema Pruritus	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
DAY / ORAL		Urticaria		Zyban Tablet - Zyban	SS		ORAL
UNK / UNK /							
ORAL							

Date:02/01/99ISR Number: 3395351-6Report Type:Periodic Company Report #A0073513  
 Age:45 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Disturbance In Attention Drug Ineffective Irritability	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

Mood Swings

PER DAY /

ORAL 3 MON

Wellbutrin Tablet SS

ORAL

UNK / UNK /

ORAL

Conjugated Estrogens C  
Darvocet-N C

Date:02/01/99ISR Number: 3395352-8Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #A0073518

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Vision Blurred Visual Disturbance	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

150 MG / PER

DAY / ORAL

No Concurrent  
Medication C

Date:02/01/99ISR Number: 3395353-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0073519

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Increased Appetite	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

150 MG / PER

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

DAY / ORAL

Fluoxetine  
Hydrochloride C

Date:02/01/99ISR Number: 3395354-1Report Type:Periodic  
Age:24 YR Gender:Male I/FU:I

Company Report #A0073520

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness Nervousness Tremor	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

150 MG /

TWICE PER DAY

/ ORAL

Date:02/01/99ISR Number: 3395356-5Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #A0073538

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pain In Extremity	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL

150 MG /

TWICE PER DAY

/ ORAL

Phentermine C

Date:02/01/99ISR Number: 3395357-7Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0073575

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Gingivitis	Consumer	Wellbutrin Tablet-Controlled			

150 MG /  
TWICE PER DAY  
/ ORAL  
Release PS ORAL  
Paroxetine  
Hydrochloride C

Date:02/01/99ISR Number: 3395375-9Report Type:Periodic Company Report #A0072208  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

150 MG /  
TWICE PER DAY  
/ ORAL

Date:02/01/99ISR Number: 3395376-0Report Type:Periodic Company Report #A0072223  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia Mania Tremor	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

TWICE PER DAY  
/ ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3395378-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0072304

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY / ORAL	1 YR	Agitation Drug Effect Decreased	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:02/01/99ISR Number: 3395379-6Report Type:Periodic  
 Age:41 YR Gender:Male I/FU:I

Company Report #A0072326

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL		Agitation Dizziness	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
				Diltiazem Hydrochloride Clonidine	C C		

Date:02/01/99ISR Number: 3395381-4Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #A0072327

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
TWICE PER DAY/ ORAL		Cold Sweat Irritability Tremor	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
				Paracetamol	C		

Date:02/01/99ISR Number: 3395383-8Report Type:Periodic Company Report #A0072328  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Gait Disturbance	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL				Thyroxine Sodium Lo/Ovral	C C		

Date:02/01/99ISR Number: 3395384-XReport Type:Periodic Company Report #A0072354  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthritis	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3395385-1Report Type:Periodic Company Report #A0072355  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL

150 MG/ ORAL 3 DAY

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3395387-5Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0072477

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3395388-7Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0072497

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL	10 DAY						

Date:02/01/99ISR Number: 3395389-9Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #A0072543

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence Headache Nervousness	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ PER DAY/ ORAL				Carisoprodol Tramadol Hydrochloride Brimonidine Tartrate Conjugated Estrogens	C C C C C		

Date:02/01/99ISR Number: 3395390-5Report Type:Periodic  
Age:70 YR Gender:Female I/FU:I

Company Report #A0072544

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Constipation Depression Dry Mouth	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
TWICE PER							
DAY/ ORAL							
				Simvastatin Alprazolam	C C		

Date:02/01/99ISR Number: 3395391-7Report Type:Periodic Company Report #A0072545  
Age:26 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Amnesia Memory Impairment	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
TWICE PER							
DAY/ ORAL							

Date:02/01/99ISR Number: 3395393-0Report Type:Periodic Company Report #A0072603  
Age:60 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety Tinnitus	Health Professional	Wellbutrin Tablet-Controlled			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

150MG/TWICE				Release	PS		ORAL
PER DAY/ ORAL				Fluoxetine Hydrochloride Mesalazine	C C		

Date:02/01/99ISR Number: 3395394-2Report Type:Periodic Company Report #A0072604  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nightmare	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ ORAL			Representative				

Date:02/01/99ISR Number: 3395396-6Report Type:Periodic Company Report #A0072605  
 Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Confusional State Orthostatic Hypotension	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ ORAL			Representative	Blood Pressure Medication	C		

Date:02/01/99ISR Number: 3395397-8Report Type:Periodic Company Report #A0072609  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3395399-1Report Type:Periodic Company Report #A0072610  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health Professional Other	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3395400-5Report Type:Periodic Company Report #A0072612  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
TWICE PER DAY							
/ ORAL				Estrogen Progesterone	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3395402-9Report Type:Periodic  
Age:18 YR Gender:Male I/FU:I

Company Report #A0072619

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Pain Rash Pruritic	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
TWICE PER DAY							
/ ORAL							

Date:02/01/99ISR Number: 3395404-2Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #A0073587

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chills Influenza Like Illness	Consumer	Wellbutrin Tablet- Controlled Release	PS		ORAL
150 MG/ ORAL							
		Pyrexia		Hormones	C		

Date:02/01/99ISR Number: 3395406-6Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0073593

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Loracarbef	C
Potassium Salt	C
Bumetanide	C
Carvedilol	C
Digoxin	C
Losartan Potassium	C
Lorazepam	C
Hydralazine	C
Prempro	C
Mirtazapine	C

Date:02/01/99ISR Number: 3395408-XReport Type:Periodic Company Report #A0073394  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Health	Wellbutrin Tablet -			
		Apathy	Professional	Controlled Release	PS		ORAL
300 MG/ PER		Attention					
DAY/ ORAL		Deficit/Hyperactivity		Semisodium Valproate			
		Disorder		(Formulation			
750 MG/ ORAL		Diarrhoea		Unknown)	SS		ORAL
		Hypersensitivity					
		Hypomania					

Date:02/01/99ISR Number: 3395410-8Report Type:Periodic Company Report #A0073109  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Wellbutrin Tablet -			
		Nausea		Controlled Release	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3395411-XReport Type:Periodic  
Age:19 YR Gender:Male I/FU:I

Company Report #A0073110

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Health Professional	Wellbutrin Tablet - Controlled Release	PS		ORAL
TWICE PER DAY/ ORAL							

Date:02/01/99ISR Number: 3395413-3Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0073111

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Malaise Mental Impairment	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
150 MG/ PER DAY / ORAL							

Date:02/01/99ISR Number: 3395415-7Report Type:Periodic  
Age:58 YR Gender:Male I/FU:I

Company Report #A0073112

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Increased Appetite	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
TWICE PER DAY/ ORAL							
				Diazepam	C		
				Salbutamol Sulphate	C		
				Nicotine	C		

Date:02/01/99ISR Number: 3395416-9Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #A0073113

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

	Alopecia	Consumer	Wellbutrin Tablet -		
	Glossodynia		Controlled Release	PS	ORAL
150 MG/ TWICE					
PER DAY /	Halitosis				
ORAL			Wellbutrin Tablet -		
			Controlled Release	SS	ORAL
150 MG/ TWICE					
PER DAY/ ORAL			Conjugated Estrogens	C	
			Multivitamin	C	

Date:02/01/99ISR Number: 3395418-2Report Type:Periodic Company Report #A0073115  
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Wellbutrin Tablet -			
				Controlled Release	PS		ORAL
TWICE PER							
DAY/ ORAL							

Date:02/01/99ISR Number: 3395419-4Report Type:Periodic Company Report #A0073288  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Balance Disorder	Health Professional	Wellbutrin Tablet -			
			Company Representative	Controlled Release	PS		ORAL
ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3395421-2Report Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #A0072767

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Wellbutrin Tablet -Controlled Release	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL							

Date:02/01/99ISR Number: 3395422-4Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0072768

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression Drug Ineffective	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
THREE TIMES PER DAY/ ORAL							

Estrogen	C
Thyroxine Sodium	C

Date:02/01/99ISR Number: 3395424-8Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0072770

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Health Professional	Wellbutrin Tablet- Controlled Release	PS		ORAL
TWICE PER DAY/ ORAL							

Date:02/01/99ISR Number: 3395426-1Report Type:Periodic  
Age:33 YR Gender:Male I/FU:I

Company Report #A0072984

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation	Consumer	Wellbutrin Tablet -			

150 MG /  
 TWICE PER DAY  
 / ORAL

Dizziness  
 Tremor

Controlled Release PS ORAL

No Concurrent Medication C

Date:02/01/99ISR Number: 3395427-3Report Type:Periodic Company Report #A0071750  
 Age: Gender:Unknown I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tremor	Health Professional	Wellbutrin Tablet- Controlled Release	PS		ORAL
			Company Representative	Nefazodone Hydrochloride (Formulation Unknown)	SS		

ORAL

Date:02/01/99ISR Number: 3395429-7Report Type:Periodic Company Report #A0071816  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abnormal Behaviour Insomnia	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
		Tremor		Fluoxetine Hydrochloride	C		

150 MG/ PER  
 DAY/ ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3395430-3Report Type:Periodic  
 Age:49 YR Gender:Female I/FU:I

Company Report #A0071857

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
150 MG /							
TWICE PER							
DAY/ ORAL							
				Carisoprodol	C		
				Estrogen	C		
				Omeprazole	C		
				Ibuprofen	C		

Date:02/01/99ISR Number: 3395432-7Report Type:Periodic  
 Age:53 YR Gender:Male I/FU:I

Company Report #A0071905

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Dry Mouth	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
UNK / TWICE							
PER DAY/ ORAL		Headache					
		Joint Stiffness Muscle Rigidity Tremor		Doxycycline	C		

Date:02/01/99ISR Number: 3395433-9Report Type:Periodic  
 Age:60 YR Gender:Female I/FU:I

Company Report #A0071906

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Smoker	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
TWICE PER DAY							
/ ORAL							

Date:02/01/99ISR Number: 3395434-0Report Type:Periodic Company Report #A0072002  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation Hyperhidrosis	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
150 MG/ PER							
DAY/ ORAL				Oestradiol	C		

Date:02/01/99ISR Number: 3395548-5Report Type:Periodic Company Report #M082966  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation	Consumer	Trazodone Tabs	PS		ORAL
ORAL				Wellbutrin	SS		
1 YR							

Date:02/01/99ISR Number: 3403909-0Report Type:Periodic Company Report #A0076439  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depressed Mood Irritability	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / UNK							
/ ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3403910-7Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #A0076536

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
450 MG / PER DAY/ ORAL		Insomnia Tic Weight Increased	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
60 MG/ PER DAY/ UNKNOWN				Paroxetine Hydrochloride (Formulation Unknown)	SS		
				Buspirone Hydrochloride	C		

Date:02/01/99ISR Number: 3403911-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0076540

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Headache Urticaria	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:02/01/99ISR Number: 3403912-0Report Type:Periodic  
 Age:28 YR Gender:Female I/FU:I

Company Report #A0076581

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Balance Disorder Feeling Abnormal Headache	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

Hypoaesthesia

TWICE PER DAY

/ ORAL

Paroxetine  
Hydrochloride C  
Multivitamin C

Date:02/01/99ISR Number: 3403913-2Report Type:Periodic

Company Report #A0076759

Age:11 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Flushing	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
100 MG / PER							
DAY / ORAL							

Date:02/01/99ISR Number: 3403914-4Report Type:Periodic

Company Report #A0076787

Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3403915-6Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #A0076788

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Agitation Disorientation Insomnia Memory Impairment	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:02/01/99ISR Number: 3403916-8Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #A0076789

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG / TWICE PER DAY / ORAL		Amnesia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
	YR			Nefazodone Hydrochloride	C		
				Buspirone Hydrochloride	C		

Date:02/01/99ISR Number: 3403917-XReport Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0076790

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Insomnia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

Alprazolam C  
 Omeprazole C  
 Conjugated Estrogens C  
 Thyroxine Sodium C  
 Vitamin E C  
 Paracetamol C  
 Dicyclomine C

Date:02/01/99ISR Number: 3403918-1Report Type:Periodic Company Report #A0076852  
 Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Fatigue Nausea Nervousness Tremor	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:02/01/99ISR Number: 3404005-9Report Type:Periodic Company Report #A0061813  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK /UNK / ORAL		Nausea	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
				Cancer Chemotherapy	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404006-0Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0061916

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK / UNK /		Dermatitis	Health	Wellbutrin Tablet	PS		ORAL
ORAL		Dyspnoea	Professional				
			Company Representative				

Date:02/01/99ISR Number: 3404007-2Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0062035

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK / UNK /		Medication Residue	Health	Wellbutrin Tablet	PS		ORAL
ORAL			Professional				

Date:02/01/99ISR Number: 3404008-4Report Type:Periodic  
 Age:63 YR Gender:Male I/FU:I

Company Report #A0062041

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG / TWICE		Photosensitivity Reaction	Health	Wellbutrin Tablet	PS		ORAL
PER DAY /			Professional				
ORAL				DiFlunisal	C		

Date:02/01/99ISR Number: 3404009-6Report Type:Periodic  
 Age:71 YR Gender:Female I/FU:I

Company Report #A0062233

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Vomiting	Health	Wellbutrin			

			Professional	(Formulation			
				Unknown)		PS	ORAL
UNK / UNK /							
ORAL	2	WK					
Date:02/01/99ISR Number: 3404010-2Report Type:Periodic Company Report #A0062240							
Age:54 YR Gender:Female I/FU:I							
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Arthralgia	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG / PER		Eye Disorder					
DAY / ORAL	1	YR					
		Muscle Spasms		Conjugated Estrogens	C		
		Pollakiuria		Loratadine	C		
		Sensation Of Pressure		Vitamin E	C		
				Multivitamin	C		
				Calcium Salt	C		

Date:02/01/99ISR Number: 3404011-4Report Type:Periodic Company Report #A0062261							
Age:64 YR Gender:Male I/FU:I							
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Condition Aggravated	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG /		Drug Interaction					
TWICE PER DAY		Herpes Simplex					
/ ORAL				Famciclovir Tablet	SS		ORAL
500 MG /							
TWICE PER DAY							
/ ORAL				Multivitamin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404012-6Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0062299

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK / UNK /		Hypoglycaemia	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL	3 WK						

Date:02/01/99ISR Number: 3404013-8Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0062334

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG /		Convulsion	Health Professional	Wellbutrin Tablet	PS		ORAL
TWICE PER DAY				Nortriptyline	C		
/ ORAL							

Date:02/01/99ISR Number: 3404014-XReport Type:Periodic  
Age:17 YR Gender:Unknown I/FU:I

Company Report #A0062375

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG / TWICE		Dermatitis	Health Professional	Wellbutrin Tablet	PS		ORAL
PER DAY /							
ORAL							

Date:02/01/99ISR Number: 3404015-1Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0062377

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK / UNK /		Hypoglycaemia	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL	3 WK						

Date:02/01/99ISR Number: 3404016-3Report Type:Periodic Company Report #A0062485  
 Age:64 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG / TWICE PER DAY /		Alopecia	Consumer	Wellbutrin Tablet	PS		ORAL
ORAL		Constipation					
UNK / UNK /		Dizziness					
ORAL		Dysphagia		Paroxetine Hydrochloride Tablet	SS		ORAL
				Fluvoxamine Maleate	C		
				Sertraline Hydrochloride	C		

Date:02/01/99ISR Number: 3404017-5Report Type:Periodic Company Report #A0062635  
 Age: Gender:Unknown I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK / UNK /		Arthralgia	Health Professional	Wellbutrin Tablet	PS		ORAL
ORAL		Dermatitis					
		Pyrexia					
		Vasculitis					



150 MG/ TWICE  
 PER DAY /  
 ORAL

Frequent Bowel Movements  
 Proctalgia

Health  
 Professional  
 Company  
 Representative

Wellbutrin Tablet  
 PS

ORAL

Date:02/01/99ISR Number: 3404022-9Report Type:Periodic Company Report #A0062800  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG / PER DAY / ORAL	2 WK	Pain In Jaw Pharyngolaryngeal Pain Urticaria	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3404023-0Report Type:Periodic Company Report #A0063752  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK/SEE TEXT/ ORAL		Hypoaesthesia	Health Professional	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3404024-2Report Type:Periodic Company Report #A0062861  
 Age:42 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG / FOUR		Drug Ineffective	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

Freedom Of Information (FOI) Report

TIMES PER DAY

ORAL

Gabapentin

C

Date:02/01/99ISR Number: 3404025-4Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0075563

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus Rash Erythematous	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ TWICE							

PER DAY /

ORAL

Levora

C

Date:02/01/99ISR Number: 3404026-6Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #A0075564

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Anxiety Decreased Appetite	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
UNK / UNK /							

UNK / UNK /

Paraesthesia

ORAL

Date:02/01/99ISR Number: 3404027-8Report Type:Periodic  
Age:60 YR Gender:Male I/FU:I

Company Report #A0075565

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Fatigue Sedation	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /							

150 MG /

TWICE PER DAY

/ ORAL

Blood Pressure Medication C

Date:02/01/99ISR Number: 3404028-XReport Type:Periodic Company Report #A0075592  
Age:50 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / IN		Crying Headache Heart Rate Increased	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
THE MORNING /		Irritability					
ORAL		Mood Altered					

Hydrochlorothiazide C  
Medroxyprogesterone C  
Conjugated Estrogens C

Date:02/01/99ISR Number: 3404029-1Report Type:Periodic Company Report #A0075593  
Age:46 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / IN		Crying Depression Insomnia	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
THE MORNING /							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL

Date:02/01/99ISR Number: 3404030-8Report Type:Periodic Company Report #A0075604  
 Age:13 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG / TWICE PER DAY / ORAL		Halitosis	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:02/01/99ISR Number: 3404031-XReport Type:Periodic Company Report #A0075614  
 Age: Gender:Unknown I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK / UNK / ORAL		Dermatitis Urticaria	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:02/01/99ISR Number: 3404032-1Report Type:Periodic Company Report #A0075615  
 Age: Gender:Unknown I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK / UNK / ORAL		Dermatitis Urticaria	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Face Oedema Pain Rash Pruritic	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /		Urticaria	Representative				
TWICE PER DAY							
/ ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
100 MG /							
TWICE PER DAY							
/ ORAL				Hydroxyzine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404035-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0076018

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /							
THREE TIMES							
PER DAY /							
ORAL							

Date:02/01/99ISR Number: 3404036-9Report Type:Periodic  
 Age:58 YR Gender:Male I/FU:I

Company Report #A0076044

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tic	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / FOUR							
TIMES PER DAY							
/ ORAL	1	YR					

Date:02/01/99ISR Number: 3404037-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0076087

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amenorrhoea Dry Mouth Headache	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
UNK / UNK /							
ORAL	3	MON					

Date:02/01/99ISR Number: 3404038-2Report Type:Periodic Company Report #A0076115  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression Suicidal	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / UNK							
/ ORAL							

Date:02/01/99ISR Number: 3404039-4Report Type:Periodic Company Report #A0076196  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation Diarrhoea Dry Mouth	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /							
THREE TIMES							
PER DAY /							
ORAL							
				Hypericum	C		

Date:02/01/99ISR Number: 3404040-0Report Type:Periodic Company Report #A0076360  
Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source
Urticaria		Health Professional

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Company Representative	Product	Role	Manufacturer	Route
UNK / UNK /			Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL						

Date:02/01/99ISR Number: 3404041-2Report Type:Periodic Company Report #A0076361  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ecchymosis	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
UNK / UNK /							
ORAL							

Date:02/01/99ISR Number: 3404042-4Report Type:Periodic Company Report #A0076362  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ecchymosis	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
UNK / UNK /							
ORAL							

Date:02/01/99ISR Number: 3404043-6Report Type:Periodic Company Report #A0076363  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Health	Wellbutrin			

UNK/ UNK/ Professional Company Release PS ORAL  
ORAL Representative

Date:02/01/99ISR Number: 3404044-8Report Type:Periodic Company Report #A0076390  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Tic	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / PER							
DAY / ORAL	7	DAY					

Date:02/01/99ISR Number: 3404046-1Report Type:Periodic Company Report #A0075058  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flushing Menopause	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL

UNK/ UNK/  
ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404047-3Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0075078

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Renal Colic Visual Disturbance	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
				Lithium Salt	C		

Date:02/01/99ISR Number: 3404049-7Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #A0075079

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL		Dermatitis Urticaria	Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:02/01/99ISR Number: 3404051-5Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #A0075197

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Rash Generalised	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
				Conjugated Estrogens Quinapril Hydrochloride	C C		

Digoxin C  
Hydrochlorothiazide C

Date:02/01/99ISR Number: 3404053-9Report Type:Periodic Company Report #A0075265  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Disturbance In Attention Feeling Abnormal	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
300 MG / PER DAY / ORAL							

Date:02/01/99ISR Number: 3404058-8Report Type:Periodic Company Report #A0075266  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Dizziness	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / TWICE PER DAY/ ORAL							
10 MG / UNK / UNKNOWN				Buspirone Hydrochloride (Formulation Unknown)	SS		

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Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404059-XReport Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0075271

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Urticaria	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
UNK/ UNK/			Representative				
ORAL							

Date:02/01/99ISR Number: 3404061-8Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #A0075299

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Overdose Pruritus	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
600 MG/ PER				Venlafaxine Hydrochloride	C		
DAY/ ORAL				Nefazodone Hydrochloride	C		

Date:02/01/99ISR Number: 3404063-1Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0075302

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth Insomnia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ IN				Nefazodone Hydrochloride	C		
THE MORNING /							
ORAL							

Date:02/01/99ISR Number: 3404070-9Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #A0075309

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia Dizziness Flushing	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY		Nausea					
/ ORAL		Tremor		Lisinopril Pravastatin Sodium Thyroxine Sodium Sulindac	C C C C		

Date:02/01/99ISR Number: 3404080-1Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0075311

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Hair Disorder Insomnia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /		Tremor					
TWICE PER							
DAY/ ORAL				Lithium Salt	C		

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Freedom Of Information (FOI) Report

Alprazolam C  
 Thyroxine Sodium C

Date:02/01/99ISR Number: 3404082-5Report Type:Periodic Company Report #A0075340  
 Age:45 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Anorexia Nasopharyngeal Disorder Nausea Panic Attack Weight Decreased	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

Ascriptin C  
 Ranitidine  
 Hydrochloride C  
 Multivitamin C

Date:02/01/99ISR Number: 3404084-9Report Type:Periodic Company Report #A0075342  
 Age:55 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK/ UNK/ ORAL	1 WK	Dermatitis	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:02/01/99ISR Number: 3404085-0Report Type:Periodic Company Report #A0075391  
 Age:65 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache Tinnitus	Health Professional	Wellbutrin Tablet-Controlled			

UNKN/ UNKN/ Release PS ORAL

ORAL

Date:02/01/99ISR Number: 3404087-4Report Type:Periodic Company Report #A0075393  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / PER							
DAY / ORAL				Sumatriptan Succinate	C		
				Midrin	C		
				Beclomethasone Dipropion	C		

Date:02/01/99ISR Number: 3404092-8Report Type:Periodic Company Report #A0075394  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ TWICE							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

PER DAY /

ORAL

Date:02/01/99ISR Number: 3404094-1Report Type:Periodic Company Report #A0075395  
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Headache Pollakiuria	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL	4	WK		Librium Venlafaxine Hydrochloride	C		

Date:02/01/99ISR Number: 3404096-5Report Type:Periodic Company Report #A0075412  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia Oedema Peripheral Rash Pruritic	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /							
TWICE PER DAY		Urticaria	Representative				
/ ORAL							

Date:02/01/99ISR Number: 3404098-9Report Type:Periodic Company Report #A0075523  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis	Health Professional	Wellbutrin Tablet-Controlled			

150 MG /  
TWICE PER  
DAY/ ORAL  
Company Representative  
Release  
PS  
ORAL

Date:02/01/99ISR Number: 3404099-0Report Type:Periodic Company Report #A0075562  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying Depression Emotional Disorder	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ PER				Vicodin Lortab	C C		

Date:02/01/99ISR Number: 3404101-6Report Type:Periodic Company Report #A0060122  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphagia Insomnia	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG / TWICE							
PER DAY /							
ORAL							



100 MG /	Abdominal Distension	Consumer	Wellbutrin Tablet	PS	ORAL
TWICE PER DAY	Constipation				
/ ORAL	Insomnia				
6 WK			Nicotine Patch	SS	
TRANSDERMAL	CONTINUOUS /				
TRANSDERMAL			Antibiotics	C	

Date:02/01/99ISR Number: 3404106-5Report Type:Periodic Company Report #A0060321  
 Age:17 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion	Health Professional Company	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL			Representative				

Date:02/01/99ISR Number: 3404107-7Report Type:Periodic Company Report #A0060323  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Health Professional Other	Wellbutrin (Formulation Unknown)	PS		ORAL
VARIABLE DOSE				Loestrin Tablet	SS		ORAL
/ ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404109-0Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #A0060364

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG / PER DAY / ORAL	2 MON	Loss Of Consciousness	Consumer	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3404110-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0060379

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Pruritus	Consumer Company Representative	Wellbutrin (Formulation Unknown)	PS		ORAL
				Quetiapine Fumarate	C		

Date:02/01/99ISR Number: 3404111-9Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0060393

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Blindness Convulsion Corneal Oedema	Health Professional Other	Wellbutrin (Formulation Unknown)	PS		ORAL
				Amantadine Hydrochloride (Formulation Unknown)	SS		

Date:02/01/99ISR Number: 3404113-2Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0060399

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Arrhythmia	Health	Wellbutrin			

	Headache	Professional	(Formulation		
	Hypoaesthesia	Other	Unknown)	PS	ORAL
THREE TIMES					
PER DAY /					
ORAL	1	YR			
			Dexfenfluramine	C	
			Buspirone		
			Hydrochloride	C	
			Phentermine	C	
			Fenfluramine Hcl	C	
			Conjugated Estrogens	C	

Date:02/01/99ISR Number: 3404114-4Report Type:Periodic Company Report #A0060406  
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Drug Ineffective Tinnitus	Consumer Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
150 MG /			Other				
TWICE PER DAY							
/ ORAL							
				Venlafaxine Hydrochloride (Formulation Unknown)	SS		ORAL
150 MG /							
TWICE PER DAY							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

/ ORAL

Date:02/01/99ISR Number: 3404115-6Report Type:Periodic Company Report #A0060420  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Wellbutrin Tablet	PS		ORAL
ORAL		Drug Ineffective					
		Insomnia					

Date:02/01/99ISR Number: 3404117-XReport Type:Periodic Company Report #A0060458  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Wellbutrin Tablet	PS		ORAL
1 TABLET/ TWICE PER DAY		Insomnia					

/ ORAL

Date:02/01/99ISR Number: 3404118-1Report Type:Periodic Company Report #A0060620  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health Professional Company Representative	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL		Convulsion					

Date:02/01/99ISR Number: 3404119-3Report Type:Periodic Company Report #A0060666  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

	Dry Mouth	Health Professional Other	Wellbutrin (Formulation Unknown)	PS	ORAL
ORAL			Nicotine (Formulation Unknown)	SS	
2 MG					

Date:02/01/99ISR Number: 3404120-XReport Type:Periodic Company Report #A0060684  
 Age:29 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG / PER DAY / ORAL		Dermatitis	Consumer	Wellbutrin Tablet	PS		ORAL
		Hypersensitivity					
		Pruritus Urticaria		Ibuprofen	C		

Date:02/01/99ISR Number: 3404121-1Report Type:Periodic Company Report #A0060765  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Pollakiuria	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404123-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0060783

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3404124-7Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0060789

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Haemolytic Anaemia Thrombocytopenia	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3404126-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0060791

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Wellbutrin Tablet	PS		ORAL
100MG ORAL		Hypersensitivity Urticaria					

Date:02/01/99ISR Number: 3404128-4Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0060836

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Health	Wellbutrin Tablet	PS		ORAL
ORAL	3 WK		Professional Company Representative				

Date:02/01/99ISR Number: 3404129-6Report Type:Periodic Company Report #A0060856  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Consumer	Wellbutrin Tablet	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3404130-2Report Type:Periodic Company Report #A0060910  
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Consumer	Wellbutrin Tablet	PS		ORAL
75MG TWICE		Headache					
PER DAY ORAL		Insomnia					
		Lethargy					
		Weight Increased					

Date:02/01/99ISR Number: 3404131-4Report Type:Periodic Company Report #A0060916  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Wellbutrin			
		Hypersensitivity		(Formulation			
		Urticaria		Unknown)	PS		ORAL
100MG ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404133-8Report Type:Periodic  
Age: Gender:Not SpecifiedFU:I

Company Report #A0060979

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health Professional Company Representative	Wellbutrin (Formulation Unknown)	PS		

Date:02/01/99ISR Number: 3404135-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0060980

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Wellbutrin Tablet	PS		ORAL
ORAL		Insomnia					

Date:02/01/99ISR Number: 3404136-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0061002

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Wellbutrin Tablet	PS		ORAL
100MG TWICE		Hypersensitivity					
PER DAY ORAL		Irritability		Wellbutrin Tablet- Controlled Release	SS		ORAL
100MG TWICE		Urticaria					
PER DAY ORAL							

Date:02/01/99ISR Number: 3404138-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0061020

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Wellbutrin Tablet	PS		ORAL
75MG TWICE							

PRT DAY ORAL

Potassium Salt C  
Propranolol C  
Hydrochlorothiazide C  
Clonazepam C

Date:02/01/99ISR Number: 3404139-9Report Type:Periodic Company Report #A0061055  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dissociation	Health	Wellbutrin Tablet	PS		ORAL
75MG ORAL			Professional	Zyban Tablet - Zyban	SS		ORAL
150MG ORAL				Wellbutrin Tablet- Controlled Release	SS		ORAL
100MG ORAL							

Date:02/01/99ISR Number: 3404141-7Report Type:Periodic Company Report #A0061097  
Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gastrointestinal Disorder	Consumer	Wellbutrin Tablet	PS		ORAL
150MG IN THE		Insomnia					
MORNING ORAL	6 WK						



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Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404142-9Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #A0061116

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG TWICE PER DAY ORAL		Cardiac Enzymes Increased Chest Pain	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3404144-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0061208

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75MG PER DAY ORAL	3 DAY	Muscle Spasms Vomiting	Health Professional	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3404145-4Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0061218

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG PER DAY ORAL		Deafness Tinnitus	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3404147-8Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #A0061276

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG TWICE		Carpal Tunnel Syndrome	Health	Wellbutrin Tablets	PS		ORAL

PER DAY ORAL  
Middle Insomnia Professional  
Pain In Extremity Cefaclor C  
Paraesthesia

Date:02/01/99ISR Number: 3404148-XReport Type:Periodic Company Report #A0061333  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity	Health	Wellbutrin Tablet	PS		ORAL
ORAL			Professional				

Date:02/01/99ISR Number: 3404149-1Report Type:Periodic Company Report #A0061385  
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Wellbutrin Tablet	PS		ORAL
100MG TWICE							
PER DAY ORAL							

Date:02/01/99ISR Number: 3404151-XReport Type:Periodic Company Report #A0061393  
Age:55 YR Gender:Male I/FU:I

Outcome  
PT  
Abnormal Dreams  
Burning Sensation  
Flushing  
Pruritus

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG THREE TIMES PER DAY ORAL		Skin Irritation Sunburn Tenderness	Consumer	Wellbutrin Tablet	PS		ORAL
				Clonidine Hydrochloride	C		
				Triamcinolone Acetonide	C		
				Diazepam	C		
				Excedrin	C		

Date:02/01/99ISR Number: 3404152-1Report Type:Periodic Company Report #A0061422  
Age:6 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75MG PER DAY ORAL		Abnormal Behaviour Aggression Dissociation Mania	Consumer	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3404154-5Report Type:Periodic Company Report #A0061437  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75MG THREE TIMES PER DAY ORAL		Agitation Condition Aggravated Dizziness Headache Insomnia	Consumer	Wellbutrin Tablet	PS		ORAL
300MG PER DAY				Wellbutrin Tablet - Controlled Release	SS		ORAL

ORAL

Date:02/01/99ISR Number: 3404187-9Report Type:Periodic Company Report #A0063431  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Spasms	Health	Wellbutrin Tablet	PS		ORAL
100 MG/PER		Speech Disorder	Professional				
DAY/ORAL				Wellbutrin Tablet-Controlled Release	SS		ORAL
150 MG/TWICE							
PER DAY/ORAL				Thyroxine Sodium	C		

Date:02/01/99ISR Number: 3404188-0Report Type:Periodic Company Report #A0063438  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG/ORAL		Insomnia		Semisodium Valproate	C		
		Nervousness		Clonazepam	C		
		Tremor					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404189-2Report Type:Periodic  
 Age:53 YR Gender:Female I/FU:I

Company Report #A0063459

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Abdominal Pain Amnesia Arrhythmia	Consumer Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
		Chest Pain	Other				
15 MG/TWICE PER DAY/ORAL		Condition Aggravated Crying Depression		Dexfenfluramine (Formulation Unknown)	SS		ORAL
		Diarrhoea					
		Dyspnoea Emotional Disorder Headache Nervousness Neurosis Oedema Palpitations Syncope Tinnitus		Diltiazem Hydrochloride Metoprolol Succinate	C C		

Date:02/01/99ISR Number: 3404190-9Report Type:Periodic  
 Age:60 YR Gender:Female I/FU:I

Company Report #A0063512

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL	18 MON	Antinuclear Antibody Fibromyalgia	Health Professional	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3404192-2Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0063851

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Nausea	Consumer	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3404193-4Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #A0063558

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
75 MG/TWICE			Consumer	Wellbutrin Tablet	PS		ORAL
PER DAY/ORAL	2	MON					
		Insomnia					
		Weight Increased		Estratest	C		
				Medroxyprogesterone			
				Ace.	C		
				Thyroxine Sodium	C		
				Simvastatin	C		

Date:02/01/99ISR Number: 3404195-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0063606

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
ORAL	2	WK	Health	Wellbutrin Tablet	PS		ORAL
			Professional	Fluoxetine			
				Hydrochloride	C		

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Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404197-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0063658

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Accommodation Disorder Amnesia Dysphagia	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
		Dyspraxia		Semisodium Valproate	C		

Date:02/01/99ISR Number: 3404198-3Report Type:Periodic  
Age:71 YR Gender:Female I/FU:I

Company Report #A0063689

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Asthenia Dyspnoea	Health Professional	Wellbutrin Tablet	PS		ORAL
		Insomnia Malaise Muscle Spasms Nightmare Tremor Vomiting		Nefazodone Hydrochloride Multivitamin	C C		

Date:02/01/99ISR Number: 3404201-0Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0064071

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG/THREE TIMES PER DAY/ORAL		Antibody Test Abnormal Eosinophilia Leukopenia	Health Professional	Wellbutrin Tablet	PS		ORAL
100 MG/THREE		Neutropenia		Wellbutrin Tablet-Controlled Release	SS		ORAL

TIMES PER

DAY/ORAL

Date:02/01/99ISR Number: 3404202-2Report Type:Periodic Company Report #A0069599  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Libido Increased	Consumer	Wellbutrin Tablet	PS		ORAL
				Sertraline Hydrochloride	C		
				Alprazolam	C		

Date:02/01/99ISR Number: 3404203-4Report Type:Periodic Company Report #A0069621  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Epistaxis	Health	Wellbutrin Tablet	PS		ORAL
ORAL			Professional	Bupropion Hydrochloride	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404205-8Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0069626

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amenorrhoea	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3404206-XReport Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #A0069628

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Health	Wellbutrin Tablet	PS		ORAL
ORAL							
			Professional Other	Hydroxyzine Hydrochloride Thyroxine Sodium	C C		

Date:02/01/99ISR Number: 3404208-3Report Type:Periodic  
Age:66 YR Gender:Female I/FU:I

Company Report #A0069666

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG/TWICE PER DAY/ORAL							

Date:02/01/99ISR Number: 3404209-5Report Type:Periodic  
Age:9 YR Gender:Male I/FU:I

Company Report #A0069703

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gynaecomastia	Health Professional Company Representative	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3404211-3Report Type:Periodic Company Report #A0069978  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperglycaemia	Consumer	Wellbutrin Tablet	PS		ORAL
150 MG/TWICE			Health				
PER DAY/ORAL			Professional	Humalog	C		
			Other	Int./Long-Acting Insulin	C		

Date:02/01/99ISR Number: 3404212-5Report Type:Periodic Company Report #A0070023  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Excitability	Health	Wellbutrin			
			Professional	(Formulation			
ORAL			Other	Unknown)	PS		ORAL
				Fluoxetine			
				Hydrochloride	C		
				Nortriptyline Hcl	C		
				Paroxetine			
				Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404213-7Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0070124

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Pain	Health	Wellbutrin Tablet	PS		ORAL
75		Menorrhagia	Professional				
MG/VARIABLE		Menstruation Irregular					
DOSE/ORAL		Pelvic Pain		Trazodone	C		
				Glucophage	C		
				Int./Long-Acting			
				Insulin	C		
				Enalapril Maleate	C		
				Hydrochlorothiazide	C		
				Fluvastatin Sodium	C		
				Loratadine	C		
				Ibuprofen	C		

Date:02/01/99ISR Number: 3404214-9Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #A0070125

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG/THREE		Insomnia					
TIMES PER		Tobacco Abuse					
DAY/ORAL				Wellbutrin			
				Tablet-Controlled			
				Release	SS		ORAL
150 MG/TWICE							
PER DAY/ORAL				Thyroxine Sodium	C		

Date:02/01/99ISR Number: 3404215-0Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0070170

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nail Discolouration	Health Professional Company	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL			Representative				

Date:02/01/99ISR Number: 3404216-2Report Type:Periodic Company Report #A0070258  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia Headache Sexual Dysfunction	Consumer	Wellbutrin Tablet Estrogen	PS C		ORAL

Date:02/01/99ISR Number: 3404217-4Report Type:Periodic Company Report #A0070325  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams Insomnia	Consumer Health Professional Other	Wellbutrin (Formulation Unknown) Nicotine (Formulation Unknown)	PS SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404218-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0070327

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache	Consumer Health Professional Other	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3404220-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0070428

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Diarrhoea	Consumer Health Professional Other	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3404221-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0070467

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG / TWICE PER DAY		Blood Pressure Increased	Health Professional	Wellbutrin Tablet	PS		ORAL
50 MG / TWICE PER DAY				Mirtazapine Tablet	SS		ORAL

Date:02/01/99ISR Number: 3404222-8Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0070468

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hyperbilirubinaemia	Health Professional	Wellbutrin (Formulation			

Unknown)

PS

ORAL

Date:02/01/99ISR Number: 3404223-XReport Type:Periodic Company Report #A0070658  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health Professional Company Representative	Wellbutrin (Formulation Unknown)	PS		ORAL
			Serum Sickness				

Date:02/01/99ISR Number: 3404224-1Report Type:Periodic Company Report #A0070703  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health Professional Company Representative	Wellbutrin Tablet	PS		ORAL
			Overdose				



Other	Amyotrophy	Health	Wellbutrin		
	Asthenia	Professional	Tablet-Controlled		
150 MG /	Conversion Disorder		Release	PS	ORAL
	Convulsion				
TWICE PER DAY	Hypoaesthesia				
/ ORAL	Livedo Reticularis		Zyban Tablet - Zyban	SS	ORAL
150 MG/ PER	Malaise				
DAY/ ORAL	Nausea		Paroxetine		
	Paralysis		Hydrochloride	C	
	Peripheral Coldness		Ipratropium Bromide	C	
	Peripheral Vascular		Budesonide	C	
	Disorder		Pain Medication	C	
	Pulse Absent		Beclomethasone		
	Speech Disorder		Dipropion	C	
	Transient Ischaemic				
	Attack				
	Tremor				

Date:02/01/99ISR Number: 3404231-9Report Type:Periodic Company Report #A0070973  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Health Professional	Wellbutrin (Formulation			

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Gait Disturbance

Health  
Professional

Wellbutrin  
(Formulation  
Unknown)

PS

Date:02/01/99ISR Number: 3404238-1Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0071158

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Coeliac Disease Constipation Frequent Bowel Movements	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3404240-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0071363

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia Eructation	Consumer	Wellbutrin Tablet Ibuprofen Fexofenadine Hydrochlorid Deconamine	PS C C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404242-3Report Type:Periodic  
 Age:74 YR Gender:Male I/FU:I

Company Report #A0065290

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG/PER DAY/ORAL		Constipation	Consumer	Wellbutrin Tablet	PS		ORAL
		Photosensitivity Reaction					
		Pruritus		Lithium Salt	C		
				Aloe	C		
				Warfarin Sodium	C		
				Thyroxine Sodium	C		
				Vitamin	C		

Date:02/01/99ISR Number: 3404244-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0065365

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Mastication Disorder Parosmia	Health Professional	Wellbutrin (Formulation Unknown)	PS		

Date:02/01/99ISR Number: 3404245-9Report Type:Periodic  
 Age:70 YR Gender:Female I/FU:I

Company Report #A0065413

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/PER DAY/ORAL		Libido Increased	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3404247-2Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0065484

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Muscle Rigidity  
Nervous System Disorder  
Tardive Dyskinesia

Health  
Professional

Wellbutrin  
(Formulation  
Unknown)

PS

Date:02/01/99ISR Number: 3404249-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0065549

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Wellbutrin (Formulation Unknown)	PS		

Date:02/01/99ISR Number: 3404250-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0066153

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Malaise	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL

ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404252-6Report Type:Periodic  
 Age:31 YR Gender:Female I/FU:I

Company Report #A0066252

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Diarrhoea Dizziness Dyspnoea	Consumer Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
			Other	Fenfluramine Hcl (Formulation Unknown)	SS		ORAL
	20 MG/THREE TIMES PER DAY/ORAL			Phentermine (Formulation Unknown)	SS		
				Nefazodone Hydrochloride (Formulation Unknown)	SS		
				Conjugated Estrogens	C		
				Conjugate Estrogens	C		

Date:02/01/99ISR Number: 3404253-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0066345

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Dermatitis Exfoliative Skin Disorder	Company Representative	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3404277-0Report Type:Periodic  
 Age:48 YR Gender:Male I/FU:I

Company Report #A0065450

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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150 MG/PER	Drug Ineffective	Consumer	Wellbutrin Tablet	PS	ORAL
DAY/ORAL	2 MON		Ritonavir	C	
			Lamivudine	C	
			Saquinavir	C	
			Co-Trimoxazole	C	

Date:02/01/99ISR Number: 3404278-2Report Type:Periodic Company Report #A0065469  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia	Consumer	Wellbutrin Tablet	PS		ORAL
225 MG/TWICE		Nausea					
PER DAY/ORAL		Tremor		Trazodone	C		
				Hydroxyzine	C		

Date:02/01/99ISR Number: 3404281-2Report Type:Periodic Company Report #A0065582  
 Age:50 YR Gender:Male I/FU:I

Outcome	PT
	Agitation
	Bruxism
	Depression
	Drug Interaction

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nervousness

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/PER DAY/ORAL		Consumer	Wellbutrin Tablet	PS		ORAL
20 MG/THREE TIMES PER DAY/ORAL	3 YR		Fluoxetine (Formulation Unknown) 10 Mg	SS		ORAL
			Anacin	C		

Date:02/01/99ISR Number: 3404283-6Report Type:Periodic Company Report #A0065745  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG/TWICE PER DAY/ORAL		Arthralgia	Health	Wellbutrin Tablet	PS		ORAL
		Back Pain	Professional				
.625 MG/TWICE PER DAY/ORAL		Neck Pain	Company Representative	Conjugated Estrogens Tablet	SS		ORAL

Date:02/01/99ISR Number: 3404285-XReport Type:Periodic Company Report #A0065866  
 Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG/SEE TEXT/ORAL		Alopecia	Consumer	Wellbutrin Tablet	PS		ORAL
				Hydroxyurea	C		

Aspirin C  
Lamotrigine C

Date:02/01/99ISR Number: 3404287-3Report Type:Periodic Company Report #A0066012  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Wellbutrin Tablet	PS		ORAL
TWICE PER							
DAY/ORAL							

Lorazepam C

Date:02/01/99ISR Number: 3404289-7Report Type:Periodic Company Report #A0066022  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG/PER							
DAY/ORAL	9 MON						

Date:02/01/99ISR Number: 3404290-3Report Type:Periodic Company Report #A0066032  
Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia	Health	Wellbutrin Tablet	PS		ORAL
150 MG/TWICE		Hyponatraemia	Professional				
PER DAY/ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404291-5Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #A0066039

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG/PER DAY/ORAL		Abnormal Dreams	Consumer	Wellbutrin Tablet	PS		ORAL
		Headache					
150 MG/TWICE PER DAY/ORAL		Mental Impairment		Zyban Tablet-Zyban	SS		ORAL
		Palpitations					
		Tinnitus		Glipizide	C		
				Amlodipine	C		
				Simvastatin	C		
				Multivitamin	C		

Date:02/01/99ISR Number: 3404293-9Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #A0066231

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG/TWICE PER DAY/ORAL		Agitation	Consumer	Wellbutrin Tablet	PS		ORAL
		Excitability					
				Alprazolam	C		

Date:02/01/99ISR Number: 3404295-2Report Type:Periodic  
Age:11 YR Gender:Male I/FU:I

Company Report #A0066294

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG/TWICE PER DAY/ORAL		Fatigue	Consumer	Wellbutrin Tablet	PS		ORAL
		Hyperaesthesia					

Date:02/01/99ISR Number: 3404297-6Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #A0066317

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Health	Wellbutrin Tablet	PS		ORAL
75 MG/THREE			Professional				
TIMES PER							
DAY/ORAL							

Ascorbic Acid	C
Iron Salt	C
Multivitamin	C
Ginkgo Biloba	C
Fluoxetine	
Hydrochloride	C

Date:02/01/99ISR Number: 3404310-6Report Type:Periodic Company Report #A0061567  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Wellbutrin			
		Hypersensitivity	Professional	(Formulation			
		Oedema Peripheral	Company	Unknown)	PS		ORAL
UNK / UNK			Representative				
/ORAL							

Date:02/01/99ISR Number: 3404313-1Report Type:Periodic Company Report #A0061568  
 Age: Gender:Unknown I/FU:I

Outcome	PT
	Dermatitis
	Hypersensitivity

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Oedema Peripheral

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
UNK/ UNK/		Health Professional Company	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL		Representative				

Date:02/01/99ISR Number: 3404314-3Report Type:Periodic Company Report #A0061569  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Hypersensitivity Oedema Peripheral	Health Professional Company	Wellbutrin (Formulation Unknown)	PS		ORAL
UNK / UNK/			Representative				
ORAL							

Date:02/01/99ISR Number: 3404315-5Report Type:Periodic Company Report #A0061605  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pollakiuria	Consumer	Wellbutrin Tablet	PS		ORAL
2 TABLET /							
PER DAY /							
ORAL							

Cephalexin C  
 Hypericum C  
 Multivitamin C

Date:02/01/99ISR Number: 3404317-9Report Type:Periodic Company Report #A0061642  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Grand Mal Convulsion	Health Professional Company	Wellbutrin (Formulation Unknown)	PS		ORAL
150 MG / UNK			Representative				
/ ORAL							

Date:02/01/99ISR Number: 3404320-9Report Type:Periodic Company Report #A0061646  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
UNK / UNK /							
ORAL							

Date:02/01/99ISR Number: 3404322-2Report Type:Periodic Company Report #A0061660  
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Interaction	Health Professional	Wellbutrin Tablet	PS		ORAL
37.5 MG /		Leukopenia					
TWICE PER				Fluoxetine Hydrochloride (Formulation			
DAY/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

20 MG / PER  
 DAY/ ORAL  
 Unknown) SS ORAL

Date:02/01/99ISR Number: 3404324-6Report Type:Periodic Company Report #A0061763  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Face Oedema	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

UNK / UNK /

ORAL

Date:02/01/99ISR Number: 3404326-XReport Type:Periodic Company Report #A0061766  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

300 MG / PER

DAY / ORAL

Fluoxetine  
Hydrochloride C

Date:02/01/99ISR Number: 3404328-3Report Type:Periodic Company Report #A0061809  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema	Health Professional	Wellbutrin Tablet	PS		ORAL

75 MG / PER

DAY / ORAL

Date:02/01/99ISR Number: 3404333-7Report Type:Periodic Company Report #A0063405  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea Vertigo	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL	3	WK					

Date:02/01/99ISR Number: 3404336-2Report Type:Periodic Company Report #A0062874  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Insomnia Mental Disorder	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
150 MG/ PER DAY / ORAL		Nausea Nightmare		Methylphenidate Hcl	C		

Date:02/01/99ISR Number: 3404338-6Report Type:Periodic Company Report #A0062964  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Face Oedema	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404339-8Report Type:Periodic Company Report #A0062965  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Prostatitis	Health Professional	Wellbutrin (Formulation Unknown)	PS		

Date:02/01/99ISR Number: 3404343-XReport Type:Periodic Company Report #A0062966  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Metrorrhagia	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

ORAL

Date:02/01/99ISR Number: 3404345-3Report Type:Periodic Company Report #A0062968  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

ORAL

Sertraline Hydrochloride

C

Date:02/01/99ISR Number: 3404348-9Report Type:Periodic Company Report #A0062986  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation Memory Impairment	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

100 MG/

TWICE PER

DAY/ ORAL

Date:02/01/99ISR Number: 3404355-6Report Type:Periodic Company Report #A0062999  
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache Photosensitivity Reaction	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

150 MG/ PER

DAY/ ORAL

Amlodipine C  
Aspirin C

Date:02/01/99ISR Number: 3404356-8Report Type:Periodic Company Report #A0063824  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Health Professional	Wellbutrin (Formulation Unknown)	PS		

75 MG/ TWICE

PER DAY

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404358-1Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #A0063838

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Health Professional Company	Wellbutrin (Formulation Unknown)	PS		ORAL
150 MG / PER DAY / ORAL			Representative	Vicodin Clonazepam	C C		

Date:02/01/99ISR Number: 3404359-3Report Type:Periodic  
Age:34 YR Gender:Male I/FU:I

Company Report #A0063228

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose ORAL		Headache Influenza Like Illness Pharyngolaryngeal Pain	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
				Sertraline Hydrochloride	C		

Date:02/01/99ISR Number: 3404362-3Report Type:Periodic  
Age:27 YR Gender:Male I/FU:I

Company Report #A0063229

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose ORAL		Dermatitis Hypersensitivity Insomnia	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
		Night Sweats Pain Pruritus Stevens-Johnson Syndrome Swelling					

Date:02/01/99ISR Number: 3404363-5Report Type:Periodic Company Report #A0062890

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ecchymosis	Health	Wellbutrin Tablet	PS		ORAL
PER DAY /		Gingival Bleeding	Professional				
ORAL	1 YR	Platelet Count Decreased	Company Representative				

Date:02/01/99ISR Number: 3404365-9Report Type:Periodic Company Report #A0062891

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ecchymosis	Health	Wellbutrin Tablet	PS		ORAL
PER DAY/ ORAL	3 MON	Gingival Bleeding	Professional				
		Platelet Count Decreased	Company Representative				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404366-0Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0062969

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Akathisia	Health	Wellbutrin Tablet	PS		ORAL
350 MG/ PER			Professional				
DAY/ ORAL							

Date:02/01/99ISR Number: 3404368-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0063094

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG/ PER							
DAY/ ORAL	1 WK			Lithium Salt	C		

Date:02/01/99ISR Number: 3404370-2Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0063128

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Health	Wellbutrin Tablet	PS		ORAL
ORAL							
		International Normalised Ratio Increased	Professional Company	Warfarin Sodium Tablet	SS		ORAL
ORAL			Representative Other				

Date:02/01/99ISR Number: 3404372-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0063269

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Wellbutrin Tablet	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3404373-8Report Type:Periodic Company Report #A0063321  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Motor Dysfunction	Health	Wellbutrin Tablet	PS		ORAL
100 MG/ TWICE			Professional				
PER DAY/ ORAL			Company Representative				

Date:02/01/99ISR Number: 3404375-1Report Type:Periodic Company Report #A0063338  
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dissociation	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG/ TWICE		Dry Mouth					
PER DAY/ ORAL		Emotional Disorder		Wellbutrin			
		Insomnia		Tablet-Controlled			
150 MG/ TWICE				Release	SS		ORAL
PER DAY/ ORAL				Hyoscyamine Sulphate	C		
				Tamsulosin Hcl	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404560-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0074188

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK/UNK/ORAL	2 YR	Vitiligo	Consumer	Lamictal Tablet	PS		ORAL
450 MG/PER				Wellbutrin Tablet-Controlled Release	SS		ORAL
DAY/ORAL	3 MON			Methylphenidate Hcl Gabapentin	C C		

Date:02/01/99ISR Number: 3404602-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0073972

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Angioneurotic Oedema	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
				Amlodipine Multiple Medication	C C		

Date:02/01/99ISR Number: 3404604-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0060062

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Chills Pain	Consumer	Wellbutrin (Formulatin Unknown)	PS		ORAL
		Urinary Tract Infection					

Date:02/01/99ISR Number: 3404606-8Report Type:Periodic  
 Age:20 YR Gender:Male I/FU:I

Company Report #A0060069

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG/PER DYA /ORAL		Coordination Abnormal Hypoaesthesia Pyrexia Tachycardia Vomiting	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3404608-1Report Type:Periodic Company Report #A0060083  
 Age:51 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG / TWICE PER DAY / ORAL		Tinnitus	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

Alprazolam C

Date:02/01/99ISR Number: 3404610-XReport Type:Periodic Company Report #A0073393  
 Age:45 YR Gender:Unknown I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia	Health Professional	Wellbutrin (Formulation			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL  
 Unknown) PS ORAL

Date:02/01/99ISR Number: 3404614-7Report Type:Periodic Company Report #A0073396  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tic Tremor	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
300 MG /PER							
DAY /ORAL				Paroxetine Hydrochloride	C		

Date:02/01/99ISR Number: 3404616-0Report Type:Periodic Company Report #A0073521  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased Dizziness Panic Reaction	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL		Urticaria					

Date:02/01/99ISR Number: 3404618-4Report Type:Periodic Company Report #A0073555  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Metrorrhagia	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3404619-6Report Type:Periodic Company Report #A0073664  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abnormal Faeces Irritability	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL

ORAL

Date:02/01/99ISR Number: 3404621-4Report Type:Periodic Company Report #A0073738  
 Age:18 YR Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash Erythematous	Health Professional	Wellbutrin (Formulatin Unknown)	PS		ORAL
300 MG / PER DAY / ORAL							
					Multiple Medication	C	

Date:02/01/99ISR Number: 3404623-8Report Type:Periodic Company Report #A0073782  
 Age:36 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Menstruation Irregular	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
100 MG /TWICE PER DAY / ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404626-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0059499

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash Pruritic	Health Professional Company Representative	Wellbutrin (Formulation Unknown)	PS		

Date:02/01/99ISR Number: 3404627-5Report Type:Periodic  
Age:62 YR Gender:Male I/FU:I

Company Report #A0059970

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alcohol Interaction Convulsion	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
				Ethanol (Formulation Unknown)	SS		

Date:02/01/99ISR Number: 3404628-7Report Type:Periodic  
Age:54 YR Gender:Male I/FU:I

Company Report #A0059972

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypertension	Health Professional	Wellbutrin (Formulation Unknown)	PS		

Date:02/01/99ISR Number: 3404629-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0059974

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation Hypoaesthesia Paraesthesia	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

300 MG /PER

DAY / ORAL

Date:02/01/99ISR Number: 3404631-7Report Type:Periodic Company Report #A0059464  
Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Wellbutrin Tablet	PS		
75 MG / PER		Influenza Like Illness					
DAY / ORAL	5	Tremor		Docusate	C		
	DAY	Visual Disturbance					

Date:02/01/99ISR Number: 3404632-9Report Type:Periodic Company Report #A0059980  
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nervousness	Consumer	Wellbutrin Tablet	PS		ORAL
150 MG /PER							
DAY /ORAL				Blood Pressure Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404634-2Report Type:Periodic  
 Age:26 YR Gender:Female I/FU:I

Company Report #A0059987

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG /PER DAY / ORAL		Constipation	Consumer	Wellbutrin Tablet	PS		ORAL
300 MG /PER DAY / ORAL		Urticaria		Wellbutrin Tablet-Controlled Release	SS		ORAL
				Birth Control	C		

Date:02/01/99ISR Number: 3404636-6Report Type:Periodic  
 Age:34 YR Gender:Female I/FU:I

Company Report #A0073945

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG /PER DAY/ ORAL		Anxiety	Consumer	Wellbutrin Tablet	PS		ORAL
		Confusional State		Paroxetine Hydrochloride	C		
		Dry Mouth Headache Insomnia Nausea Nightmare Visual Acuity Reduced					

Date:02/01/99ISR Number: 3404637-8Report Type:Periodic  
 Age:39 YR Gender:Female I/FU:I

Company Report #A0073958

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
THREE TIMES PER DAY/ORAL		Visual Disturbance		Wellbutrin Tablet	PS		ORAL
				Semisodium Valproate	C		

Fluticasone  
Propionate C  
Beclomethasone  
Dipropion C

Date:02/01/99ISR Number: 3404639-1Report Type:Periodic Company Report #A0074208  
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG/ TWICE		Insomnia					
PER DAY/ ORAL				Piroxicam	C		

Date:02/01/99ISR Number: 3404641-XReport Type:Periodic Company Report #A0074256  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred	Health	Wellbutrin Tablet	PS		ORAL
100 MG /			Professional				
TWICE PER							
DAY/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404643-3Report Type:Periodic  
Age:35 YR Gender:Male I/FU:I

Company Report #A0074390

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Tablet	PS		ORAL
150 MG/ THREE		Drug Interaction	Professional				
TIMES PER DAY		Primary Cerebellar					
ORAL		Degeneration					
		Tremor		Lithium Salt (Formulation Unknown)	SS		

Date:02/01/99ISR Number: 3404645-7Report Type:Periodic  
Age:11 YR Gender:Male I/FU:I

Company Report #A0074422

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Tablet	PS		ORAL
75 MG/ THREE		Arthralgia	Professional				
TIMES PER		Dermatitis					
DAY/ ORAL		Pruritus					
		Rash Maculo-Papular		Clonazepam Dexamphetamine Sulphate	C C		

Date:02/01/99ISR Number: 3404646-9Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0074827

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Tablet	PS		ORAL
75 MG/ THREE		Amnesia	Professional				
TIMES PER DAY			Company				
/ ORAL			Representative	Risperidone	C		

Date:02/01/99ISR Number: 3404648-2Report Type:Periodic Company Report #A0074909  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Burning Sensation	Consumer	Wellbutrin Tablet	PS		ORAL
150 MG/ TWICE		Paraesthesia					
PER DAY/ ORAL				Amitriptyline Hcl	C		

Date:02/01/99ISR Number: 3404649-4Report Type:Periodic Company Report #A0074910  
Age:10 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Condition Aggravated	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG/ ORAL		Drug Ineffective Tic		Methylphenidate Hcl	C		

Date:02/01/99ISR Number: 3404651-2Report Type:Periodic Company Report #A0075056  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Asthenia	Health	Wellbutrin Tablet	PS		ORAL
100 MG/ TWICE		Fatigue	Professional				
PER DAY /		Rhabdomyolysis					
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404652-4Report Type:Periodic  
Age:82 YR Gender:Male I/FU:I

Company Report #A0075307

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fear	Consumer	Wellbutrin Tablet	PS		ORAL
150 MG/ PER DAY/ ORAL		Hyperhidrosis		Sertraline Hydrochloride	C		

Date:02/01/99ISR Number: 3404653-6Report Type:Periodic  
Age:28 YR Gender:Male I/FU:I

Company Report #A0075343

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG/ THREE TIMES PER DAY ORAL				Zyban Tablet - Zyban	SS		ORAL
150 MG/ TWICE PER DAY/ ORAL				Amoxicillin	C		
				Prednisone	C		
				Acyclovir	C		
				Salbutamol Sulphate	C		
				Ipratropium Bromide	C		
				Diazepam	C		

Date:02/01/99ISR Number: 3404660-3Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0074234

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Health Professional Company	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

150 MG/ ORAL			Representative	Zyban Tablet - Zyban	C		ORAL
Date:02/01/99ISR Number: 3404661-5Report Type:Periodic Company Report #A0074237							
Age:51 YR Gender:Female I/FU:I							
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Anxiety Dysgeusia	Health Professional Company	Wellbutrin (Formulation Unknown)	PS		ORAL
150 MG/ ORAL			Representative	Zyban Tablet - Zyban	SS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:02/01/99ISR Number: 3404663-9Report Type:Periodic Company Report #A0074391							
Age: Gender:Unknown I/FU:I							
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Dermatitis Tremor	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404664-0Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0074393

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoaesthesia Hypoaesthesia Oral Miosis	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL							

Date:02/01/99ISR Number: 3404665-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0074512

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysmenorrhoea	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3404668-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0074709

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							
			Other	Nicotine	C		

Date:02/01/99ISR Number: 3404669-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0074823

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Erythematous Urticaria	Health Professional	Wellbutrin (Formulation Unknown)	PS		

TOPICAL 21 MG TOPICAL

Nicotine Patch

SS

Date:02/01/99ISR Number: 3404670-6Report Type:Periodic Company Report #A0074908  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia Areata	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL	2 WK			Nicotine	C		

Date:02/01/99ISR Number: 3404671-8Report Type:Periodic Company Report #A0075085  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lymphoproliferative Disorder	Health Professional	Wellbutrin (Formulation Unknown)	PS		
				Sertraline Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3404673-1Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0075264

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Vision Blurred	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3405406-5Report Type:Periodic  
 Age:41 YR Gender:Male I/FU:I

Company Report #A0058693

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ SINGLE DOSE/ ORAL		Anxiety Apathy Hallucination Hypoaesthesia Nausea Paranoia Tremor	Consumer	Wellbutrin Tablet   Paroxetine Hydrochloride Alprazolam Ethanol	PS   C C C		ORAL

Date:02/01/99ISR Number: 3405408-9Report Type:Periodic  
 Age:71 YR Gender:Female I/FU:I

Company Report #A0058828

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG/PER DAY/ ORAL		Headache Influenza Like Illness Insomnia Tremor	Consumer	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3405410-7Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:I

Company Report #A0058930

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
SEE TEXT/ORAL			Aortic Valve Incompetence	Health	Wellbutrin Tablet	PS		ORAL
VARIABLE			Cardiac Valve Disease Mitral Valve Incompetence Palpitations	Professional Other	Phentermine (Formulation Unknown)	SS		ORAL
DOSE/ ORAL			Tricuspid Valve Incompetence		Fenfluramine Hcl (Formulation Unknown)	SS		ORAL
20 MG/ THREE TIMES PER DAY/ ORAL					Dexfenfluramine Capsule	SS		ORAL
15 MG/ TWICE PER DAY/ ORAL					Phentermine Hydrochloride (Formulation Unknown)	SS		ORAL
30 MG/ PER DAY/ ORAL								

Date:02/01/99ISR Number: 3405411-9Report Type:Periodic Company Report #A0058964  
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL			Anorgasmia Urticaria	Consumer	Wellbutrin Tablet	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3405412-0Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0058976

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oedema	Consumer	Wellbutrin Tablet	PS		ORAL
		Pain In Extremity		Nefazodone Hydrochloride	C		
				Loratadine	C		

Date:02/01/99ISR Number: 3405413-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0058989

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Confusional State	Consumer	Wellbutrin Tablet	PS		ORAL
		Dizziness					
		Dry Mouth					
		Elevated Mood					
		Insomnia					
		Nightmare					
		Sedation					

Date:02/01/99ISR Number: 3405416-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0059236

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pollakiuria	Health	Wellbutrin Tablet	PS		ORAL
		Premature Ejaculation	Professional				
			Company Representative				

Date:02/01/99ISR Number: 3405417-XReport Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0059285

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Influenza	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG, THREE		Vomiting					
TIMES PER							
DAY, ORAL				Alprazolam	C		
				Zolpidem Tartrate	C		

Date:02/01/99ISR Number: 3405419-3Report Type:Periodic Company Report #A0059444  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vertigo	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG, PER							
DAY, ORAL				Fluoxetine			
				Hydrochloride	C		
				Clonazepam	C		

Date:02/01/99ISR Number: 3405420-XReport Type:Periodic Company Report #A0059370  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Health	Wellbutrin			
		International Normalised	Professional	(Formulation			
		Ratio Increased	Other	Unknown)	PS		ORAL
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Warfarin Sodium  
(Formulation  
Unknown) SS

Date:02/01/99ISR Number: 3405421-1Report Type:Periodic Company Report #A0059383  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3405422-3Report Type:Periodic Company Report #A0059390  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health Professional Company Representative	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3405424-7Report Type:Periodic Company Report #A0059392  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health Professional Company Representative	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3405425-9Report Type:Periodic Company Report #A0059414  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Aggression	Consumer	Wellbutrin			
		Blood Pressure Decreased	Health	(Formulation			
150 MG, THREE		Irritability	Professional	Unknown)	PS		ORAL
TIMES PER		Personality Disorder	Other				
DAY, ORAL	5	MON		Propranolol	C		

Date:02/01/99ISR Number: 3405427-2Report Type:Periodic Company Report #A0059440  
Age:52 YR Gender:Female I/FU:I

Outcome

- PT
- Anxiety
- Arthralgia
- Chest Pain
- Cough
- Dyspnoea
- Haemorrhage
- Hallucination, Auditory
- Influenza Like Illness
- Insomnia
- Joint Stiffness
- Lethargy



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
		Muscle Disorder Myalgia Pain					
100 MG, TWICE PER DAY, ORAL		Palpitations Panic Disorder Pruritus	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
		Rash Generalised					
		Tremor		Amitriptyline Hcl	C		
		Weight Decreased		Lorazepam	C		
				Thyroxine Sodium	C		
				Conjugated Estrogens	C		

Date:02/01/99ISR Number: 3405429-6Report Type:Periodic Company Report #A0059259  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Liver Function Test Abnormal	Health Professional Company	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL			Representative				

Date:02/01/99ISR Number: 3405431-4Report Type:Periodic Company Report #A0059280  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Weight Increased	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL				Paroxetine Hydrochloride	C		
				Sertraline Hydrochloride	C		
				Fluoxetine Hydrochloride	C		
				Bupirone Hydrochloride	C		

Lorazepam	C
Dalmane	C
Simvastatin	C
Famotidine	C
Loratadine	C
Docusate Sodium	C

Date:02/01/99ISR Number: 3405432-6Report Type:Periodic  
 Age:51 YR Gender:Male I/FU:I

Company Report #A0059171

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Dissociation	Consumer Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
			Other	Alprazolam	C		
				Buspirone			
				Hydrochloride	C		
				Temazepam	C		
				Lansoprazole	C		
				Frusemide	C		
				Zolpidem Tartrate	C		
				Potassium Salt	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nortriptyline Hcl C  
 Lortab C  
 Sertraline  
 Hydrochloride C  
 Ipratropium Bromide C  
 Salbutamol Sulphate C  
 Acetylcysteine C  
 Ibuprofen C

Date:02/01/99ISR Number: 3405433-8Report Type:Periodic Company Report #A0058684  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Acne	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL				Lithium Salt	C		

Date:02/01/99ISR Number: 3405435-1Report Type:Periodic Company Report #A0058791  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Gastric Disorder	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3405437-5Report Type:Periodic Company Report #A0057573  
 Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Excitability	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG /							
TWICE PER DAY							

/ ORAL

Omeprazole C  
Dyazide C  
Lorazepam C

Date:02/01/99ISR Number: 3405438-7Report Type:Periodic Company Report #A0057770  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Epistaxis	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG /							
TWICE PER DAY							
/ ORAL							

Date:02/01/99ISR Number: 3405439-9Report Type:Periodic Company Report #A0058424  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Drug Ineffective Libido Decreased	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3405440-5Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0058144

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3405441-7Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0058264

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3405442-9Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0058325

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Testicular Swelling	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL	3 WK						

Date:02/01/99ISR Number: 3405443-0Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0058496

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3405444-2Report Type:Periodic Company Report #A0058604  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dermatitis	Consumer	Wellbutrin			
		Dry Mouth		(Formulation			
ORAL	18	Ecchymosis		Unknown)	PS		ORAL
		Keratoconjunctivitis					
		Sicca					

Date:02/01/99ISR Number: 3405445-4Report Type:Periodic Company Report #A0058667  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Ecchymosis	Health	Wellbutrin			
			Professional	(Formulation			
ORAL			Other	Unknown)	PS		ORAL
				Warfarin Sodium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3405446-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0058578

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperproteinaemia	Health	Wellbutrin Tablet	PS		ORAL
150 MG / PER		Insomnia	Professional				
DAY / ORAL		Jaundice					
		Muscle Spasms					
		Oedema Peripheral					
		Pharyngeal Oedema					
		Urticaria					

Date:02/01/99ISR Number: 3405448-XReport Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #A0058427

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Consumer	Wellbutrin Tablet	PS		ORAL
ORAL		Nausea		Wellbutrin Tablet-Controlled Release	SS		ORAL
ORAL							

Date:02/01/99ISR Number: 3405449-1Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0058326

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG /							
TWICE PER DAY							
/ ORAL				Cimetidine	C		
				Clonidine			
				Hydrochloride	C		

Date:02/01/99ISR Number: 3405451-XReport Type:Periodic Company Report #A0058410  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal	Consumer	Wellbutrin Tablet	PS		ORAL
150 MG /		Hyperhidrosis					
TWICE PER DAY		Insomnia					
/ ORAL		Nausea					

Date:02/01/99ISR Number: 3405452-1Report Type:Periodic Company Report #A0058418  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Health	Wellbutrin Tablet	PS		ORAL
150 MG/ IN			Professional				
THE MORNING /							
ORAL							

Date:02/01/99ISR Number: 3405453-3Report Type:Periodic Company Report #A0057844  
Age: Gender:Female I/FU:I

Outcome	PT
	Agitation
	Crying
	Depression
	Disorientation



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Dizziness Dry Mouth Tinnitus	Report Source	Product	Role	Manufacturer	Route
150 MG /		Urticaria	Consumer	Wellbutrin Tablet	PS		ORAL
TWICE PER DAY							
/ ORAL							

Guaiphenesin	C
Ibuprofen	C

Date:02/01/99ISR Number: 3405454-5Report Type:Periodic Company Report #A0057934  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Health	Wellbutrin Tablet	PS		ORAL
150 MG /		Tremor	Professional				
TWICE PER DAY							
/ ORAL							

Vaginal Haemorrhage

Date:02/01/99ISR Number: 3405455-7Report Type:Periodic Company Report #A0057991  
 Age:75 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction	Consumer	Wellbutrin Tablet	PS		ORAL
150 MG /							
THREE TIMES							
DA / ORAL							

Donepezil Hcl	C
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Date:02/01/99ISR Number: 3405456-9Report Type:Periodic Company Report #A0058002  
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG /		Anxiety					
THREE TIMES		Arthralgia					
PER DA / ORAL		Asthenia					
		Chest Pain					
		Dizziness					
		Faecal Incontinence					
		Insomnia					
		Malaise					
		Movement Disorder					
		Nausea					
		Tongue Ulceration					

Date:02/01/99ISR Number: 3405457-0Report Type:Periodic Company Report #A0058044  
Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation	Health	Wellbutrin Tablet	PS		ORAL
100 MG /		Ear Disorder	Professional				
THREE TIMES							
DA / ORAL				Gabapentin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3405458-2Report Type:Periodic  
 Age:51 YR Gender:Female I/FU:I

Company Report #A0058296

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Emotional Disorder	Consumer	Wellbutrin Tablet	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Simvastatin	C		
				M.V.I.	C		

Date:02/01/99ISR Number: 3405459-4Report Type:Periodic  
 Age:23 YR Gender:Female I/FU:I

Company Report #A0075392

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Acne	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG / TWICE							
PER DAY /							
ORAL							
				Clonazepam	C		

Date:02/01/99ISR Number: 3405461-2Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0075566

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus	Health	Wellbutrin Tablet	PS		ORAL
75 MG / PER							
DAY / ORAL			Professional				

Date:02/01/99ISR Number: 3405462-4Report Type:Periodic  
 Age:13 YR Gender:Unknown I/FU:I

Company Report #A0075605

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Anxiety	Health	Wellbutrin Tablet	PS		ORAL
75 MG / SEE		Chest Pain	Professional				
TEXT / ORAL							

Date:02/01/99ISR Number: 3405464-8Report Type:Periodic Company Report #A0075938  
 Age: Gender:Unknown I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Vision Blurred	Health	Wellbutrin Tablet	PS		ORAL
ORAL			Professional	Wellbutrin Tablet-Controlled Release	SS		ORAL
ORAL							

Date:02/01/99ISR Number: 3405465-XReport Type:Periodic Company Report #A0075991  
 Age:12 YR Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Condition Aggravated	Health	Wellbutrin Tablet	PS		ORAL
75 MG/ TWICE		Tic	Professional				
PER DAY /							
ORAL	6 MON			Dexamphetamine Sulphate (Formulation Unknown)	SS		ORAL
2.5 MG / AT							
NIGHT/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3405466-1Report Type:Periodic  
 Age:75 YR Gender:Male I/FU:I

Company Report #A0076190

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG /		Headache	Consumer	Wellbutrin Tablet	PS		ORAL
TWICE PER DAY							
/ ORAL							

Date:02/01/99ISR Number: 3405467-3Report Type:Periodic  
 Age:46 YR Gender:Male I/FU:I

Company Report #A0076523

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG / PER		Dizziness	Consumer	Wellbutrin Tablet	PS		ORAL
DAY / ORAL							
Drug Interaction							
Influenza Like Illness							
Adderall							
C							

Date:02/01/99ISR Number: 3405469-7Report Type:Periodic  
 Age:40 YR Gender:Male I/FU:I

Company Report #A0078107

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Dermatitis	Health	Wellbutrin Tablet	PS		ORAL
Professional							

Date:02/01/99ISR Number: 3405470-3Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0078280

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Dermatitis	Health	Wellbutrin Tablet	PS		ORAL
Professional							

Date:02/01/99ISR Number: 3405471-5Report Type:Periodic Company Report #A0078281  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Wellbutrin Tablet	PS		ORAL
ORAL			Professional	Wellbutrin Tablet-Controlled Release	SS		ORAL
ORAL							

Date:02/01/99ISR Number: 3405472-7Report Type:Periodic Company Report #A0078531  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG / TWICE		Dizziness					
PER DAY/ ORAL		Drug Interaction		Guaiphenesin (Formulation Unknown)	SS		
ORAL				Guaiphenesin Caplet	SS		ORAL
				Nicotine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3405474-0Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0078282

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

ORAL

Date:02/01/99ISR Number: 3405475-2Report Type:Periodic  
 Age:13 YR Gender:Male I/FU:I

Company Report #A0078111

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Cough	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

ORAL

Date:02/01/99ISR Number: 3405477-6Report Type:Periodic  
 Age:53 YR Gender:Female I/FU:I

Company Report #A0077288

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypertension	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

ORAL

Date:02/01/99ISR Number: 3405478-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0076083

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Mydriasis	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

ORAL

Date:02/01/99ISR Number: 3405480-6Report Type:Periodic Company Report #A0076150  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL			Other	Nicotine	C		

Date:02/01/99ISR Number: 3405481-8Report Type:Periodic Company Report #A0075414  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nervousness	Consumer Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL			Other	Nicotine	C		

Date:02/01/99ISR Number: 3405483-1Report Type:Periodic Company Report #A0075507  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source
		Palpitations	Health Professional



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Company Representative	Product	Role	Manufacturer	Route
ORAL			Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3405484-3Report Type:Periodic Company Report #A0075512  
 Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL			Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
				Famotidine	C		
				Atorvastatin Calcium	C		
				Atenolol	C		
				Aspirin	C		

Date:02/01/99ISR Number: 3410842-7Report Type:Periodic Company Report #A0068113  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL			Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
				Diuretic	C		

Date:02/01/99ISR Number: 3410864-6Report Type:Periodic Company Report #A0068255  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Consumer Health	Wellbutrin (Formulation			

ORAL		Professional	Unknown)	PS	ORAL
		Other	Lithium Salt (Formulation Unknown)	SS	
			Nicotine (Formulation Unknown)	SS	

Date:02/01/99ISR Number: 3410867-1Report Type:Periodic Company Report #A0068377  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Micturition Urgency Urinary Retention	Health Professional Company Representative	Wellbutrin (Formulation Unknown)	PS		
2	WK						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3410870-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0068401

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Amnesia Derealisation Disorientation	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3410872-5Report Type:Periodic  
 Age:50 YR Gender:Female I/FU:I

Company Report #A0068402

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG/TWICE PER DAY/ORAL		Back Pain Cough Dyspnoea Gastrointestinal Disorder Pain In Extremity Tinnitus	Consumer	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3410875-0Report Type:Periodic  
 Age:52 YR Gender:Female I/FU:I

Company Report #A0068404

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
TWICE PER DAY/ORAL		Hypercholesterolaemia	Consumer	Wellbutrin Tablet	PS		ORAL
				Lorazepam	C		
				Betaxolol Hydrochloride	C		
				Clonazepam	C		

Date:02/01/99ISR Number: 3410878-6Report Type:Periodic  
 Age:54 YR Gender:Female I/FU:I

Company Report #A0068439

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG/TWICE							
PER DAY/ORAL							

- Isosorbide C
- Nitroglycerin C
- Loratadine C
- Lansoprazole C
- Atenolol C
- Amlodipine C
- Estropipate C
- Medroxyprogesterone
- Ace. C
- Clonazepam C

Date:02/01/99ISR Number: 3410880-4Report Type:Periodic Company Report #A0068797  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Testicular Pain	Health	Wellbutrin Tablet	PS		ORAL
150 MG/ORAL			Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3410887-7Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #A0068810

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Wellbutrin Tablet	PS		ORAL
150 MG/THREE		Rash Erythematous					
TIMES PER		Rash Vesicular					
DAY/ORAL				Lamictal Tablet	SS		ORAL
75 MG/PER							
DAY/ORAL				Nortriptyline	C		
				Thyroid Medication	C		
				Methylphenidate Hcl	C		
				M.V.I	C		
				Lansoprazole	C		
				Sucralfate	C		

Date:02/01/99ISR Number: 3410891-9Report Type:Periodic  
Age:73 YR Gender:Female I/FU:I

Company Report #A0068886

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Difficulty In Walking	Health	Wellbutrin Tablet	PS		ORAL
ORAL		Hypoaesthesia	Professional				

Date:02/01/99ISR Number: 3410893-2Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0068923

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema	Health	Wellbutrin Tablet	PS		ORAL
150 MG/ORAL			Professional				

Date:02/01/99ISR Number: 3410894-4Report Type:Periodic Company Report #A0068969  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG/PER		Dysphagia					
DAY/ORAL		Hypoglycaemia					
		Insomnia					
		Muscle Rigidity					
		Swelling					

Date:02/01/99ISR Number: 3410897-XReport Type:Periodic Company Report #A0069031  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Wellbutrin Tablet	PS		ORAL
ORAL		Tachycardia					

Date:02/01/99ISR Number: 3410899-3Report Type:Periodic Company Report #A0069238  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL	1	MON					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3410901-9Report Type:Periodic Company Report #A0069270  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL	6 DAY						

Date:02/01/99ISR Number: 3410904-4Report Type:Periodic Company Report #A0069271  
 Age:12 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Health	Wellbutrin Tablet	PS		ORAL
1 TABLET/PER		Tremor	Professional				
DAY/ORAL							

Date:02/01/99ISR Number: 3410905-6Report Type:Periodic Company Report #A0069302  
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG/ORAL		Pollakiuria		Pemoline	C		

Date:02/01/99ISR Number: 3410906-8Report Type:Periodic Company Report #A0069304  
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Influenza Like Illness	Consumer	Wellbutrin Tablet	PS		ORAL
150 MG/TWICE		Nausea					
PER DAY/ORAL							

Date:02/01/99ISR Number: 3410909-3Report Type:Periodic Company Report #A0069317  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Unevaluable Event	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
150 MG/ORAL							

Date:02/01/99ISR Number: 3410910-XReport Type:Periodic Company Report #A0069408  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Euphoric Mood	Consumer Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
150 MG/PER			Other				
DAY/ORAL							

Date:02/01/99ISR Number: 3410915-9Report Type:Periodic Company Report #A0064376  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Liver Function Test Abnormal	Health Professional	Wellbutrin ( Formulation Unknown)	PS		ORAL
ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3410918-4Report Type:Periodic Company Report #A0064394  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Agitation Feeling Abnormal Formication	Health Professional Company  Representative	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3410919-6Report Type:Periodic Company Report #A0064538  
 Age: Gender:Unknown I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Drug Interaction Muscle Twitching	Health Professional	Wellbutrin (Formulation Unknown)	PS		
				Paroxetine Hydrochloride	SS		

Date:02/01/99ISR Number: 3410921-4Report Type:Periodic Company Report #A0064593  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Excitability	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3410923-8Report Type:Periodic Company Report #A0064667  
 Age:37 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG / PER		Diarrhoea	Consumer	Wellbutrin Tablet	PS		ORAL

DAY / ORAL  
Dizziness  
Paraesthesia  
Muscle Relaxant  
Pain Medication  
C  
C

Date:02/01/99ISR Number: 3410925-1Report Type:Periodic Company Report #A0064749  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction Haemorrhage Prothrombin Time	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL		Prolonged		Warfarin Sodium	SS		

Date:02/01/99ISR Number: 3410927-5Report Type:Periodic Company Report #A0064754  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression Aphasia Dysphemia	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
100 MG / THREE TIME PER DAY / ORAL		Irritability					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3410929-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0064798

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Aggression	Health Professional Company	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL			Representative	Fluoxetine Hydrochloride	C		

Date:02/01/99ISR Number: 3410931-7Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0064799

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypoaesthesia	Health Professional	Wellbutrin Tablet	PS		ORAL
37.5 MG / PER DAY / ORAL		Hypoaesthesia Oral					
		Muscle Spasms		Verapamil Hydrochloride	C		
		Muscle Twitching		Lisinopril	C		
		Paraesthesia		Aspirin	C		
				Zolpidem Tartrate	C		
				Sulphasalazine	C		
				Vitamin E	C		

Date:02/01/99ISR Number: 3410933-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0064805

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Uterine Haemorrhage	Health Professional Company Representative	Wellbutrin (Formulation Unknown)	PS		

Date:02/01/99ISR Number: 3410934-2Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0064824

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG /		Arthralgia					
THREE TIMES		Bladder Pain					
PER DAY /		Breast Engorgement					
ORAL		Chondropathy		Multivitamin	C		
		Dyspnoea		Conjugated Estrogens	C		
		Fatigue					
		Hormone Level Abnormal					
		Hypertension					
		Joint Swelling					
		Myalgia					
		Oedema Peripheral					
		Pruritus					
		Tachycardia					
		Vaginal Pain					
		Vulvovaginal Dryness					



ORAL	Tooth Discolouration	Health	Wellbutrin	PS	ORAL
		Professional	Atorvastatin Calcium	C	
			Clarithromycin	C	
			Simvastatin	C	

Date:02/01/99ISR Number: 3410946-9Report Type:Periodic Company Report #A0065011  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Health	Wellbutrin	PS		ORAL
ORAL	Motor Dysfunction	Professional Company Representative	Paroxetine Hydrochloride	C		

Date:02/01/99ISR Number: 3410947-0Report Type:Periodic Company Report #A0065016  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Consumer	Wellbutrin	PS		ORAL
ORAL	Fatigue					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3410949-4Report Type:Periodic  
 Age:49 YR Gender:Female I/FU:I

Company Report #A0065149

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	75 MG / TWICE	Sedation	Health	Wellbutrin Tablet	PS		ORAL
	PER DAY /		Professional				
	ORAL			Gabapentin	C		
				Clonazepam	C		

Date:02/01/99ISR Number: 3410951-2Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0065252

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flushing	Health	Wellbutrin Tablet	PS		ORAL
	ORAL		Professional Company Representative				

Date:02/01/99ISR Number: 3410954-8Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0065279

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Wellbutrin Tablet	PS		ORAL
	ORAL		Professional Company Representative				

Date:02/01/99ISR Number: 3410957-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0071612

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Dry Mouth  
Nervous System Disorder  
Sleep Disorder  
Consumer  
Wellbutrin  
(Formulation  
Unknown)  
PS  
ORAL  
TWICE PER DAY  
/ ORAL

Date:02/01/99ISR Number: 3410959-7Report Type:Periodic Company Report #A0071645  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL	6 WK						

Date:02/01/99ISR Number: 3410961-5Report Type:Periodic Company Report #A0071649  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Acne Cystic	Consumer	Wellbutrin Tablet	PS		ORAL
50 MG / TWICE							
PER DAY /							
ORAL				Loratadine	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3410962-7Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0071675

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG /		Irritability					
TWICE PER DAY							
/ ORAL							

Date:02/01/99ISR Number: 3410965-2Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0071678

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Wellbutrin Tablet	PS		ORAL
PER DAY /		Disturbance In Attention					
ORAL		Weight Increased					

Date:02/01/99ISR Number: 3410966-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0071740

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Health	Wellbutrin Tablet	PS		ORAL
100 MG /		Mania	Professional				
THREE TIMES		Psychotic Disorder					
PER DAY /		Tremor					
ORAL				Lithium Salt	C		
				Haloperidol	C		

Date:02/01/99ISR Number: 3410968-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0071768

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased	Consumer	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3410970-6Report Type:Periodic Company Report #A0071904  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression Agitation Alopecia Fatigue Hair Colour Changes	Consumer	Wellbutrin (Formulation Unknown)	PS		

Date:02/01/99ISR Number: 3410971-8Report Type:Periodic Company Report #A0071907  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Wellbutrin Tablet	PS		ORAL

100 MG / PER  
DAY / ORAL

Date:02/01/99ISR Number: 3410972-XReport Type:Periodic Company Report #A0071981  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness Dyskinesia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Feeling Abnormal

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
75 MG / TWICE PER DAY / ORAL		Health Professional	Wellbutrin Tablet	PS		ORAL
			Sertraline Hydrochloride	C		

Date:02/01/99ISR Number: 3410974-3Report Type:Periodic Company Report #A0072240  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eye Movement Disorder	Health Professional Other	Wellbutrin (Formulation Unknown)	PS		ORAL
150 MG / PER DAY / ORAL				Risperidone Tablet	SS		ORAL
.5 MG / TWICE PER DAY / ORAL							

Date:02/01/99ISR Number: 3410976-7Report Type:Periodic Company Report #A0072299  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis	Health Professional	Wellbutrin Tablet	PS		ORAL
75 MG / PER DAY / ORAL				Venlafaxine Hydrochloride Tablet	SS		ORAL
ORAL							

Date:02/01/99ISR Number: 3410977-9Report Type:Periodic Company Report #A0072301  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State	Health Professional Company	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL			Representative				

Date:02/01/99ISR Number: 3410979-2Report Type:Periodic Company Report #A0072613  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain	Health Professional	Wellbutrin Tablet	PS		ORAL
75 MG / TWICE PER DAY /							
ORAL							

Date:02/01/99ISR Number: 3410981-0Report Type:Periodic Company Report #A0072615  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Wellbutrin Tablet	PS		ORAL
150 MG / ORAL			Health Professional	Conjugated Estrogens Tablet	SS		ORAL
625 MG / PER DAY / ORAL			Other	Multivitamin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3410983-4Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0072683

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Medication Error	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/01/99ISR Number: 3410984-6Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0072769

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG / TWICE PER DAY / ORAL		Constipation Dry Mouth Insomnia Irritability Mydriasis Urine Abnormality Urine Odour Abnormal	Health Professional	Wellbutrin Tablet	PS		ORAL

Date:02/01/99ISR Number: 3410985-8Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0072946

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
375 MG / PER DAY / ORAL		Alopecia	Health Professional Company Representative	Wellbutrin	PS		ORAL

Date:02/01/99ISR Number: 3410986-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0072972

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Consumer	Wellbutrin	PS		ORAL
300 MG / PER		Mania					
DAY / ORAL				Conjugated Estrogens	SS		

Date:02/01/99ISR Number: 3410988-3Report Type:Periodic Company Report #A0073114  
 Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Decreased	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG / PER		Gastrointestinal Disorder					
DAY / ORAL		Headache		Semisodium Valproate	C		
		Hypersensitivity		Clonazepam	C		
		Swelling					
		Vision Blurred					

Date:02/01/99ISR Number: 3414870-7Report Type:Periodic Company Report #A0066792  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Wellbutrin Tablet 75	PS		ORAL
2 TABLET /		Pruritus		Mg			
TWICE PER DAY				Salbutamol Sulphate	C		
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3414871-9Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #A0066378

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Abdominal Distension Fatigue Irritability Libido Decreased Malaise Nausea	Consumer	Wellbutrin Tablet   Gemfibrozil Propranolol Hydrochloride Oestradiol Phentermine	PS   C C C C		ORAL

Date:02/01/99ISR Number: 3414872-0Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0066387

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG / THREE TIMES PER DAY/ ORAL		Abdominal Distension Constipation Flatulence Headache Pain	Consumer	Wellbutrin Tablet   No Concurrent Medication	PS   C		ORAL

Date:02/01/99ISR Number: 3414873-2Report Type:Periodic  
Age:73 YR Gender:Male I/FU:I

Company Report #A0066396

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER / DAY/ ORAL		Pollakiuria	Consumer	Wellbutrin Tablet  Lorazepam Glipizide	PS  C C		ORAL

Date:02/01/99ISR Number: 3414874-4Report Type:Periodic Company Report #A0066502  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG / TWICE		Psychomotor Hyperactivity					
PER DAY /							
ORAL							

Date:02/01/99ISR Number: 3414876-8Report Type:Periodic Company Report #A0066559  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Health	Wellbutrin Tablet	PS		ORAL
75 MG/ PER		Agitation	Professional				
DAY / ORAL		Anxiety		No Concurrent Medication	C		

Date:02/01/99ISR Number: 3414877-XReport Type:Periodic Company Report #A0066580  
Age:8 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG / PER		Psychomotor Hyperactivity					
DAY / ORAL				Ascorbic Acid	C		
				Ca Salt + Mg Salt	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3414878-1Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0066634

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Health	Wellbutrin Tablet	PS		ORAL
UNK/ UNK/		Headache	Professional				
ORAL			Company Representative	Wellbutrin Tablet-Controlled Release	SS		ORAL
UNK /UNK /							
ORAL							

Date:02/01/99ISR Number: 3414879-3Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0066718

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Health	Wellbutrin Tablet	PS		ORAL
UNK / UNK/		Pain In Extremity	Professional				
ORAL			Company Representative				

Date:02/01/99ISR Number: 3414880-XReport Type:Periodic  
Age:55 YR Gender:Male I/FU:I

Company Report #A0066850

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Wellbutrin Tablet	PS		ORAL
37.5 MG / PER		Arthralgia					
DAY/ ORAL		Headache		Interferon	C		
		Nasal Congestion		Doxazosin Mesylate	C		
		Nausea		Dexamphetamine	C		
		Sinus Congestion		Vicoprofen	C		
		Tinnitus		Zolpidem Tartrate	C		
		Visual Disturbance		Tinnitus	C		

Nausea

C

Date:02/01/99ISR Number: 3414881-1Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #A0066940

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fibromyalgia	Health	Wellbutrin Tablet	PS		ORAL
150 MG/ THREE		Myalgia	Professional				
TIMES PER DA							
ORAL							

Date:02/01/99ISR Number: 3414882-3Report Type:Periodic  
Age:26 YR Gender:Male I/FU:I

Company Report #A0066996

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gynaecomastia	Health	Wellbutrin Tablet	PS		ORAL
100 MG /			Professional				
THREE TIMES							
PER DA ORAL							

Antihistamine	C
Loratadine	C
Triamcinolone	
Acetonide	C
Budesonide	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3414883-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0066803

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK/ UNK/		Hypersensitivity	Health Professional Other	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3414884-7Report Type:Periodic  
 Age:44 YR Gender:Male I/FU:I

Company Report #A0066826

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK/ UNK/ORAL	1 YR	Drug Interaction Libido Decreased Orgasm Abnormal	Consumer Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
UNK / UNK /		Sedation	Other	Fluoxetine Hydrochloride Tablet	SS		ORAL
ORAL	4 YR						
UNK/ UNK/				Trazodone Tablet	SS		ORAL
ORAL							

Date:02/01/99ISR Number: 3414885-9Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0066783

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK/ UNK/		Dermatitis	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3414886-0Report Type:Periodic Company Report #A0066635  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Visual Acuity Reduced	Health Professional Company	Wellbutrin (Formulation Unknown)	PS		ORAL
UNK/ UNK/			Representative				
ORAL							

Date:02/01/99ISR Number: 3414887-2Report Type:Periodic Company Report #A0066645  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
UNK/ PER DAY							
/ ORAL							

Date:02/01/99ISR Number: 3414888-4Report Type:Periodic Company Report #A0066541  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Visual Disturbance	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
100 MG/ TWICE							
PER DAY/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3414889-6Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0066487

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oedema Peripheral	Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3414890-2Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0066383

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tremor	Health Professional Company Representative	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3414950-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0067680

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Menstruation Irregular	Health Professional	Wellbutrin (Formulation Unknown)	PS		

Date:02/01/99ISR Number: 3414951-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0067712

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity	Consumer	Wellbutrin (Formulation Unknown) Zantac Tablet	PS SS		ORAL
ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
150 MG TWICE PER DAY ORAL							
				Venlafaxine Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3414954-3Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0067091

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Wellbutrin Tablet	PS		ORAL
ORAL			Health Professional Other	Nicotine (Formulation Unknown)	SS		

Date:02/01/99ISR Number: 3414955-5Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #A0067139

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Burning Sensation	Health	Wellbutrin Tablet	PS		ORAL
300 MG ORAL			Professional	Wellbutrin Tablet-Controlled Release	SS		ORAL
ORAL		Dermatitis Flushing Lichen Planus		Corticosteroid	C		

Date:02/01/99ISR Number: 3414956-7Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #A0067225

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Wellbutrin Tablet	PS		ORAL
50 MG PER DAY		Sedation					
ORAL							

Date:02/01/99ISR Number: 3414957-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0067277

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Wellbutrin Tablet	PS		ORAL
ORAL							

Date:02/01/99ISR Number: 3414958-0Report Type:Periodic Company Report #A0067281  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Muscle Contractions	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG THREE		Involuntary					
TIMES PER DAY		Myalgia					
ORAL							
				Oestradiol	C		
				Multivitamin	C		

Date:02/01/99ISR Number: 3414959-2Report Type:Periodic Company Report #A0067326  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness	Health	Wellbutrin Tablet	PS		ORAL
75 MG PER DAY		Vertigo	Professional				
ORAL	3 DAY						

Date:02/01/99ISR Number: 3414960-9Report Type:Periodic Company Report #A0067505  
Age:17 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Interaction	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG TWICE							
PER DAY ORAL							
				Ranitidine			



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride C  
Anesthetic C

Date:02/01/99ISR Number: 3414961-0Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0067533

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dyspnoea	Consumer	Wellbutrin Tablet	PS		ORAL
VARIABLE DOSE							
ORAL							
				Wellbutrin Tablet-Controlled Release	SS		ORAL
150 MG AT							
NIGHT ORAL							

Date:02/01/99ISR Number: 3414962-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0067651

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG TWICE							
PER DAY ORAL		Anxiety					
				Lithium Salt	C		
				Lamotrigine	C		
				Venlafaxine			
				Hydrochloride	C		
				Thyroxine Sodium	C		

Date:02/01/99ISR Number: 3414963-4Report Type:Periodic  
Age:34 YR Gender: I/FU:I

Company Report #A0067653

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Amnesia	Consumer	Wellbutrin Tablet	PS		ORAL
150 MG TWICE							

PER DAY ORAL

Date:02/01/99ISR Number: 3414964-6Report Type:Periodic Company Report #A0067757  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperglycaemia	Consumer	Wellbutrin Tablet	PS		ORAL
1TABLET PER							
DAY ORAL 2 DAY							

Date:02/01/99ISR Number: 3414965-8Report Type:Periodic Company Report #A0067794  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Health	Wellbutrin Tablet	PS		ORAL
150 MG TWICE							
PER DAY ORAL							
				Amitriptyline Hcl	C		

Date:02/01/99ISR Number: 3414966-XReport Type:Periodic Company Report #A0067876  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Wellbutrin Tablet	PS		ORAL
100 MG TWICE							
PER DAY ORAL							
				Methylphenidate Hcl	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/99ISR Number: 3414967-1Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #A0067885

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG TWICE							
PER DAY ORAL							
				Wellbutrin Tablet-Controlled Release	SS		ORAL
150 MG TWICE							
PER DAY ORAL							

Date:02/01/99ISR Number: 3414968-3Report Type:Periodic  
Age:61 YR Gender:Male I/FU:I

Company Report #A0067891

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lethargy	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG THREE							
TIMES PER DAY							
ORAL							
				Piroxicam	C		
				Hyzaar	C		
				Quinine Sulphate	C		
				Multivitamin	C		
				Amitriptyline Hcl	C		

Date:02/01/99ISR Number: 3414969-5Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #A0067960

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred	Consumer	Wellbutrin Tablet	PS		ORAL
75 MG THREE							
TIMES PER DAY							

ORAL

Lorazepam

C

Date:02/02/99ISR Number: 3190464-6Report Type:Direct  
Age:45 YR Gender: I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged		Grand Mal Convulsion	Zyban	PS		

Date:02/04/99ISR Number: 3192336-XReport Type:Expedited (15-DaCompany Report #A0065665  
Age:45 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 75 MG/TWICE PER DAY/ ORAL		Health Professional	Wellbutrin	PS		ORAL
	Convulsion					
	Dermatitis		Clonazepam	C		
	Dry Mouth		Carisoprodol	C		
	Jaundice		Botulinum Toxin	C		
	Loss Of Consciousness					
	Weight Decreased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/04/99ISR Number: 3196585-6Report Type:Periodic  
Age:41 YR Gender:Male I/FU:I

Company Report #M087478

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Grand Mal Convulsion	Health	Serzone Tabs	PS		ORAL
150 MG QHR			Professional				
ORAL				Wellbutrin	SS		ORAL
150 MG BID							
ORAL							

Date:02/04/99ISR Number: 3197911-4Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0058679

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Burning Sensation	Health	Wellbutrin	PS		ORAL
100 MG ORAL			Professional				
		Dry Skin					
		Dyspnoea					
		Eyelid Oedema					
		Face Oedema					
		Pharyngeal Oedema					
		Pruritus					
		Tongue Oedema					
		Urticaria					

Date:02/04/99ISR Number: 3197918-7Report Type:Periodic  
Age:32 YR Gender:Male I/FU:I

Company Report #A0058587

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Intentional Misuse	Health	Wellbutrin	PS		ORAL
ORAL			Professional				
Initial or Prolonged		Loss Of Consciousness		Zidovudine	C		
		Suicide Attempt		Lamivudine	C		
				Indinavir Sulfate	C		
				Trazodone	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Tablet	PS		ORAL
75 MG	TWICE		Professional				
PER DAY	ORAL						
				Donepezil Hcl	C		
				Quinapril			
				Hydrochloride	C		
				Paroxetine			
				Hydrochloride	C		
				Aspirin	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Amnesia	Health	Wellbutrin Tablet	PS		ORAL
150 MG	THREE		Professional				
TIMES PER DAY		Choking					
ORAL		Confusional State					
		Convulsion		Sertraline			
				Hydrochloride	C		
				Tamoxifen	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/04/99ISR Number: 3296857-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #M076943

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Health	Serzone Tabs	PS		ORAL
300-400 MG		Drug Interaction	Professional				
ORAL				Wellbutrin	SS		ORAL
ORAL							

Date:02/04/99ISR Number: 3297053-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #M089705

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyskinesia	Health	Serzone Tabs	PS		ORAL
500MG QD ORAL 3 WK			Professional	Wellbutrin	SS		
150MG BID							

Date:02/04/99ISR Number: 3300316-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #M078898

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Serzone Tabs	PS		ORAL
300-400 MG		Drug Interaction					
ORAL				Wellbutrin	SS		ORAL
ORAL							

Date:02/04/99ISR Number: 3300318-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #M078899

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Health	Serzone Tabs	PS		ORAL
300-400 MG							

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL				Professional				
ORAL					Wellbutrin	SS		ORAL
Date:02/04/99ISR Number: 3300489-5Report Type:Periodic Company Report #M083831								
Age: Gender:Female I/FU:I								
ORAL	50 MG	QD	Condition Aggravated	Consumer	Serzone Tabs	50 Mg	PS	ORAL
ORAL		2 WK	Tremor					
ORAL					Wellbutrin	SS		ORAL
					Zocor	C		
					Dayhist	C		
Date:02/04/99ISR Number: 3303157-9Report Type:Periodic Company Report #M082468								
Age: Gender: I/FU:I								
ORAL		2 MON	Dermatitis	Health	Serzone Tabs		PS	ORAL
ORAL		1 MON	Pruritus	Professional	Wellbutrin		SS	ORAL
Date:02/04/99ISR Number: 3303350-5Report Type:Periodic Company Report #M075277								
Age: Gender:Female I/FU:I								
ORAL	100 MG	BID	Chills	Consumer	Serzone Tabs	100 Mg	PS	ORAL
ORAL			Paraesthesia					
ORAL			Vasodilatation		Wellbutrin		SS	ORAL
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/04/99ISR Number: 3303662-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #M078897

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300-400 MG		Dizziness	Health	Serzone Tabs	PS		ORAL
ORAL		Drug Interaction	Professional				
ORAL				Wellbutrin	SS		ORAL

Date:02/04/99ISR Number: 3303735-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #M077598

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Galactorrhoea	Health	Serzone Tabs	PS		ORAL
ORAL			Professional	Wellbutrin	SS		ORAL

Date:02/04/99ISR Number: 3304353-7Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #M081635

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
50-100 MG BID		Condition Aggravated	Health	Serzone Tabs	PS		ORAL
ORAL	12 DAY	Emotional Disorder	Professional				
300 MG HS		Insomnia		Wellbutrin	SS		ORAL
ORAL		Libido Increased					

Date:02/04/99ISR Number: 3305811-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #M079632

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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50 MG ORAL; 1	Amblyopia	Consumer	Serzone Tabs 50mg	PS	ORAL
DOSES	Headache				
ORAL	Nausea		Wellbutrin	SS	ORAL

Date:02/04/99ISR Number: 3405202-9Report Type:Periodic Company Report #98USA11257  
 Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
DAILY, ORAL		Hallucination Insomnia Overdose	Health Professional	Ritalin Tablet (Methylphenidate Hydrochloride)	PS		ORAL
RESPIRATORY (INHALATION)	INHALATION	Psychotic Disorder		Marijuana (Cannabis)	SS		
ORAL				Wellbutrin Tablet (Amfebutamone Hydrochloride)	SS		ORAL

Date:02/05/99ISR Number: 3192891-XReport Type:Expedited (15-DaCompany Report #A0061783  
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG/TWICE		Ecchymosis	Health	Zyban	PS		ORAL
PER DAY/ ORAL		Thrombocytopenia	Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/08/99ISR Number: 3193360-3Report Type:Direct  
Age:49 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 150MG PO BID		Rash Erythematous		Zyban 150mg	PS	Glaxo Wellcome	ORAL
Intervention to Prevent Permanent Impairment/Damage		Rash Pruritic					

Date:02/08/99ISR Number: 3193901-6Report Type:Expedited (15-DaCompany Report #A0081197  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG TWICE		Dehydration	Foreign	Zyban	PS		ORAL
Initial or Prolonged PER DAY ORAL		Kidney Infection	Consumer				
		Pneumonia		Fluoxetine Hydrochloride	C		

Date:02/08/99ISR Number: 3193902-8Report Type:Expedited (15-DaCompany Report #A0081366  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150 MG PER		Amblyopia	Foreign	Zyban	PS		ORAL
DAY ORAL		Diplopia	Health				
		Dizziness Fall Visual Disturbance	Professional	Oral Contraceptive	C		

Date:02/10/99ISR Number: 3194839-0Report Type:Direct  
Age:55 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 2 MG DAILY Initial or Prolonged ORAL	Back Pain Blood Creatine Phosphokinase Increased	Perphenazine	PS	ORAL
150 MG BID ORAL	Chills Dyspnoea Neuroleptic Malignant Syndrome Pyrexia Respiratory Distress Sepsis	Bupropion	SS	ORAL

Date:02/10/99ISR Number: 3195526-5Report Type:Expedited (15-DaCompany Report #A0081113  
Age:50 YR Gender:Female I/FU:I

Outcome Hospitalization - Initial or Prolonged	PT Amnesia Dysgraphia Dysphagia Facial Palsy Hearing Impaired Hemiparesis Hypoaesthesia Pruritus Speech Disorder Transient Ischaemic Attack
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Urticaria

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Consumer	Wellbutrin	PS		ORAL

Date:02/10/99ISR Number: 3195529-0Report Type:Expedited (15-DaCompany Report #A0077703  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG TWICE PER DAY ORAL		Agitation Convulsion Nausea Oral Intake Reduced Staring	Health Professional Company Representative	Wellbutrin	PS		ORAL

Date:02/11/99ISR Number: 3195948-2Report Type:Expedited (15-DaCompany Report #A0081071  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Respiratory Distress Urticaria	Health Professional	Wellbutrin Aspirin	PS C		

Date:02/11/99ISR Number: 3196526-1Report Type:Periodic Company Report #A0063369  
Age:14 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG / PER DAY / ORAL		Grand Mal Convulsion	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL

Date:02/12/99ISR Number: 3197976-XReport Type:Expedited (15-DaCompany Report #A0080414  
Age:34 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG/TWICE	Arthralgia	Foreign	Zyban	PS		ORAL
Initial or Prolonged PER DAY/ORAL	Oedema Peripheral	Health				
Disability	Serum Sickness Swelling Urticaria	Professional	Lansoprazole Cisapride	C C		

Date:02/12/99ISR Number: 3198077-7Report Type:Expedited (15-DaCompany Report #A0081119  
Age:32 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG TWICE	Arthralgia	Health	Zyban Tablet - Zyban	PS		ORAL
Initial or Prolonged PER DAY ORAL 1 WK	Difficulty In Walking	Professional				
Disability	Dry Skin Leukocytosis Oedema Peripheral Pruritus Serum Sickness Urticaria	Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/16/99ISR Number: 3198497-0Report Type:Direct  
 Age:35 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG BID PO Initial or Prolonged		Arthralgia	Health Professional	Zyban	PS		ORAL

Date:02/16/99ISR Number: 3198559-8Report Type:Direct  
 Age:17 YR Gender: I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 200MG PO Q AM		Fatigue		Wellbutrin	PS		ORAL

Date:02/16/99ISR Number: 3198585-9Report Type:Direct  
 Age:50 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG 2X'S Other ORAL		Anxiety Arthralgia Arthritis Depression Joint Swelling Muscle Rigidity Muscle Spasms Oedema Peripheral Pain		Zyban	PS	Glaxo Wellcome	ORAL

Date:02/16/99ISR Number: 3199364-9Report Type:Expedited (15-DaCompany Report #A0081946  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - ORAL	Abdominal Pain Upper	Consumer	Zyban Tablet - Zyban	PS	ORAL
Initial or Prolonged	Decreased Appetite Diarrhoea Dysgeusia Hepatitis Hypersensitivity Jaundice				

Date:02/17/99ISR Number: 3200550-XReport Type:Expedited (15-DaCompany Report #9904510  
 Age:14 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 50.00 MG Hospitalization - TOTAL:DAILY	Depression	Consumer	Zoloft Tablets	PS		ORAL
Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage	Drug Ineffective Suicide Attempt		Prozac Effexor Wellbutrin Depakote Buspar Unknown Medications	SS SS SS C C C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/18/99ISR Number: 3203086-5Report Type:Expedited (15-DaCompany Report #A0062241

Age: Gender: I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly	Duration Complications Of Maternal Exposure To Therapeutic Drugs	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/TWICE PER DAY ORAL	Exomphalos Heart Disease Congenital					

Date:02/18/99ISR Number: 3203091-9Report Type:Expedited (15-DaCompany Report #A0081061

Age:26 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - SINGLE	Duration Convulsion	Health	Wellbutrin Tablet	PS		ORAL
Initial or Prolonged DOSE/ORAL	Electrocardiogram Qt	Professional				
SINGLE	Prolonged Intentional Misuse	Other	Antidepressant Tablet	SS		ORAL
DOSE/ORAL	Suicide Attempt		Ethanol (Formulation Unknown)	SS		

Date:02/18/99ISR Number: 3203094-4Report Type:Expedited (15-DaCompany Report #A0082079

Age:69 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG/PER	Duration Balance Disorder	Consumer	Zyban Tablet - Zyban	PS		ORAL
Initial or Prolonged DAY/ORAL	Confusional State					
	Feeling Abnormal Nausea		Vitamin E Calcium Salt	C C		

Date:02/19/99ISR Number: 3203570-4Report Type:Expedited (15-DaCompany Report #A0074981  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Health	Wellbutrin			
		Haematoma	Professional	Tablet-Controlled			
		International Normalised	Company	Release	PS		ORAL
150MG/ORAL							
		Ratio Increased	Representative	Warfarin Sodium			
		Joint Swelling		Tablet	SS		ORAL
10MG/ALTERNAT							
		Pain In Extremity					
E DAYS/ORAL							
		Prothrombin Time Ratio		Warfarin Sodium			
		Increased		Tablet	SS		ORAL
7.5							
MG/ALTERNATE							
DAYS/ORAL							

Date:02/22/99ISR Number: 3204842-XReport Type:Expedited (15-DaCompany Report #A0080063  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Dizziness	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE							
		Insomnia	Professional				
PER DAY/ORAL							
		Muscle Rigidity		Multivitamin	C		
		Muscle Spasms					
		Myalgia					
		Nervousness					
		Panic Attack					
		Visual Disturbance					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/22/99ISR Number: 3204849-2Report Type:Expedited (15-DaCompany Report #A0082610  
Age:55 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY ORAL	Angioneurotic Oedema	Foreign Health Professional	Zyban Tablet - Zyban	PS		ORAL
	Rash Erythematous Tongue Oedema					

Date:02/22/99ISR Number: 3204908-4Report Type:Expedited (15-DaCompany Report #A0082630  
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL Initial or Prolonged	Hypoaesthesia	Health Professional Company Representative	Zyban Tablet - Zyban	PS		ORAL

Date:02/22/99ISR Number: 3204910-2Report Type:Expedited (15-DaCompany Report #A0082661  
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL Initial or Prolonged	Colitis Diarrhoea Haemorrhagic	Health Professional Company Representative	Zyban Tablet - Zyban	PS		ORAL

Date:02/22/99ISR Number: 3204944-8Report Type:Expedited (15-DaCompany Report #A0079791  
Age:64 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG /	Aneurysm	Consumer	Zyban Tablet - Zyban	PS		ORAL

Initial or Prolonged Cerebrovascular Accident  
TWICE PER DAY

Hemiplegia

/ ORAL

Digoxin	C
Frusemide	C
Diclofenac Sodium	C
Warfarin Sodium	C

Date:02/22/99ISR Number: 3204956-4Report Type:Expedited (15-DaCompany Report #A0081197

Age:56 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY	Chills Dehydration Kidney Infection	Foreign Health Professional	Zyban Tablet - Zyban	PS		ORAL
/ ORAL	Malaise Myalgia Oliguria Pneumonia Pyelonephritis Pyrexia		Fluoxetine Hydrochloride	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/22/99ISR Number: 3205582-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #8-98348-072A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Health	Effexor Xr			
300 MG;	ONCE	Condition Aggravated	Professional	Capsules	PS		ORAL
DAILY;	ORAL	Depression					
300 MG;		Myalgia		Wellbutrin			
DAILY;	ORAL	Thinking Abnormal		(Bupropion)	SS		ORAL

Date:02/22/99ISR Number: 3401571-4Report Type:Periodic  
Age:31 YR Gender:Male I/FU:I

Company Report #WAES 98030804

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/		Drug Interaction	Consumer	Tab Zocor Unk	PS		
BID/PO				Tab Zyban 150 Mg	SS		ORAL

Date:02/23/99ISR Number: 3202972-XReport Type:Expedited (15-DaCompany Report #A0077674  
Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Health	Zyban Tablet - Zyban	PS		ORAL
UNKNOWN	ORAL		Professional				

Date:02/23/99ISR Number: 3204642-0Report Type:Expedited (15-DaCompany Report #MPI-97418(1)  
Age:9 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged	Aggression Agitation Intermittent Explosive Disorder	Health Professional	Methylphenidate Tablets 10 Mg (Methylphenidate Hydrochloride 10 Mg)	PS	ORAL
55 MG, PO	Memory Impairment Parent-Child Problem		Clonidine Hydrochloride (Clonidine Hydrochloride)	SS	ORAL
0.1 MG PO			Bupropion (Amfebutamone)	SS	ORAL
150 MG PO					

Date:02/24/99ISR Number: 3206129-8Report Type:Direct  
Age:47 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Angioneurotic Oedema Oedema Peripheral		Zyban - 150 Mg Tab (Glaxo Wellcome)	PS	Glaxo Wllome	ORAL
150MG BY MOUTH DAILY	10 DAY	Pain In Extremity Pruritus Rash Erythematous Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/24/99ISR Number: 3427033-6Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #A0076399

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gingival Bleeding	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Insomnia					
PER DAY ORAL				Nitroglycerin	C		

Date:02/25/99ISR Number: 3208776-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #9835125

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zoloft Tablets	PS		ORAL
Other		Erectile Dysfunction					
25 MG TOTAL		Libido Decreased		Wellbutrin	SS		ORAL
DAILY ORAL							
ORAL							

Date:02/25/99ISR Number: 3208806-1Report Type:Periodic  
Age:26 YR Gender:Female I/FU:F

Company Report #9719922

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Health	Zoloft	PS		ORAL
Other		Cardiovascular Disorder	Professional	Redux	SS		ORAL
ORAL							
30.00 MG							
TOTAL:DAILY:0							
RAL				Ambien	SS		ORAL
ORAL				Wellbutrin	SS		ORAL
ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain	Consumer	Zoloft Tablets	PS		ORAL
DAILY:ORAL		Agitation		Wellbutrin	SS		ORAL
ORAL		Anxiety		Remeron	SS		ORAL
ORAL		Arthralgia		Vitamins	C		
		Coordination Abnormal		Xanax	C		
		Dizziness					
		Drug Ineffective					
		Erectile Dysfunction					
		Increased Appetite					
		Insomnia					
		Libido Decreased					
		Listless					
		Mania					
		Nervousness					
		Oedema Peripheral					
		Paraesthesia					
		Sexual Dysfunction					
		Weight Increased					

Outcome	PT
Other	Abnormal Dreams
	Alopecia
	Constipation
	Depersonalisation



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100.00 MG		Dizziness Drug Ineffective Insomnia	Consumer	Zoloft Tablets	PS		ORAL
TOTAL: DAILY:		Male Sexual Dysfunction	Health				
ORAL		Nocturia	Professional				
87.50 MG		Weight Decreased		Wellbutrin	SS		
TOTAL: BID							

Date:02/25/99ISR Number: 3210871-2Report Type:Periodic Company Report #9800713  
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
50.00 MG		Dermatitis	Health	Zoloft Tablets	PS		ORAL
TOTAL: DAILY:		Hyperhidrosis	Professional				
ORAL				Wellbutrin	SS		ORAL
ORAL							

Date:02/25/99ISR Number: 3211066-9Report Type:Periodic Company Report #9801452  
 Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100.00 MG		Anorexia	Consumer	Zoloft Tablets	PS		ORAL
TOTAL: DAILY:		Diarrhoea	Health				
ORAL		Nausea	Professional				
				Paxil	SS		
				Wellbutrin	SS		
				Hytrin	C		

Date:02/25/99ISR Number: 3212165-8Report Type:Periodic Company Report #9728129  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Dizziness	Health	Zoloft Tablets	PS		ORAL
ORAL							
20.00 MG		Drug Ineffective	Professional	Prozac	SS		ORAL
		Erectile Dysfunction					
TOTAL:DAILY:0							
RAL							
				Wellbutrin	SS		ORAL
ORAL							
				Parnate	SS		ORAL
ORAL							

Date:02/25/99ISR Number: 3212466-3Report Type:Periodic Company Report #JAUSA-34478  
Age:11 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Muscle Contractions Involuntary	Health Professional	Risperdal (Risperidone)	PS	Janssen	ORAL
.5 MG DAILY							
ORAL/ .5 MG 2							
DAILY ORAL							
				Wellbutrin (Amfebutamone)	SS		ORAL
150 MG DAILY							
ORAL/ 100 MG							
1 DAILY ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/25/99ISR Number: 3213420-8Report Type:Periodic  
 Age:65 YR Gender:Male I/FU:I

Company Report #JAUSA-35125

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia	Health Professional	Risperdal (Risperidone), Janssen Tablet	PS	Janssen	ORAL
ORAL							
				Klonopin (Clonazepam) Tablet	SS		ORAL
ORAL							
				Paxil (Paroxetine)	SS		ORAL
TABLET ORAL							
				Wellbutrin (Amfebutamone)	SS		
TABLET							

Date:02/25/99ISR Number: 3213792-4Report Type:Periodic  
 Age:46 YR Gender:Female I/FU:I

Company Report #9808807

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Consumer	Zoloft Tablets	PS		ORAL
50.00 MG							
		Chest Pain					
TOTAL DAILY							
		Drug Ineffective					
ORAL							
		Dry Mouth		Wellbutrin	SS		ORAL
150.00 MG							
		Headache					
TOTAL DAILY							
ORAL							

Date:02/25/99ISR Number: 3217332-5Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #9810197

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Health Professional	Zoloft Tablets	PS		
		Drug Interaction		Aricept	SS		

Company  
Representative

Wellbutrin

SS

Date:02/25/99ISR Number: 3217596-8Report Type:Periodic Company Report #9807191

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depersonalisation	Health	Zoloft Tablets	PS		ORAL
50.00 MG		Dizziness	Professional				
TOTAL:		Drug Ineffective					
DAILY:ORAL		Ejaculation Disorder		Wellbutrin	SS		ORAL
75.00 MG		Insomnia					
TOTAL:DAILY:O		Thinking Abnormal					
RAL		Vertigo		Depakote	C		

Date:02/25/99ISR Number: 3217904-8Report Type:Periodic Company Report #9810841

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anorgasmia	Consumer	Zoloft Tablets	PS		ORAL
50.00 MG		Depression					
TOTAL; DAILY;		Drug Ineffective					
ORAL		Dyspepsia		Remeron	SS		ORAL
ORAL		Weight Increased		Wellbutrin	SS		ORAL
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3208619-0Report Type:Expedited (15-DaCompany Report #A0081366  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Amblyopia	Foreign	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER DAY / ORAL		Diplopia	Health				
		Dizziness	Professional	Oral Contraceptive	C		
		Fall					
		Visual Disturbance					

Date:02/26/99ISR Number: 3213518-4Report Type:Periodic Company Report #8-98237-039A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Asthenia	Health	Effexor	PS		ORAL
UP TO 400 MG DAILY ORAL; TAPERED FROM 400 MG; 275 MG DAILY; 50	57 DAY	Depression	Professional				
		Drug Withdrawal Syndrome					
		Hypotonia					
		Nausea					
		Nervousness		Wellbutrin (Bupropion)	SS		ORAL
		Trismus		Trazodone Trilafon (Perphenazine)	C		
ORAL					C		

Date:02/26/99ISR Number: 3214455-1Report Type:Periodic Company Report #8-98023-015D  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anorexia	Consumer	Effexor	PS		ORAL
37.5 MG DAILY							

ORAL

Asthenia

Confusional State  
Depression  
Dry Mouth  
Nausea  
Sedation  
Yawning

Klonopin SS  
Wellbutrin Sr SS  
Alesse C

Date:02/26/99ISR Number: 3217191-0Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #8-98089-039A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UP TO 225 MG		White Blood Cell Disorder	Consumer	Effexor	PS		ORAL

DAILY ORAL

ORAL

Wellbutrin Tablet	SS		ORAL
Cortisone Injection			
Epidural	C		
Diuretic	C		
Posicur	C		
Wellbutrin	C		

Date:02/26/99ISR Number: 3424057-XReport Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #001-0945-980438

Outcome	PT	Report Source
Drug Interaction Grand Mal Convulsion		Health Professional Company

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Representative

Dose	Duration	Product	Role	Manufacturer	Route
2400 MG (1200 MG, BID), PER ORAL		Neurontin (Gabapentin)	PS		ORAL
375 (DAILY), PER ORAL		Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL
		Prozac Klonopin	C C		

Date:03/01/99ISR Number: 3208520-2Report Type:Direct  
Age:23 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - TWO TIMES Initial or Prolonged DAILY 1 MON Disability		Grand Mal Convulsion	Consumer	Zyban Bupropion Hydrochloride	PS		

Date:03/01/99ISR Number: 3209111-XReport Type:Expedited (15-DaCompany Report #A0078454  
Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability TABLET, 150MG Other / TWICE PER		Dermatitis Exfoliative Retinal Vein Thrombosis	Foreign Health	Zyban - Zyban	PS		ORAL

DAY / ORAL

Date:03/01/99ISR Number: 3209118-2Report Type:Expedited (15-DaCompany Report #A0076073

Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged TABLET, 150		Electroencephalogram Abnormal Grand Mal Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
Other MG / TWICE		Medication Error	Representative				
PER DAY /							
ORAL				Ethanol (Formulation Unknown)	SS		ORAL
ORAL				Glucophage	C		

Date:03/01/99ISR Number: 3209132-7Report Type:Expedited (15-DaCompany Report #A0081417

Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability TABLET, 150		Hypoaesthesia	Foreign	Zyban Tablet - Zyban	PS		ORAL
MG / TWICE		Myalgia	Health				
PER DAY /		Paraesthesia	Professional				
ORAL							

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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3209137-6Report Type:Expedited (15-DaCompany Report #A0065816

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression	Health	Zyban Tablet - Zyban	PS		ORAL
TABLET, 150		Agitation	Professional				
MG / ORAL		Amnesia		Medroxyprogesterone			
		Anger		Ace.	C		
		Anxiety					
		Breast Pain					
		Disorientation					
		Disturbance In Attention					
		Dizziness					
		Feeling Jittery					
		Fluid Retention					
		Galactorrhoea					
		Insomnia					
		Migraine					
		Nausea					
		Paranoia					
		Psychomotor Hyperactivity					
		Suicidal Ideation					
		Tinnitus					

Date:03/01/99ISR Number: 3209143-1Report Type:Expedited (15-DaCompany Report #A0082234

Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Arthralgia	Foreign	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dysphagia	Health				
TWICE PER DAY		Dyspnoea	Professional				
/ ORAL		Headache	Company				
		Influenza Like Illness	Representative				
		Joint Swelling					
		Pain					
		Pharyngolaryngeal Pain					
		Respiratory Distress					
		Urticaria					

Date:03/01/99ISR Number: 3209455-1Report Type:Expedited (15-DaCompany Report #A0082611  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amenorrhoea	Consumer	Wellbutrin			
		Amnesia		Tablet-Controlled			
		Convulsion		Release	PS		ORAL
1 TABLET/ TWICE PER DAY/ ORAL		Head Injury		Aspirin	C		

Date:03/01/99ISR Number: 3218031-6Report Type:Periodic Company Report #A0074913  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150 MG; Initial or Prolonged TWICE PER DAY; ORAL		Convulsion	Health Professional Company Representative	Zyban	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3218032-8Report Type:Periodic  
 Age:29 YR Gender:Female I/FU:I

Company Report #A0074712

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG; Initial or Prolonged TWICE PER DAY;		Dyspnoea Swelling Urticaria	Consumer	Zyban Tablet	PS		ORAL

Date:03/01/99ISR Number: 3218033-XReport Type:Periodic  
 Age:31 YR Gender:Male I/FU:I

Company Report #A0074630

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG; TWICE PER DAY;		Convulsion	Consumer	Zyban Tablet	PS		ORAL
				Tramadol Hydrochloride Hydrocodone	C C		

Date:03/01/99ISR Number: 3218034-1Report Type:Periodic  
 Age:40 YR Gender:Male I/FU:F

Company Report #A0070651

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG; Initial or Prolonged TWICE PER Other DAY;		Convulsion	Health Professional Company Representative	Zyban Tablet	PS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Zyban Tablet	PS		ORAL
150 MG;		Muscle Twitching	Professional				
TWICE PER		Vision Blurred					
DAY;	ORAL						
				Oxybutynin	C		
				Cisapride	C		
				Carbamazepine	C		
				Gabapentin	C		
				Atorvastatin Calcium	C		
				Setraline			
				Hydrochloride	C		
				Prempro	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anxiety	Health	Zyban Tablet	PS		ORAL
150 MG;		Chest Pain	Professional				
Initial or Prolonged							
TWICE PER							
DAY;	ORAL						
				Conjugated Estrogens	C		
				Medroxyprogesterone			
				Ace.	C		
				Multivitamin	C		
				Vitamin E	C		
				Ascorbic Acid	C		
				Aspirin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3218037-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0082071

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Study	Zyban Tablet	PS		ORAL
150 MG;			Consumer				
ORAL							

Date:03/01/99ISR Number: 3218038-9Report Type:Periodic  
 Age:51 YR Gender:Female I/FU:I

Company Report #A0081611

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dyspnoea	Consumer	Zyban Tablet	PS		ORAL
150 MG;							
Initial or Prolonged		Throat Tightness					
TWICE PER		Urticaria					
DAY; ORAL				Weight Loss Medication	C		

Date:03/01/99ISR Number: 3218039-0Report Type:Periodic  
 Age:31 YR Gender:Male I/FU:I

Company Report #A0070268

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspnoea	Consumer	Zyban Tablet	PS		ORAL
150 MG;							
TWICE PER		Eyelid Oedema					
DAY; ORAL		Face Oedema					
		Pruritus		.....	C		
		Urticaria					

Date:03/01/99ISR Number: 3218040-7Report Type:Periodic  
 Age:32 YR Gender:Male I/FU:I

Company Report #A0080724

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG; Initial or Prolonged TWICE PER DAY; ORAL		Arthralgia Dermatitis Swelling Urticaria	Consumer	Zyban Tablet	PS		ORAL

Date:03/01/99ISR Number: 3218041-9Report Type:Periodic Company Report #A0080138  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Convulsion	Health Professional Company Representative	Zyban Tablet	PS		ORAL

Date:03/01/99ISR Number: 3218042-0Report Type:Periodic Company Report #A0080112  
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG; Initial or Prolonged TWICE PER Other DAY; ORAL		Bronchospasm Chest Discomfort Hypersensitivity Oedema Mouth Urticaria	Health Professional Company Representative	Zyban Tablet Nicotine	PS C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3218043-2Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0079702

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban Tablet	PS		ORAL
Other		Convulsion					
150 MG;		Syncope	Professional				
TWICE PER			Company				
DAY;	ORAL		Representative	Cetirizine			
				Hydrochloride	C		
				Fluoxymesterone	C		

Date:03/01/99ISR Number: 3218044-4Report Type:Periodic  
Age:35 YR Gender:Male I/FU:I

Company Report #A0079559

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban Tablet	PS		ORAL
Hospitalization -		Dermatitis					
150 MG;			Professional				
Initial or Prolonged			Company				
TWICE PER			Representative				
DAY;	ORAL						

Date:03/01/99ISR Number: 3218045-6Report Type:Periodic  
Age:39 YR Gender:Unknown I/FU:I

Company Report #A0079375

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban Tablet	PS		ORAL
Hospitalization -		Dermatitis					
150 MG;			Professional				
Initial or Prolonged		Dyspnoea	Company				
TWICE PER			Representative				
DAY;	ORAL						
	3	DAY					
		Hypersensitivity					
		Oedema Mouth					
		Periorbital Oedema					
		Throat Tightness					
		Urticaria					

Date:03/01/99ISR Number: 3218046-8Report Type:Periodic Company Report #A0079407  
Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Dermatitis	Health	Zyban Tablet	PS		ORAL
150 MG; ORAL						
Initial or Prolonged		Professional				

Date:03/01/99ISR Number: 3218047-XReport Type:Periodic Company Report #A0079414  
Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Dermatitis	Health	Zyban Tablet	PS		ORAL
150 MG;						
Initial or Prolonged		Professional				
ORAL						

Date:03/01/99ISR Number: 3218048-1Report Type:Periodic Company Report #A0079094  
Age:33 YR Gender:Female I/FU:I

Outcome	PT
Life-Threatening	Anaphylactic Reaction
Hospitalization -	Angioneurotic Oedema
Initial or Prolonged	Bronchospasm
	Dyspnoea
	Erythema Multiforme
	Face Oedema
	Oedema Peripheral



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Freedom Of Information (FOI) Report

		Pruritus Tongue Oedema Urticaria	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Health	Zyban	PS		ORAL
150 MG;			Professional				
TWICE PER							
DAY;	ORAL						

Date:03/01/99ISR Number: 3218612-XReport Type:Periodic Company Report #A0078496  
 Age:58 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Convulsion	Health	Zyban Tablet - Zyban	PS		ORAL
Dose			Professional	Doxepin	C		
Hospitalization -	2 MON		Company	Diazepam	C		
ORAL			Representative				
Initial or Prolonged							

Date:03/01/99ISR Number: 3218616-7Report Type:Periodic Company Report #A0078477  
 Age:32 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Hypersensitivity	Consumer	Zyban Tablet - Zyban	PS		ORAL
Dose							
Hospitalization -							
150 MG /TWICE							
Initial or Prolonged							
PER DAY/	ORAL						

Date:03/01/99ISR Number: 3218619-2Report Type:Periodic Company Report #A0078288  
 Age:23 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Grand Mal Convulsion	Consumer	Zyban Tablet - Zyban	PS		ORAL
Dose							
Other							
150 MG/TWICE							
PER DAY/	ORAL						

Date:03/01/99ISR Number: 3218624-6Report Type:Periodic Company Report #A0078186  
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Musculoskeletal Pain	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE			Professional				
Other				Nicotine	C		
PER DAY/ORAL				Iron Salt	C		

Date:03/01/99ISR Number: 3218626-XReport Type:Periodic Company Report #A0078110  
 Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Convulsion	Consumer	Zyban Tablet -			
Initial or Prolonged		Tremor		Zyban	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL							

Date:03/01/99ISR Number: 3218630-1Report Type:Periodic Company Report #A0078076  
 Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source
Hospitalization -	Dermatitis	Health
Initial or Prolonged		Professional
		Company

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Representative

Dose	Duration	Product	Role	Manufacturer	Route
ORAL		Zyban Tablet - Zyban	PS		ORAL

Date:03/01/99ISR Number: 3218633-7Report Type:Periodic Company Report #A0078075  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL Initial or Prolonged			Health  Professional Company Representative	Zyban Tablet - Zyban	PS		ORAL

Date:03/01/99ISR Number: 3218634-9Report Type:Periodic Company Report #A0078073  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG, / Initial or Prolonged ORAL			Health  Professional  Company Representative	Zyban Tablet - Zyban	PS		ORAL

Date:03/01/99ISR Number: 3218635-0Report Type:Periodic Company Report #A0077711  
 Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other  150 MG/TWICE  PER DAY/ORAL			Health Professional  Company	Zyban Tablet - Zyban	PS		ORAL

Representative

Date:03/01/99ISR Number: 3218637-4Report Type:Periodic Company Report #A0077362  
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG /TWICE Initial or Prolonged PER DAY/ORAL		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Nausea		Tramadol			
		Sedation		Hydrochloride	C		
		Weight Decreased		Alprazolam	C		
				Conjugated Estrogens	C		
				Iron Salt	C		
				Multivitamin	C		

Date:03/01/99ISR Number: 3218641-6Report Type:Periodic Company Report #A0077347  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Convulsion	Health	Zyban Tablet - Zyban	PS		ORAL
			Professional Company Representative				

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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3218644-1Report Type:Periodic  
Age:33 YR Gender:Male I/FU:I

Company Report #A0077336

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL	Hypersensitivity	Other	Zyban Tablet - Zyban	PS		ORAL
	Oedema Peripheral					
	Rash Erythematous		Claritin - D	C		
	Urticaria		Finasteride	C		
	Vomiting					

Date:03/01/99ISR Number: 3218648-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0077155

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL	Stevens-Johnson Syndrome	Health	Zyban Tablet - Zyban	PS		ORAL
		Professional				
			Gabapentin Capsule	SS		ORAL

Date:03/01/99ISR Number: 3218649-0Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0077046

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG/TWICE PER DAY/ORAL	Pharyngeal Oedema	Consumer	Zyban Tablet - Zyban	PS		ORAL
	Pruritus					
	Rash Erythematous		Donnatal	C		
	Swelling					
	Throat Tightness					
	Urticaria					

Date:03/01/99ISR Number: 3218653-2Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0076779

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Discomfort	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/	TWICE						
		Face Oedema	Professional				
PER DAY/	ORAL						
		Tongue Oedema					
		Urticaria					

Date:03/01/99ISR Number: 3218657-XReport Type:Periodic Company Report #A0076534  
Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Accidental Overdose	Health	Zyban Tablet - Zyban	PS		ORAL
300 MG/	TWICE						
		Convulsion	Professional				
PER DAY/	ORAL						
		Dermatitis					
		Urticaria					

Date:03/01/99ISR Number: 3218660-XReport Type:Periodic Company Report #A0076427  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Hypersensitivity	Consumer	Zyban Tablet -Zyban	PS		ORAL
ORAL							
Initial or Prolonged							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3218663-5Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0076211

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - ORAL Initial or Prolonged		Consumer	Zyban Tablet -Zyban	PS		ORAL

Date:03/01/99ISR Number: 3218666-0Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #A0075090

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL	Chest Pain Hypoaesthesia Myocardial Infarction	Consumer	Zyban Tablet -Zyban	PS		ORAL
			Atorvastatin Calcium	C		
			Aspirin	C		
			Nitroglycerin	C		

Date:03/01/99ISR Number: 3218670-2Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0075018

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL	Convulsion	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:03/01/99ISR Number: 3426816-6Report Type:Periodic  
Age:69 YR Gender:Female I/FU:I

Company Report #A0077030

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose 150 MG /	Flushing Insomnia	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Night Sweats

TWICE PER DAY

ORAL

Ibuprofen C  
Zolpidem Tartrate C

Date:03/01/99ISR Number: 3426822-1Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #A0077031

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Erythema Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE PER DAY ORAL		Urticaria					

Date:03/01/99ISR Number: 3426825-7Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #A0077032

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY/ORAL							

Carisoprodol C  
Ibuprofen C  
Ulcer Medication C  
Diuretic C



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3426828-2Report Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #A0077033

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Dermatitis Disorientation  Pruritus Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3426837-3Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0077034

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Insomnia Major Depression  Palpitations	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Birth Control	C		

Date:03/01/99ISR Number: 3426840-3Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #A0077066

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Arthralgia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3426845-2Report Type:Periodic  
Age:59 YR Gender:Female I/FU:I

Company Report #A0077067

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nausea	Consumer	Bupropion			

150 MG TWICE  
 PER DAY ORAL

Tremor

Hydrochloride PS Glaxo Wellcome Inc ORAL

Combivent C  
 Salbutamol Sulphate C  
 Loratadine C  
 Triamcinolone  
 Acetonide C  
 Guaphenesin C  
 Paracetamol C  
 Naproxen Sodium C

Date:03/01/99ISR Number: 3426849-XReport Type:Periodic Company Report #A0077068  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150MG PER DAY							
ORAL							

Date:03/01/99ISR Number: 3426854-3Report Type:Periodic Company Report #A0077071  
 Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia Insomnia	Consumer	Zyban Tablet- Zyban	PS	Pharmacia And Upjohn Co	ORAL
150 MG TWICE							
PER DAY ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

150 MG/TWICE  
 PER DAY /ORAL  
 Nicotine C Glaxo Wellcome Inc ORAL

Date:03/01/99ISR Number: 3426857-9Report Type:Periodic Company Report #A0077072  
 Age:55 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Agitation Dermatitis	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Nervousness					
ORAL		Tremor					

Mellaril C  
 Theophylline C  
 Prednisone C  
 Oxygen C

Date:03/01/99ISR Number: 3426858-0Report Type:Periodic Company Report #A0077073  
 Age:61 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE		Skin Odour Abnormal Sweat Gland Disorder	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ORAL		Urine Odour Abnormal					

Theophylline C

Date:03/01/99ISR Number: 3426860-9Report Type:Periodic Company Report #A0077075  
 Age:47 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Decreased Appetite	Consumer	Bupropion			

150 MG TWICE	Insomnia		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL						
Date:03/01/99ISR Number: 3426864-6Report Type:Periodic Company Report #A0077076						
Age:	Gender:Male	I/FU:I				
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Abdominal Pain	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc
150 MG PER						ORAL
DAY ORAL						

Date:03/01/99ISR Number: 3426866-XReport Type:Periodic Company Report #A0077077						
Age:48 YR	Gender:Female	I/FU:I				
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Crying Major Depression	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc
150 MG TWICE						ORAL
PER DAY ORAL				Paroxetine Hydrochloride	C	
				Alprazolam	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3426869-5Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0077078

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hunger Mood Swings	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3426874-9Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0077079

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia Mouth Ulceration	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAYORAL							

Date:03/01/99ISR Number: 3426876-2Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0077081

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
				Benazepril Hydrochloride	C		
				Thyroxine Sodium	C		
				Atorvastatin Calcium	C		

Date:03/01/99ISR Number: 3426882-8Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0077082

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Pruritic	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL				Echinacea	C		

Date:03/01/99ISR Number: 3426888-9Report Type:Periodic Company Report #A0077083  
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Skin Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL		Urticaria		Hrt	C		

Date:03/01/99ISR Number: 3426892-0Report Type:Periodic Company Report #A0077084  
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							



150 MG TWICE  
PER DAY ORAL

Date:03/01/99ISR Number: 3426922-6Report Type:Periodic Company Report #A0076189  
Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Urticaria					

DOSE ORAL

Date:03/01/99ISR Number: 3426924-XReport Type:Periodic Company Report #A0076191  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypersensitivity	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Pruritus	Professional				
		Swelling		Diclofenac Sodium	C		
		Urticaria		Naproxen Sodium	C		

Date:03/01/99ISR Number: 3426926-3Report Type:Periodic Company Report #A0076192  
Age:33 YR Gender:Female I/FU:I

Outcome PT  
Insomnia  
Pruritus



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Rash Papular

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL			Ibuprofen	C		

Date:03/01/99ISR Number: 3426929-9Report Type:Periodic Company Report #A0076193  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Tachycardia					
PER DAY ORAL							

Date:03/01/99ISR Number: 3426931-7Report Type:Periodic Company Report #A0076194  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Insomnia					
PER DAY ORAL		Pharyngolaryngeal Pain		Triphasil	C		

Date:03/01/99ISR Number: 3426934-2Report Type:Periodic Company Report #A0076195  
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Insomnia					
DAY ORAL		Musculoskeletal Stiffness		Cefaclor	C		

Date:03/01/99ISR Number: 3426936-6Report Type:Periodic Company Report #A0076197  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Swelling					
PER DAY ORAL		Urticaria					

Date:03/01/99ISR Number: 3426940-8Report Type:Periodic Company Report #A0076199  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG SINGLE		Feeling Abnormal					
DOSE ORAL		Flatulence		Thyroxine Sodium	C		
		Headache					
		Mental Impairment					
		Nausea					

Date:03/01/99ISR Number: 3426943-3Report Type:Periodic Company Report #A0076200  
Age:60 YR Gender:Male I/FU:I

Outcome	PT
	Constipation
	Flatulence

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Weight Increased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL			Vitamin E Aspirin	C C		

Date:03/01/99ISR Number: 3426946-9Report Type:Periodic Company Report #A0076201  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL				Omeprazole	C		

Date:03/01/99ISR Number: 3426951-2Report Type:Periodic Company Report #A0076202  
Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Hyperhidrosis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL							

Date:03/01/99ISR Number: 3426955-XReport Type:Periodic Company Report #A0076203  
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Diarrhoea	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL		Dizziness	Professional				

Feeling Hot  
Flatulence  
Haemorrhoids  
Ill-Defined Disorder  
Nausea

Verapamil C  
Lisinopril C  
Atorvastatin Calcium C

Date:03/01/99ISR Number: 3426958-5Report Type:Periodic Company Report #A0076204  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Dermatitis					
PER DAY ORAL		Pruritus		Medroxyprogesterone Ace.	C		

Date:03/01/99ISR Number: 3426962-7Report Type:Periodic Company Report #A0076205  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Dry Mouth					
PER DAY ORAL		Ear Discomfort		Nifedipine	C		
		Migraine					
		Pharyngolaryngeal Pain					
		Rhinorrhoea					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3426966-4Report Type:Periodic Company Report #A0076206  
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Insomnia					
PER DAY ORAL		Tremor					

Date:03/01/99ISR Number: 3426967-6Report Type:Periodic Company Report #A0076207  
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Pruritus					
PER DAY ORAL				Thyroid Medication	C		
				Magnesium Salt	C		
				Multivitamin	C		

Date:03/01/99ISR Number: 3427023-3Report Type:Periodic Company Report #A0076208  
 Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3427025-7Report Type:Periodic Company Report #A0076209  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							

Pruritus  
PER DAY ORAL  
Urticaria

Date:03/01/99ISR Number: 3427026-9Report Type:Periodic Company Report #A0076210  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyslexia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							

PER DAY ORAL

Date:03/01/99ISR Number: 3427030-0Report Type:Periodic Company Report #A0076383  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Difficulty In Walking	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
ORAL							
		Dysgraphia					
		Feeling Drunk		Prempro	C		
		Speech Disorder					
		Vertigo					

Date:03/01/99ISR Number: 3427036-1Report Type:Periodic Company Report #A0076409  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gingival Bleeding	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							

PER DAY ORAL

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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427037-3Report Type:Periodic Company Report #A0076410  
 Age:40 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Gingival Bleeding	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427041-5Report Type:Periodic Company Report #A0076412  
 Age:35 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Rash Erythematous					
		Urticaria		Antibiotics	C		
				Miconazole Nitrate	C		
				Tylenol With Codeine	C		

Date:03/01/99ISR Number: 3427043-9Report Type:Periodic Company Report #A0076413  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Heart Rate Increased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Hyperhidrosis					
				Aspirin	C		
				Vitamin	C		

Date:03/01/99ISR Number: 3427046-4Report Type:Periodic Company Report #A0076414  
 Age:50 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE			DERMATITIS	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL			PRURITUS					
			TINNITUS		Simvastatin	C		
Date:03/01/99ISR Number: 3427051-8Report Type:Periodic				Company Report #A0076415				
Age:43 YR Gender:Male I/FU:I								
150 MG TWICE			Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL			Insomnia					
Date:03/01/99ISR Number: 3427055-5Report Type:Periodic				Company Report #A0076416				
Age:66 YR Gender:Female I/FU:I								
150 MG TWICE			Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL			Vomiting					
					Salbutamol Sulphate	C		
Date:03/01/99ISR Number: 3427062-2Report Type:Periodic				Company Report #A0076417				
Age:20 YR Gender:Female I/FU:I								
150 MG TWICE			Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL			Pharyngolaryngeal Pain					
					Birth Control	C		

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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427064-6Report Type:Periodic Company Report #A0076421  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3427068-3Report Type:Periodic Company Report #A0076521  
 Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
		Nausea		Birth Control	C		

Date:03/01/99ISR Number: 3427071-3Report Type:Periodic Company Report #A0076522  
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3427082-8Report Type:Periodic Company Report #A0076524  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
		Hyperhidrosis					
		Insomnia					

Memory Impairment  
Nervousness  
Tachycardia  
Tremor

Date:03/01/99ISR Number: 3427086-5Report Type:Periodic Company Report #A0076525  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Epistaxis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Headache					
DAY ORAL							

Date:03/01/99ISR Number: 3427088-9Report Type:Periodic Company Report #A0076526  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Face Oedema	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Hypoaesthesia Oral					
PER DAY ORAL		Pruritus					
		Urticaria					

Date:03/01/99ISR Number: 3427091-9Report Type:Periodic Company Report #A0076527  
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Libido Increased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427093-2Report Type:Periodic  
Age:68 YR Gender:Male I/FU:I

Company Report #A0075291

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Dyspraxia Movement Disorder	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Company Representative	Probenecid Clobetasol Priopionate Atenolol Atorvastatin Calcium Insulin Thyroxine Sodium Lisinopril Trazodone Hydrochloride Diazepam Aspirin	C C C C C C C C C		

Date:03/01/99ISR Number: 3427096-8Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #A0075292

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Oral Contraceptive	C		

Date:03/01/99ISR Number: 3427098-1Report Type:Periodic  
Age:54 YR Gender:Male I/FU:I

Company Report #A0075293

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Dermatitis Pain Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Clidinium Bromide	C		

Swelling  
Urticaria

Lansoprazole

C

Date:03/01/99ISR Number: 3427100-7Report Type:Periodic Company Report #A0075294  
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICED		Mouth Ulceration					
PER DAY ORAL		Pain		Diuretic	C		
				Multivitamin	C		

Date:03/01/99ISR Number: 3427103-2Report Type:Periodic Company Report #A0075295  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decreased Appetite	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Nausea					
DAY ORAL				Azithromycin	C		



150 MG TWICE	Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL	Dizziness					
	Dry Mouth		Hydrochlorothiazide	C		
	Gingival Bleeding		Microdiol	C		
	Increased Appetite					
	Insomnia					
	Toothache					

Date:03/01/99ISR Number: 3427123-8Report Type:Periodic Company Report #A0775301  
 Age:69 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Asthenia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL		Chest Pain	Professional				
		Dizziness		Atorvastatin Calcium	C		
		Insomnia					
		Palpitations					

Date:03/01/99ISR Number: 3427126-3Report Type:Periodic Company Report #A0075303  
 Age:49 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL		Swelling					
				Zestoretic	C		
				Atorvastatin Calcium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Ethanol C

Date:03/01/99ISR Number: 3427129-9Report Type:Periodic Company Report #A0075304  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER							
DAY ORAL							

Lisinopril C

Date:03/01/99ISR Number: 3427132-9Report Type:Periodic Company Report #A0075305  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Erythematous	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Urticaria					
PER DAY ORAL							

Date:03/01/99ISR Number: 3427133-0Report Type:Periodic Company Report #A0075306  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Face Oedema	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Pruritus					
PER DAY ORAL							

Date:03/01/99ISR Number: 3427136-6Report Type:Periodic Company Report #A0075308  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG TWICE	Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL	Fatigue					
	Irritability		Multivitamin	C		
	Mood Altered					

Date:03/01/99ISR Number: 3427138-XReport Type:Periodic Company Report #A0075310  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL		Disturbance In Attention					
		Hyperhidrosis		Buspirone Hydrochloride	C		

Date:03/01/99ISR Number: 3427142-1Report Type:Periodic Company Report #A0075312  
 Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Confusional State	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL		Nausea					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427146-9Report Type:Periodic  
Age:60 YR Gender:Male I/FU:I

Company Report #A0075313

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL		Dermatitis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Pruritus	Professional	Comtrex	C		
		Purpura		Ibuprofen	C		
		Rash Macular					
		Rash Papular					
		Urticaria					

Date:03/01/99ISR Number: 3427151-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0075341

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Creatinine Renal Clearance Decreased	Consumer	Zyban	PS	Glaxo Wellcome Inc	
		Drug Interaction		Thyroxine Sodium Tablet	SS		

Date:03/01/99ISR Number: 3427154-8Report Type:Periodic  
Age:23 YR Gender:Male I/FU:I

Company Report #A0075375

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Dizziness		Zyban	PS	Glaxo Wellcome Inc	ORAL
		Dry Mouth					
		Dysgeusia					
		Epistaxis					
		Hallucination					
		Nausea					
		Palpitations					
		Pollakiuria					
		Tremor					

Date:03/01/99ISR Number: 3427167-6Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #A0075038

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL		Dermatitis Nasal Congestion	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427171-8Report Type:Periodic Company Report #A0075039  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
10 DAY		Dry Mouth Irritability	Consumer	Zyban	PS	Glaxo Wellcome Inc	

Date:03/01/99ISR Number: 3427172-XReport Type:Periodic Company Report #A0075041  
 Age:49 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Tension	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Blood Pressure Medication Ibuprofen	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427175-5Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0075047

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3427178-0Report Type:Periodic  
Age:69 YR Gender:Female I/FU:I

Company Report #A0075050

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Anxiety	Professional				
PER DAY/ORAL		Dizziness		Paroxetine			
		Dry Mouth		Hydrochloride	C		
		Dry Throat		Trazodone	C		
		Headache		Alprazolam	C		
		Insomnia		Herbal Medication	C		
		Nausea		Multivitamin	C		
		Nervousness		Fluticasone			
		Tremor		Propionate	C		
				Salbutamol Sulphate	C		

Date:03/01/99ISR Number: 3427180-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0075051

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL							

Date:03/01/99ISR Number: 3427182-2Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #A0075052

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL				Oral Contraceptive	C		

Date:03/01/99ISR Number: 3427184-6Report Type:Periodic Company Report #A0075053  
 Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nervousness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Tremor					
DAY/ ORAL		Vomiting		Tramadol Hydrochloride	C		

Date:03/01/99ISR Number: 3427186-XReport Type:Periodic Company Report #A0075055  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Gastrooesophageal Reflux					
PER DAY/ ORAL		Disease					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427188-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0075074

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dysgeusia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/	ORAL						

Date:03/01/99ISR Number: 3427190-1Report Type:Periodic  
 Age:39 YR Gender:Female I/FU:I

Company Report #A0075075

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Condition Aggravated					
PER DAY/	ORAL						
		Confusional State		Omeprazole	C		
		Dermatitis		Oral Contraceptive	C		
		Diarrhoea					
		Electric Shock					
		Feeling Abnormal					
		Hyperhidrosis					
		Smoker					
		Swelling					
		Tachycardia					
		Tremor					

Date:03/01/99ISR Number: 3427192-5Report Type:Periodic  
 Age:41 YR Gender:Female I/FU:I

Company Report #A0075076

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/	TWICE						
PER DAY/	ORAL						
				No Concurrent Medication	C		

Date:03/01/99ISR Number: 3427194-9Report Type:Periodic Company Report #A0075077  
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Hyperhidrosis	Professional				
PER DAY/ ORAL		Nausea		Cimetidine	C		
		Vomiting		Donnatal	C		

Date:03/01/99ISR Number: 3427217-7Report Type:Periodic Company Report #A0075109  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL							

Date:03/01/99ISR Number: 3427219-0Report Type:Periodic Company Report #A0075189  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL				Semisodium Valproate	C		
				Paroxetine			
				Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427221-9Report Type:Periodic Company Report #A0075190  
 Age:39 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY / ORAL		Agitation Dissociation Disturbance In Attention Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427224-4Report Type:Periodic Company Report #A0075191  
 Age:60 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Abnormal Faeces Diarrhoea Urine Abnormality	Consumer	Zyban ....	PS C	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427226-8Report Type:Periodic Company Report #A0075200  
 Age:23 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Ear Disorder Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427229-3Report Type:Periodic Company Report #A0075289  
 Age:28 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /TWICE		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

PER DAY/ ORAL

Insomnia

Sleep Talking Birth Control C

Sleep Walking

Tachycardia

Tremor

Date:03/01/99ISR Number: 3427230-XReport Type:Periodic Company Report #A0075290

Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

150 MG/ TWICE

PER DAY/ ORAL

Conjugated Estrogens C

Ascorbic Acid C

Date:03/01/99ISR Number: 3427257-8Report Type:Periodic Company Report #A0076528

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Pain Upper	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL

150 MG ORAL

Professional  
Company  
Representative



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427259-1Report Type:Periodic Company Report #A0076529  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diplopia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL			Professional Company Representative				

Date:03/01/99ISR Number: 3427261-XReport Type:Periodic Company Report #A0076557  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Pruritus	Professional				
PER DAY ORAL		Swelling		Thyroxine Sodium	C		

Date:03/01/99ISR Number: 3427262-1Report Type:Periodic Company Report #A0076558  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER				Multivitamin	C		
DAY ORAL							

Date:03/01/99ISR Number: 3427264-5Report Type:Periodic Company Report #A0076559  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							

MONTHS

MON

Unknown Patch

SS

Date:03/01/99ISR Number: 3427266-9Report Type:Periodic  
Age:82 YR Gender:Female I/FU:I

Company Report #A0076560

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Agitation					
PER DAY ORAL		Arthropathy					
		Fall		Amitriptyline Hcl	C		
				Famotidine	C		
				Fluticasone			
				Propionate	C		

Date:03/01/99ISR Number: 3427269-4Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0076561

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Headache					
PER DAY ORAL		Insomnia					
				Ortho-Novum	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427271-2Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0076562

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Ibuprofen C

Date:03/01/99ISR Number: 3427274-8Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0076563

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Pruritus	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL

Medroxyprogesterone  
Ace. C

Date:03/01/99ISR Number: 3427275-XReport Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0076564

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL	4 WK	Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Multivitamin C

Date:03/01/99ISR Number: 3427278-5Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0076969

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Hair Texture Abnormal	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

PER DAY ORAL

Multivitamin C  
Mesalazine C

Date:03/01/99ISR Number: 3427281-5Report Type:Periodic Company Report #A0076970  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Bronchitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Pneumonia					
PER DAY ORAL		Upper Respiratory Tract Infection		Thyroxine Sodium Loestrin Omeprazole	C C C		

Date:03/01/99ISR Number: 3427282-7Report Type:Periodic Company Report #A0076971  
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain Upper	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Anxiety					
PER DAY ORAL		Insomnia Malaise Nervousness Tremor		Conjugated Estrogens Cimetidine Oxybutynin Hydrochloride Multivitamin	C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427284-0Report Type:Periodic  
 Age:42 YR Gender:Female I/FU:I

Company Report #A0076972

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
				Aspirin	C		
				Clopidogrel			
				Bisulphate	C		
				Omeprazole	C		

Date:03/01/99ISR Number: 3427286-4Report Type:Periodic  
 Age:45 YR Gender:Female I/FU:I

Company Report #A0076973

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
		Dysgeusia					
		Headache		Famotidine	C		
		Insomnia		Cetirizine			
		Tremor		Hydrochloride	C		
				Alendronate Sodium	C		
				Atorvastatin Calcium	C		
				Triamcinolone			
				Acetonide	C		
				Naproxen	C		
				Salbutamol Sulphate	C		
				Ipratropium Bromide	C		

Date:03/01/99ISR Number: 3427288-8Report Type:Periodic  
 Age:65 YR Gender:Female I/FU:I

Company Report #A0076974

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fatigue	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
		Sensation Of Pressure					

Date:03/01/99ISR Number: 3427289-XReport Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #A0076975

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG	TWICE		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY	ORAL	Tremor					
				Ziac	C		
				Alprazolam	C		
				Bumetanide	C		
				Antibiotics	C		

Date:03/01/99ISR Number: 3427290-6Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #A0076976

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG	TWICE		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY	ORAL	Tremor					
				Estropipate	C		
				Progesterone	C		
				Aspirin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427291-8Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0076977

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Nervousness					
PER DAY ORAL							

Tamoxifen C

Date:03/01/99ISR Number: 3427292-XReport Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0076978

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Diarrhoea					
PER DAY ORAL		Dizziness		Ranitidine			
		Nausea		Hydrochloride	C		
		Tremor		Conjugated Estrogens	C		

Date:03/01/99ISR Number: 3427294-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0075390

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ UKN /							
ORAL							

Date:03/01/99ISR Number: 3427295-5Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #A0075419

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	
UNK	24 DAY						

Emotional Disorder  
Headache  
Irritability  
Lethargy  
Major Depression  
Weight Increased

Date:03/01/99ISR Number: 3427297-9Report Type:Periodic Company Report #A0075524  
Age:49 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dermatitis Fatigue Malaise Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427298-0Report Type:Periodic Company Report #A0075525  
Age:45 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Sumatriptan Succinate	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427300-6Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0075526

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Hypertension					
TWICE PER DAY		Lethargy					
/ ORAL		Syncope					

Date:03/01/99ISR Number: 3427301-8Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #A0075527

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	
150 MG /		Headache					
TWICE PER DAY		Tremor					
/ ORAL							

Date:03/01/99ISR Number: 3427303-1Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0075528

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Thyroxine Sodium	C		

Date:03/01/99ISR Number: 3427308-0Report Type:Periodic  
Age:52 YR Gender:Male I/FU:I

Company Report #A0075529

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Headache	Professional				
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3427310-9Report Type:Periodic Company Report #A0075530  
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Unevaluable Event	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3427311-0Report Type:Periodic Company Report #A0075531  
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decreased Appetite	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Dry Mouth					
TWICE PER DAY		Dysgeusia					
/ ORAL		Headache					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427357-2Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #A0076565

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Effect Decreased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
				Glucophage	C		
				Lisinopril	C		
				Warfarin Sodium	C		
				Glipizide	C		

Date:03/01/99ISR Number: 3427361-4Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0076566

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anorexia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER							
DAY ORAL		Insomnia					
		Major Depression					

Date:03/01/99ISR Number: 3427363-8Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0076567

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL		Eye Disorder					
		Eye Irritation		Nyquil	C		
		Hypersensitivity		Tylenol Sinus			
		Pruritus		Medication	C		
		Swelling					

Date:03/01/99ISR Number: 3427373-0Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0076568

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Insomnia Nervousness Palpitations	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427374-2Report Type:Periodic Company Report #A0076569  
 Age:36 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Dermatitis Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427376-6Report Type:Periodic Company Report #A0076570  
 Age:44 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Dissociation Insomnia Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427377-8Report Type:Periodic  
Age:69 YR Gender:Male I/FU:I

Company Report #A0076571

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Agitation					
PER DAY ORAL		Dizziness					
		Insomnia		Triamterene	C		
		Panic Attack		Digoxin	C		
		Syncope		Oxpentifylline	C		
				Lisinopril	C		
				Vitamin	C		
				Aspirin	C		
				Defibrillation	C		

Date:03/01/99ISR Number: 3427380-8Report Type:Periodic  
Age:59 YR Gender:Male I/FU:I

Company Report #A0076572

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	
		Insomnia		Atorvastatin Calcium	C		
				Aspirin	C		
				Ginkgo Biloba	C		

Date:03/01/99ISR Number: 3427381-XReport Type:Periodic  
Age:64 YR Gender:Male I/FU:I

Company Report #A0076573

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Erectile Dysfunction					
PER DAY ORAL				Buspirone			
				Hydrochloride	C		
				Hydroxyzine	C		

Date:03/01/99ISR Number: 3427383-3Report Type:Periodic Company Report #A0076574  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Confusional State					
PER DAY ORAL		Nausea		Ibuprofen	C		
		Stomach Discomfort		Beclomethasone Dipropion.	C		

Date:03/01/99ISR Number: 3427384-5Report Type:Periodic Company Report #A0076575  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Dissociation					
PER DAY ORAL		Hyperhidrosis					
		Nausea					
		Paranoia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427386-9Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #A0076610

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Anorexia Arrhythmia Dysgeusia Dyspnoea Headache Tachycardia Tremor	Consumer	Zyban  Amitriptyline Hcl	PS  C	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427388-2Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #A0076617

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Agitation Arrhythmia Constipation Dry Mouth Dysgeusia Dyspnoea Headache Hyperhidrosis Tremor Vision Blurred	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427390-0Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #A0076656

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Chest Pain Headache Nausea	Consumer	Zyban  Multivitamin	PS  C	Glaxo Wellcome Inc	ORAL

Vomiting

Date:03/01/99ISR Number: 3427392-4Report Type:Periodic Company Report #A0076716  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Joint Stiffness	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL		Joint Swelling	Professional Company Representative				

Date:03/01/99ISR Number: 3427393-6Report Type:Periodic Company Report #A0076724  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accommodation Disorder	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Attention					
PER DAY ORAL		Deficit/Hyperactivity Disorder Depression Disturbance In Attention					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427394-8Report Type:Periodic Company Report #A0076735  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Nausea					
PER DAY ORAL		Nervousness					
		Tremor					

Date:03/01/99ISR Number: 3427396-1Report Type:Periodic Company Report #A0076739  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Company	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL		Nightmare	Representative				

Date:03/01/99ISR Number: 3427398-5Report Type:Periodic Company Report #A0076851  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Pruritus					
PER DAY ORAL		Urticaria		Entex La	C		
				Beclomethasone	C		
				Dipropion.			

Date:03/01/99ISR Number: 3427399-7Report Type:Periodic Company Report #A0076855  
 Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							

PER DAY ORAL

Enalapril Maleate C

Date:03/01/99ISR Number: 3427400-0Report Type:Periodic Company Report #A0076884  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Major Depression					

PER DAY ORAL

Date:03/01/99ISR Number: 3427401-2Report Type:Periodic Company Report #A0076888  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Dyspepsia					
PER DAY ORAL		Nausea		Actifed	C		

Date:03/01/99ISR Number: 3427402-4Report Type:Periodic Company Report #A0076889  
Age:40 YR Gender:Female I/FU:I

Outcome	PT
	Abdominal Pain Upper
	Face Oedema
	Fatigue

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Freedom Of Information (FOI) Report

Oedema Peripheral

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL			Thyroxine Sodium	C		

Date:03/01/99ISR Number: 3427403-6Report Type:Periodic Company Report #A0076890  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL				Thyroxine Sodium	C		

Date:03/01/99ISR Number: 3427404-8Report Type:Periodic Company Report #A0076891  
 Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Condition Aggravated	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL		Insomnia		Lovastatin	C		
		Tremor					

Date:03/01/99ISR Number: 3427405-XReport Type:Periodic Company Report #A0076892  
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Aggression	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
OER DAY ORAL		Personality Change	Professional				

Diltiazem C  
Glucophage C  
Glibenclamide C

Date:03/01/99ISR Number: 3427406-1Report Type:Periodic Company Report #A0076893  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nervousness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Tension					
PER DAY ORAL							

Date:03/01/99ISR Number: 3427407-3Report Type:Periodic Company Report #A0076894  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Hypersensitivity					
PER DAY ORAL		Urticaria					

Date:03/01/99ISR Number: 3427408-5Report Type:Periodic Company Report #A0076895  
Age:43 YR Gender:Male I/FU:I

Outcome	PT
	Aggression
	Depression
	Emotional Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Paranoia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Health Professional	Zyban  Ethanol (Formulation Unknown) Cyclobenzaprine Hcl	PS  SS C	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427409-7Report Type:Periodic Company Report #A0076896  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 150 MG ORAL		Decreased Appetite	Other	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427410-3Report Type:Periodic Company Report #A0076897  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 150 MG ORAL		Drug Ineffective	Other	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427411-5Report Type:Periodic Company Report #A0076898  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 150 MG ORAL	2 WK	Erythema Hypersensitivity Pruritus	Other	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427412-7Report Type:Periodic Company Report #A0076899  
Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Effect Decreased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Insomnia					
		Nervousness					

Date:03/01/99ISR Number: 3427413-9Report Type:Periodic Company Report #A0076900  
Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Disturbance In Attention	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427414-0Report Type:Periodic Company Report #A0076901  
Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Pollakiuria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Sexual Dysfunction					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427415-2Report Type:Periodic Company Report #A0076902  
 Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Papular	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
				Vicodin	C		
				Diazepam	C		

Date:03/01/99ISR Number: 3427416-4Report Type:Periodic Company Report #A0076903  
 Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chills	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							
		Erythema Multiforme	Professional				
		Joint Stiffness					
		Joint Swelling					
		Oedema Peripheral					
		Urticaria					

Date:03/01/99ISR Number: 3427417-6Report Type:Periodic Company Report #A0076904  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
		Cold Sweat					
		Diarrhoea					
		Malaise					
		Tachycardia					
		Vomiting					

Date:03/01/99ISR Number: 3427418-8Report Type:Periodic Company Report #A0076905  
 Age:25 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Blood Pressure Systolic	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL		Increased	Professional				
		Headache					
		Heart Rate Increased					

Date:03/01/99ISR Number: 3427419-XReport Type:Periodic Company Report #A0076968  
 Age:34 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Oedema Peripheral	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL		Pruritus					
		Rash Generalised					
		Urticaria					

Date:03/01/99ISR Number: 3427421-8Report Type:Periodic Company Report #A0078538  
 Age:36 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL		Anxiety					
		Dermatitis		Ortho-Novum	C		
		Hyperhidrosis		Ascorbic Acid	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427422-XReport Type:Periodic Company Report #A0078539  
 Age:46 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL				Estrogen	C		

Date:03/01/99ISR Number: 3427423-1Report Type:Periodic Company Report #A0078566  
 Age:48 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							

Date:03/01/99ISR Number: 3427424-3Report Type:Periodic Company Report #A0078818  
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Depression					
PER DAY ORAL							

Date:03/01/99ISR Number: 3427425-5Report Type:Periodic Company Report #A0078819  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL				Hyzaar	C		
				Prempro	C		
				Etodolac	C		

Date:03/01/99ISR Number: 3427426-7Report Type:Periodic Company Report #A0059124  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	

Date:03/01/99ISR Number: 3427427-9Report Type:Periodic Company Report #A0060052  
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Pruritus					
PER DAY ORAL							

Conjugated Estrogens	C
Centrum	C
Medroxyprogesterone	
Ace.	C

Date:03/01/99ISR Number: 3427428-0Report Type:Periodic Company Report #A0064579  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Creatinine	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Increased	Professional				
DAY ORAL		Malaise		Ibuprofen (Formulation			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Unknown) C

Date:03/01/99ISR Number: 3427429-2Report Type:Periodic Company Report #A0066450  
 Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Ear Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Headache					
PER DAY ORAL		Musculoskeletal Stiffness		Bupropion			
		Paraesthesia		Hydrochloride	C		
				Vitamin E	C		
				Multivitamin	C		

Date:03/01/99ISR Number: 3427430-9Report Type:Periodic Company Report #A0070235  
 Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Nervous System Disorder	Professional				
PER DAY ORAL		Oral Intake Reduced					

Date:03/01/99ISR Number: 3427432-2Report Type:Periodic Company Report #A0070280  
 Age:75 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tremor		Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL				Digoxin	C		
				Atorvastatin Calcium	C		
				Warfarin Sodium	C		
				Losartan Potassium	C		

Date:03/01/99ISR Number: 3427434-6Report Type:Periodic Company Report #A0070823  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Headache	Professional				
DAY ORAL		Insomnia					

Date:03/01/99ISR Number: 3427436-XReport Type:Periodic Company Report #A0071074  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blister	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL		Dizziness	Professional				
		Hypertension					
		Joint Stiffness					
		Pain In Extremity					
		Skin Exfoliation					
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427437-1Report Type:Periodic  
Age:47 YR Gender:Female I/FU:F

Company Report #A0071181

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Agitation	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Constipation	Professional				
		Disorientation		Nifedipine	C		
		Dry Mouth					
		Hyperhidrosis					
		Insomnia					
		Nausea					

Date:03/01/99ISR Number: 3427438-3Report Type:Periodic  
Age:68 YR Gender:Female I/FU:F

Company Report #A0071715

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Arrhythmia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Balance Disorder	Professional				
		Blood Pressure Increased		Conjugated Estrogens	C		
		Circulatory Collapse					
		Dizziness					
		Malaise					
		Vision Blurred					

Date:03/01/99ISR Number: 3427440-1Report Type:Periodic  
Age:56 YR Gender:Male I/FU:F

Company Report #A0072154

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Disorientation	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Dizziness	Professional				
		Insomnia		Carvedilol	C		
				Simvastatin	C		
				Digoxin	C		
				Aspirin	C		

Date:03/01/99ISR Number: 3427442-5Report Type:Periodic Company Report #A0072548  
Age:69 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dry Mouth	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWIC		Insomnia	Professional				
PER DAY ORAL		Pollakiuria		Simvastatin Amlodipine	C C		

Date:03/01/99ISR Number: 3427444-9Report Type:Periodic Company Report #A0073460  
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Cold Sweat	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Dizziness	Professional				
PER DAY ORAL		Fall Headache Vision Blurred Vomiting		Ranitidine Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427446-2Report Type:Periodic  
Age:36 YR Gender:Female I/FU:F

Company Report #A0073583

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Erythema	Professional				
PER DAY ORAL		Pruritus		Birth Control	C		

Date:03/01/99ISR Number: 3427448-6Report Type:Periodic  
Age:34 YR Gender:Male I/FU:F

Company Report #A0073595

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Insomnia					
PER DAY ORAL		Pruritus					

Date:03/01/99ISR Number: 3427450-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0078435

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Other	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							

Date:03/01/99ISR Number: 3427452-8Report Type:Periodic  
Age:69 YR Gender:Male I/FU:I

Company Report #A0078436

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Nausea					
PER DAY ORAL				Metoprolol Tartrate	C		
				Diltiazem			

Hydrochloride C  
Aspirin C

Date:03/01/99ISR Number: 3427455-3Report Type:Periodic Company Report #A0078437  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperglycaemia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
				Amitriptyline Hcl	C		
				Glucophage	C		
				Lisinopril	C		

Date:03/01/99ISR Number: 3427456-5Report Type:Periodic Company Report #A0078438  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Decreased Appetite					
PER DAY ORAL		Dyspepsia					
				Lotrel	C		
				Oxpentifylline	C		



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Date:03/01/99ISR Number: 3427457-7Report Type:Periodic Company Report #A0078439  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Headache					
PER DAY ORAL		Nervousness					

Date:03/01/99ISR Number: 3427459-0Report Type:Periodic Company Report #A0078440  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE				Nicotine	C		
PER DAY ORAL				Prinzide	C		

Date:03/01/99ISR Number: 3427461-9Report Type:Periodic Company Report #A0078441  
 Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gingival Recession	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Tooth Abscess					
PER DAY ORAL				Multivitamins	C		

Date:03/01/99ISR Number: 3427463-2Report Type:Periodic Company Report #A0078442  
 Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							

PER DAY ORAL

Lansoprazole C  
Losartan Potassium C  
Prednisone C

Date:03/01/99ISR Number: 3427465-6Report Type:Periodic Company Report #A0078443  
Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Capillary Leak Syndrome	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL		Hypersensitivity Swelling	Professional Company Representative				

Date:03/01/99ISR Number: 3427467-XReport Type:Periodic Company Report #A0078450  
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Major Depression	Professional				
PER DAY ORAL			Company Representative	Conjugated Estrogens	C		
				Simvastatin	C		
				Ranitidine			
				Hydrochloride	C		
				Amlodipine	C		

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Date:03/01/99ISR Number: 3427469-3Report Type:Periodic  
Age:65 YR Gender:Male I/FU:I

Company Report #A0078527

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Confusional State Dissociation Nervousness	Consumer	Zyban  Warfarin Sodium Ibuprofen	PS  C C	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427472-3Report Type:Periodic  
Age:58 YR Gender:Male I/FU:I

Company Report #A0078528

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Dry Mouth Dysgeusia Insomnia	Consumer	Zyban  Diltiazem Hydrochloride Simvastatin Alprazolam Paracetamol	PS  C C C C	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427474-7Report Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #A0078529

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Orgasm Abnormal Sexual Dysfunction	Consumer	Zyban  Nortriptyline Hcl	PS  C	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427476-0Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #A0078530

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Lethargy					
PER DAY ORAL				Simvastatin	C		
				Omeprazole	C		

Date:03/01/99ISR Number: 3427477-2Report Type:Periodic Company Report #A0078532  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperaesthesia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Pain					
PER DAY ORAL							

Date:03/01/99ISR Number: 3427479-6Report Type:Periodic Company Report #A0078533  
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Skin	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Erythema					
DAY ORAL		Skin Disorder					



TWICE PER DAY	Chest Discomfort	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL	Chest Pain	Professional				
	Excitability					
	Headache					
	Nausea					

Date:03/01/99ISR Number: 3427488-7Report Type:Periodic Company Report #A0076979  
 Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Disturbance In Attention	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Dry Mouth					
DAY/ ORAL		Headache		Ibuprofen	C		
		Insomnia					

Date:03/01/99ISR Number: 3427489-9Report Type:Periodic Company Report #A0076980  
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Flatulence	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Nausea	Professional				
DAY/ ORAL				Antacid	C		
				Librax	C		



150 MG/ ORAL 17 DAY Drug Ineffective Consumer Zyban PS Glaxo Wellcome Inc ORAL  
Irritability  
Mood Altered

Date:03/01/99ISR Number: 3427504-2Report Type:Periodic Company Report #A0076985  
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hallucination	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Irritability					
DAY/ ORAL		Nervousness					

Date:03/01/99ISR Number: 3427506-6Report Type:Periodic Company Report #A0076986  
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Hyperhidrosis					
PER DAY/ ORAL		Nervousness		Nizatidine	C		
				Isosorbide			
				Mononitrate	C		
				Enalapril Maleate	C		
				Simvastatin	C		
				Aspirin	C		
				Frusemide	C		
				Potassium Chloride	C		



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Date:03/01/99ISR Number: 3427507-8Report Type:Periodic  
 Age:58 YR Gender:Male I/FU:I

Company Report #A0076987

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Anxiety Decreased Appetite Disorientation Faeces Discoloured Gout Personality Change	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3427509-1Report Type:Periodic  
 Age:48 YR Gender:Male I/FU:I

Company Report #A0076988

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Tremor Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Fexofenadine Hydrochlorid	C		

Date:03/01/99ISR Number: 3427511-XReport Type:Periodic  
 Age:46 YR Gender:Female I/FU:I

Company Report #A0077002

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Cefuroxime Axetil	C		

Date:03/01/99ISR Number: 3427512-1Report Type:Periodic  
 Age:37 YR Gender:Female I/FU:I

Company Report #A0077003

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL				Thyroxine Sodium	C		

Date:03/01/99ISR Number: 3427513-3Report Type:Periodic Company Report #A0077004  
 Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL		Constipation	Professional				
		Dry Mouth					
		Insomnia					
		Irritability					
		Muscle Twitching					
		Restlessness					

Date:03/01/99ISR Number: 3427514-5Report Type:Periodic Company Report #A0077005  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL		Insomnia					

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Date:03/01/99ISR Number: 3427515-7Report Type:Periodic Company Report #A0077006  
 Age:51 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Confusional State	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL		Disorientation					
		Insomnia		Nabumetone	C		

Date:03/01/99ISR Number: 3427516-9Report Type:Periodic Company Report #A0077025  
 Age:52 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Burning Sensation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAILY/ ORAL		Oedema Peripheral					
		Pruritus		Conjugated Estrogen	C		
		Urticaria		Amitriptyline Hcl	C		

Date:03/01/99ISR Number: 3427517-0Report Type:Periodic Company Report #A0077026  
 Age:55 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3427518-2Report Type:Periodic Company Report #A0077027  
 Age:41 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Bipolar I Disorder	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Suicidal Ideation

PER DAY/ ORAL

Nicotine C  
 Ketorolac No. 3 C  
 Zolpidem Tartrate C

Date:03/01/99ISR Number: 3427519-4Report Type:Periodic Company Report #A0077028  
 Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disturbance In Attention	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Erythema					
DAY/ ORAL		Vomiting		Nefazodone Hydrochloride	C		

Date:03/01/99ISR Number: 3427520-0Report Type:Periodic Company Report #A0077029  
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disturbance In Attention	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Dry Mouth					
PER DAY/ ORAL		Insomnia					

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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427865-4Report Type:Periodic  
Age:26 YR Gender:Male I/FU:F

Company Report #A0073729

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Agitation	Professional				
DAY / ORAL							

Date:03/01/99ISR Number: 3427867-8Report Type:Periodic  
Age:40 YR Gender:Female I/FU:F

Company Report #A0073753

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Tremor					
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3427870-8Report Type:Periodic  
Age:53 YR Gender:Female I/FU:F

Company Report #A0073775

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /			Professional				
TIWICE PER DAY							
/ ORAL							

Propranolol	C
Medroxyprogesterone	
Ace.	C
Dicyclomine	C
Hypericum	C
Ginkgo Biloba	C
Stool Softener	C
Polycarbophil	
Calcium	C
Multivitamin	C

Date:03/01/99ISR Number: 3427875-7Report Type:Periodic Company Report #A0073815  
Age:73 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dry Mouth	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Insomnia	Professional				
TWICE PER DAY							
/ ORAL							
				Oxpentifylline	C		
				Dipyridamole	C		
				Maxzide	C		
				Diltiazem			
				Hydrochloride	C		
				Nitroglycerin	C		
				Aspirin	C		
				Glibenclamide	C		

Date:03/01/99ISR Number: 3427879-4Report Type:Periodic Company Report #A0073822  
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dry Mouth	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Urticaria	Professional				
TWICE PER DAY							
/ ORAL							
				Lisinopril	C		

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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3427882-4Report Type:Periodic Company Report #A0073823  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tic	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /			Professional				
TWICE PER DAY			Company				
/ ORAL			Representative				

Date:03/01/99ISR Number: 3427886-1Report Type:Periodic Company Report #A0073834  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Joint Swelling	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL		Urticaria					

Date:03/01/99ISR Number: 3427890-3Report Type:Periodic Company Report #A0073837  
 Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE			Professional				
PER DAY /							
ORAL							

Date:03/01/99ISR Number: 3427893-9Report Type:Periodic Company Report #A0073850  
 Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							

DAY / ORAL  
 Dysgeusia Professional  
 Hypoaesthesia  
 Diltiazem  
 Hydrochloride C  
 Paroxetine  
 Hydrochloride C  
 Hydrochlorothiazide C  
 Conjugated Estrogens C  
 Vitamin C

Date:03/01/99ISR Number: 3427897-6Report Type:Periodic Company Report #A0074067  
 Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG / PER		Dry Mouth	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY / ORAL		Headache	Professional				
		Hyperhidrosis					
		Insomnia					
		Nausea					
		Sedation					
		Sinus Headache					

Date:03/01/99ISR Number: 3427903-9Report Type:Periodic Company Report #A0074207  
 Age:65 YR Gender:Female I/FU:F

Outcome PT  
 Difficulty In Walking  
 Hypersensitivity  
 Insomnia



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Dose	Duration	Nail Discolouration Pain Pruritus	Report Source	Product	Role	Manufacturer	Route
150 MG /		Rash Erythematous	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Swelling	Professional				
/ ORAL		Tachycardia		Conjugated Estrogens	C		

Date:03/01/99ISR Number: 3427907-6Report Type:Periodic Company Report #A0074222  
Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Psychomotor Hyperactivity	Professional				
DAY / ORAL							

Date:03/01/99ISR Number: 3427913-1Report Type:Periodic Company Report #A0074966  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nightmare	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 / TWICE			Professional				
PER DAY /			Company				
ORAL			Representative				

Date:03/01/99ISR Number: 3427916-7Report Type:Periodic Company Report #A0077085  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG	TWICE	Conjunctival Oedema	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Urticaria					
PER DAY	ORAL						

Date:03/01/99ISR Number: 3427918-0Report Type:Periodic Company Report #A0077086  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG	TWICE	Dermatitis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY	ORAL	Eyelid Oedema	Professional				
		Face Oedema		Nicotine	C		
		Hypersensitivity					
		Pruritus					
		Rash Erythematous					
		Rash Generalised					
		Urticaria					

Date:03/01/99ISR Number: 3427920-9Report Type:Periodic Company Report #A0077141 Age:39 YR Gender:Female I/FU:I
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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG		Arthralgia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Chest Pain					
ORAL				Herbal Medication	C		

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Date:03/01/99ISR Number: 3427923-4Report Type:Periodic Company Report #A0077145  
 Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG	TWICE						
PER DAY	ORAL						

Date:03/01/99ISR Number: 3427925-8Report Type:Periodic Company Report #A0077154  
 Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG	TWICE						
PER DAY	ORAL			Salbutamol Sulphate	C		

Date:03/01/99ISR Number: 3427929-5Report Type:Periodic Company Report #A0077233  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL	7 DAY	Dyspnoea					
		Speech Disorder					
		Throat Tightness					

Date:03/01/99ISR Number: 3427932-5Report Type:Periodic Company Report #A0077271  
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG	TWICE						
PER DAY	ORAL	Urticaria					

Date:03/01/99ISR Number: 3427936-2Report Type:Periodic Company Report #A0077337  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Other	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG	ORAL						

Date:03/01/99ISR Number: 3427938-6Report Type:Periodic Company Report #A0077338  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphemia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG	TWICE	Excitability					
PER DAY	ORAL	Headache					
		Insomnia					
		Tinnitus					

Date:03/01/99ISR Number: 3427940-4Report Type:Periodic Company Report #A0077339  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dissociation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG	PER	Sedation					
DAY	ORAL						

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Date:03/01/99ISR Number: 3431025-0Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #A0076174

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ PER							
DAY/ ORAL							

Fosinopril Sodium C

Date:03/01/99ISR Number: 3431027-4Report Type:Periodic  
Age:73 YR Gender:Male I/FU:I

Company Report #A0076175

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE			Professional				
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3431028-6Report Type:Periodic  
Age:53 YR Gender:Male I/FU:I

Company Report #A0076176

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Influenza Like Illness Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3431031-6Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #A0076177

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Rash Generalised	Professional				
PER DAY/ ORAL							

TRANSDERMAL 21 MG/ AS Nicotine Patch SS  
REQUIRED/  
TRANSDERMAL

Date:03/01/99ISR Number: 3431033-XReport Type:Periodic Company Report #A0076178  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Health Professional	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE				Simvastatin	C		
PER DAY/ ORAL				Estrostep 21	C		

Date:03/01/99ISR Number: 3431037-7Report Type:Periodic Company Report #A0076179  
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE				Atorvastatin Calcium	C		
PER DAY/ ORAL				Moduretic	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3431042-0Report Type:Periodic  
Age:25 YR Gender:Male I/FU:I

Company Report #A0076180

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ ORAL		Chest Discomfort	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/ PER DAY/ ORAL			Professional Company Representative	Wellbutrin (Formulation Unknown)	SS		ORAL

Date:03/01/99ISR Number: 3431045-6Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report # A0076181

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Dermatitis Urticaria	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:03/01/99ISR Number: 3431064-XReport Type:Periodic  
Age:69 YR Gender:Male I/FU:I

Company Report #A0076182

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Dry Mouth Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
				Polycarbophil Calcium Multivitamin	C C		

Date:03/01/99ISR Number: 3431074-2Report Type:Periodic  
Age:58 YR Gender:Male I/FU:I

Company Report #A0076183

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							
				Thyroxine Sodium	C		
				Digoxin	C		
				Blood Pressure Medication	C		

Date:03/01/99ISR Number: 3431077-8Report Type:Periodic Company Report #A0080697  
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depressed Mood	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							
		Dizziness					
		Epileptic Aura		Thyroxine Sodium	C		
		Insomnia		Premphase	C		
		Pallor		Pravastatin Sodium	C		
		Paralysis		Naproxen	C		
				Montelukast Sodium	C		
				Aspirin	C		
				Fluticasone			
				Propionate	C		
				Salmeterol Xinafoate	C		
				Salbutamol Sulphate	C		
				Methylprednisolone	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3431081-XReport Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #A0080698

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Insomnia					
		Rash Papular		Tetracycline Loestrin	C C		

Date:03/01/99ISR Number: 3431088-2Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0080699

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Pyrexia	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Urticaria					

Date:03/01/99ISR Number: 3431117-6Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #A0080700

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL		Disturbance In Attention	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Dizziness					
		Headache Vomiting					

Date:03/01/99ISR Number: 3431125-5Report Type:Periodic  
Age:68 YR Gender:Female I/FU:I

Company Report #A0080701

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Drug Dependence	Consumer	Zyban Tablet - Zyban	PS		ORAL

PER DAY/ ORAL

Fluticasone  
Propionate C  
Salbutamol Sulphate C  
Ipratropium Bromide C  
Budesonide C  
Glibenclamide C

Date:03/01/99ISR Number: 3431130-9Report Type:Periodic Company Report #A0080703  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ PER		Dizziness					
DAY/ ORAL		Eye Disorder Nightmare		Alprazolam	C		

Date:03/01/99ISR Number: 3431133-4Report Type:Periodic Company Report #A0080707  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ PER		Migraine					
DAY/ ORAL				Estrogen	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3431136-XReport Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0080710

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Arthralgia	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:03/01/99ISR Number: 3431140-1Report Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #A0080711

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Back Pain Pollakiuria	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:03/01/99ISR Number: 3431143-7Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0080713

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL		Crying Depression Irritability	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:03/01/99ISR Number: 3431146-2Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #A0080715

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Dizziness Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3431149-8Report Type:Periodic Company Report #A0080741  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diplopia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL		Dizziness Sensation Of Pressure Tremor					

Date:03/01/99ISR Number: 3431152-8Report Type:Periodic Company Report #A0080792  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diabetes Mellitus	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL	1 MON	Inadequate Control	Professional	Acarbose Glipizide Glucophage	C C C		

Date:03/01/99ISR Number: 3431157-7Report Type:Periodic Company Report #A0080794  
Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG SEE		Overdose					
TEXT ORAL		Tremor		Ortho Cyclen	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3431171-1Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #A0080795

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG	TWICE	Tension					
PER DAY	ORAL			Prempro	C		

Date:03/01/99ISR Number: 3431174-7Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0080796

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG		Dry Mouth					
TWICE PER DAY		Dysphagia					
ORAL		Foreign Body Trauma		Loratadine	C		
		Throat Irritation					

Date:03/01/99ISR Number: 3431196-6Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0080797

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Palpitations	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
.5 TABLET							
TWICE PER DAY							
ORAL				Sertraline			
				Hydrochloride	C		
				Diphenhydramine Hcl	C		
				Theophylline	C		
				Antihypertensive	C		

Date:03/01/99ISR Number: 3431208-XReport Type:Periodic Company Report #A0080798  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL				Sertaline Hydrochloride	C		
				Fexofenadine Hydrochlorid	C		

Date:03/01/99ISR Number: 3431228-5Report Type:Periodic Company Report #A0080799  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							
				Wellbutrin Tablet - Controlled Release	SS		ORAL
150 MG ORAL							

Date:03/01/99ISR Number: 3431231-5Report Type:Periodic Company Report #A0080800  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Effect Decreased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL				Atorvastatin Calcium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Prempro

C

Date:03/01/99ISR Number: 3431238-8Report Type:Periodic Company Report #A0080801  
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER							
DAY ORAL							

Hydroxyzine  
Theophylline

C  
C

Date:03/01/99ISR Number: 3431241-8Report Type:Periodic Company Report #A0080802  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL		Nervousness					

Date:03/01/99ISR Number: 3431245-5Report Type:Periodic Company Report #A0080803  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							

Date:03/01/99ISR Number: 3431247-9Report Type:Periodic Company Report #A0080804  
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							

Face Oedema  
Joint Swelling  
Penile Swelling  
Pruritus

Date:03/01/99ISR Number: 3431254-6Report Type:Periodic Company Report #A0080805  
Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Excitability					
DAY ORAL		Nervousness					

Date:03/01/99ISR Number: 3431294-7Report Type:Periodic Company Report #A0080808  
Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Pollakiuria					
PER DAY ORAL		Tremor		Clarithromycin	C		
				Nicotine	C		
				Salbutamol Sulphate	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3431297-2Report Type:Periodic Company Report #A0080867  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Throat	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Nasal Dryness					
PER DAY ORAL		Sleep Disorder					

Date:03/01/99ISR Number: 3431298-4Report Type:Periodic Company Report #A0080868  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG		Hypersensitivity					
TWICE PER DAY		Pruritus					
ORAL				Ibuprofen	C		
				Multivitamin	C		

Date:03/01/99ISR Number: 3431299-6Report Type:Periodic Company Report #A0080870  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG; ORAL	1 YR						

Date:03/01/99ISR Number: 3431301-1Report Type:Periodic Company Report #A0080871  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							

Urticaria

DAY; ORAL 5 DAY

Date:03/01/99ISR Number: 3431308-4Report Type:Periodic Company Report #A0080872  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Mental Disorder					
TWICE PER							

DAY; ORAL

Date:03/01/99ISR Number: 3431329-1Report Type:Periodic Company Report #A0080873  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG; ORAL		Hypersensitivity Pain In Extremity					

Date:03/01/99ISR Number: 3431332-1Report Type:Periodic Company Report #A0080874  
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Nervousness					
DAY; ORAL	3 DAY	Tremor		Omeprazole	C		
				Hormones	C		
				Vitamin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3431335-7Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0080875

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICER PER DAY; ORAL	3 DAY						

Date:03/01/99ISR Number: 3431336-9Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0080876

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY; ORAL		Constipation Disturbance In Attention Dizziness Headache Insomnia		Medroxyprogesterone Ace.	C		

Date:03/01/99ISR Number: 3431337-0Report Type:Periodic  
Age:72 YR Gender:Female I/FU:I

Company Report #A0080877

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY; ORAL		Insomnia Visual Disturbance		Vitamin	C		

Date:03/01/99ISR Number: 3431338-2Report Type:Periodic Company Report #A0080878  
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER							
DAY; ORAL							

Date:03/01/99ISR Number: 3431340-0Report Type:Periodic Company Report #A0080879  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							
DAY; ORAL							
Dizziness							
Headache							
Insomnia							
Irritability							
Major Depression							
Mental Impairment							
Nervousness							
Tension							
Tobacco Abuse							

Date:03/01/99ISR Number: 3431343-6Report Type:Periodic Company Report #A0080880  
Age: Gender:Female I/FU:I

Outcome	PT
	Accommodation Disorder
	Convulsion

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Discomfort Dry Mouth Hypersensitivity	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER		Muscle Rigidity	Professional				
DAY; ORAL		Muscle Spasms					
		Myalgia					
		Panic Disorder		Magnesium Salt	C		
		Syncope					
		Tetany					
		Tremor					
		Visual Disturbance					

Date:03/01/99ISR Number: 3431348-5Report Type:Periodic Company Report #A0080881  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER		Pruritus					
DAY; ORAL		Swelling					
		Urticaria					

Date:03/01/99ISR Number: 3431350-3Report Type:Periodic Company Report #A0080882  
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER		Acute Stress Disorder	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY; ORAL		Decreased Appetite					
		Eczema		Salbutamol Sulphate	C		
		Tachycardia		Asmacort	C		
				Guaiphenesin	C		
				Prempro	C		

Date:03/01/99ISR Number: 3431351-5Report Type:Periodic Company Report #A0080883  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER							
DAY; ORAL				Birth Control	C		

Date:03/01/99ISR Number: 3431356-4Report Type:Periodic Company Report #A0080884  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / SEE		Disturbance In Attention					
TEXT; ORAL		Insomnia		Etodolace	C		
				Atenolol	C		
				Conjugated Estrogens	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3431359-XReport Type:Periodic Company Report #A0080885  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER							
DAY; ORAL				Premphase	C		

Date:03/01/99ISR Number: 3431366-7Report Type:Periodic Company Report #A0080886  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG; ORAL		Myalgia	Professional Company Representative				

Date:03/01/99ISR Number: 3431369-2Report Type:Periodic Company Report #A0080887  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vertigo	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG; ORAL		Vomiting	Professional Company Representative				

Date:03/01/99ISR Number: 3431384-9Report Type:Periodic Company Report #A0080888  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG; ORAL  
Emotional Disorder      Health      Zyban      PS      Glaxo Wellcome Inc      ORAL  
Professional  
Company  
Representative

Date:03/01/99    ISR Number: 3431387-4    Report Type:Periodic      Company Report #A0080991  
Age:46 YR    Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Dry Skin					
TWICE PER		Face Oedema					
DAY; ORAL		Pruritus		Tramadol			
		Urticaria		Hydrochloride	C		
				Cyclobenzaprine Hcl	C		
				Conjugated Estrogens	C		

Date:03/01/99    ISR Number: 3431396-5    Report Type:Periodic      Company Report #A0075532  
Age:40 YR    Gender:Female      I/FU:I

Outcome	PT
	Abdominal Distension
	Anxiety
	Constipation
	Fluid Retention
	Insomnia



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Irritability Nervousness Skin Disorder	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE	PER DAY/ORAL		Consumer	Zyban Tablet-Zyban	PS		ORAL
				Frusemide	C		
				Cisapride	C		
				Psyllium Husk	C		
				Omeprazole	C		
				Lisinopril	C		
				Estrogen	C		
				Progesterone	C		
				Multivitamin	C		
				Thyroxine Sodium	C		
				Atorvastatin Calcium	C		
				Insulin	C		
				Timolol Maleate	C		
				Latanoprost	C		

Date:03/01/99ISR Number: 3431400-4Report Type:Periodic Company Report #A0075533  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE	PER DAY/ORAL	Insomnia	Consumer	Zyban Tablet-Zyban	PS		ORAL

Date:03/01/99ISR Number: 3431403-XReport Type:Periodic Company Report #A0075534  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/PER DAY/ORAL		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Blood Pressure Decreased					
		Dyspnoea		Verapamil			
		Headache		Hydrochloride	C		

Beclomethasone  
Dipropion C  
Salbutamol Sulphate C

Date:03/01/99ISR Number: 3431406-5Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #A0075535

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flatulence	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE		Peripheral Coldness					
PER DAY/ORAL		Skin Odour Abnormal		Wellbutrin Tablet	SS		ORAL
100 MG/ORAL				Gabapentin	C		
				Lorazepam	C		
				Pravastatin Sodium	C		
				Lactulose	C		
				Methocarbamol	C		
				Nicotine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3431417-XReport Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0075537

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/PER		Hypersensitivity					
DAY/ORAL		Syncope		Cefuroxime Axetil	C		
				Tylenol With Codeine	C		

Date:03/01/99ISR Number: 3431421-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0075538

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ORAL		Therapeutic Response Unexpected		Nicotine	C		

Date:03/01/99ISR Number: 3431427-2Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #A0075539

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE		Insomnia					
PER DAY/ORAL		Pruritus		Hydrochlorothiazide	C		
				Atenolol	C		

Date:03/01/99ISR Number: 3431436-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0075540

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ORAL							

Emotional Disorder

Date:03/01/99ISR Number: 3431438-7Report Type:Periodic Company Report #A0075541  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban Tablet-Zyban	PS		ORAL
150 MG/ORAL			Professional				

Date:03/01/99ISR Number: 3431444-2Report Type:Periodic Company Report #A0075542  
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Libido Decreased	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE		Oliguria					
PER DAY/ORAL				Atorvastatin Calcium	C		

Date:03/01/99ISR Number: 3431446-6Report Type:Periodic Company Report #A0075543  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/SINGLE		Dry Mouth					
DOSE/ORAL		Nervousness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3431457-0Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #A0075544

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/PER							
DAY/ORAL							

Triphasil C

Date:03/01/99ISR Number: 3431477-6Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0075545

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Health	Zyban Tablet-Zyban	PS		ORAL
150 MG/PER			Professional				
DAY/ORAL							

Oral Contraceptive C

Date:03/01/99ISR Number: 3431480-6Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0075546

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE		Insomnia					
PER DAY/ORAL							

Nausea  
Palpitations  
Tachycardia  
Calcium Salt C

Date:03/01/99ISR Number: 3431483-1Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0075547

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG/TWICE  
PER DAY/ORAL  
Dry Mouth  
Major Depression  
Paraesthesia  
Tremor  
Consumer  
Zyban Tablet-Zyban  
PS  
ORAL

Date:03/01/99ISR Number: 3431486-7Report Type:Periodic Company Report #A0075548  
Age:28 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE		Disturbance In Attention	Consumer	Zyban Tablet-Zyban	PS		ORAL
PER DAY/ORAL		Nervousness					
		Speech Disorder					
		Tremor					

Date:03/01/99ISR Number: 3431489-2Report Type:Periodic Company Report #A0075549  
Age:18 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE		Major Depression	Consumer	Zyban Tablet-Zyban	PS		ORAL
PER DAY/ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3431492-2Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #A0075550

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE		Decreased Appetite					
PER DAY/ORAL		Hallucination		Antibiotics	C		
		Mood Swings		Conjugated Estrogens	C		
		Nausea					
		Paraesthesia					
		Pruritus					
		Retching					

Date:03/01/99ISR Number: 3431493-4Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0075551

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Health	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE		Rash Erythematous	Professional				
PER DAY/ORAL		Urticaria					

Date:03/01/99ISR Number: 3432699-0Report Type:Periodic  
Age:34 YR Gender:Male I/FU:I

Company Report #A0075552

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Psychotic Disorder	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER			Professional				
DAY/ORAL							

Date:03/01/99ISR Number: 3432700-4Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #A0075536

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/PER							
DAY/ORAL							

Date:03/01/99ISR Number: 3432704-1Report Type:Periodic Company Report #A0075553  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Insomnia					
PER DAY/ORAL		Sedation		Naproxen	C		
				Theophylline	C		
				Glibenclamide	C		
				Salbutamol Sulphate	C		
				Fluticasone			
				Propionate	C		

Date:03/01/99ISR Number: 3432706-5Report Type:Periodic Company Report #A0075554  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Urticaria					
PER DAY/ORAL				Birth Control	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3432711-9Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #A0075555

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Fatigue Insomnia Libido Decreased Lymphadenopathy Weight Increased	Consumer	Zyban  Ascorbic Acid	PS  C	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3432714-4Report Type:Periodic  
Age:72 YR Gender:Male I/FU:I

Company Report #A0075556

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Agitation Disorientation Dizziness Dry Mouth	Health Professional	Zyban  Atenolol Prednisone	PS  C C	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3432718-1Report Type:Periodic  
Age:35 YR Gender:Male I/FU:I

Company Report #A0075557

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Dry Mouth Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3432722-3Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0075558

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Gingivitis	Health	Zyban	PS	Glaxo Wellcome Inc	

Date:03/01/99ISR Number: 3432733-8Report Type:Periodic Company Report #A0075568  
 Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Palpitations	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Urticaria	Professional				
PER DAY/ORAL							

Date:03/01/99ISR Number: 3432735-1Report Type:Periodic Company Report #A0075611  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Consumer	Zyban	PS	Glaxo Wellcome Inc	
		Headache					

Date:03/01/99ISR Number: 3432741-7Report Type:Periodic Company Report #A0076043  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT
		Face Oedema
		Headache
		Pain
		Pruritus
		Pyrexia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Rash Generalised

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ORAL						

Date:03/01/99ISR Number: 3432757-0Report Type:Periodic Company Report #A0076113  
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ORAL		Conjunctival Haemorrhage	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Professional Company Representative				

Date:03/01/99ISR Number: 3432768-5Report Type:Periodic Company Report #A0076163  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ORAL		Vision Blurred		Birth Control	C		

Date:03/01/99ISR Number: 3432775-2Report Type:Periodic Company Report #A0076165  
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ORAL		Nervousness					

Date:03/01/99ISR Number: 3432778-8Report Type:Periodic Company Report #A0076166  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Irritability					
PER DAY/ORAL		Tobacco Abuse		Simvastatin	C		
				Augmentin	C		
				Prednisone	C		

Date:03/01/99ISR Number: 3432780-6Report Type:Periodic Company Report #A0076167  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Pruritus					
PER DAY/ORAL				Metoprolol Succinate	C		
				Hydrochlorothiazide	C		

Date:03/01/99ISR Number: 3432784-3Report Type:Periodic Company Report #A0076169  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT
		Abnormal Dreams
		Conjunctival Hyperaemia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Ineffective Headache Insomnia	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Nervousness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Alprazolam	C
Clorazepate	
Dipotassium	C

Date:03/01/99ISR Number: 3432786-7Report Type:Periodic Company Report #A0076170  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ORAL			Professional				

Date:03/01/99ISR Number: 3432790-9Report Type:Periodic Company Report #A0076171  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER DAY/ORAL		Headache					

Date:03/01/99ISR Number: 3432792-2Report Type:Periodic Company Report #A0076172  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mood Altered	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE PER DAY/ORAL		Neck Pain					

Ibuprofen	C
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Date:03/01/99ISR Number: 3432795-8Report Type:Periodic Company Report #A0076173  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG.TWICE		Urticaria					
PER DAY/ORAL				Nabumetone	C		
				Thyroxine Sodium	C		
				Potassium Chloride	C		
				Amitriptyline	C		
				Diltiazem			
				Hydrochloride	C		

Date:03/01/99ISR Number: 3432811-3Report Type:Periodic Company Report #A0078830  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWIE							
PER DAY /							
ORAL				Paracetamol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3432815-0Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0078831

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE							
PER DAY /							
ORAL							
				Zestoretic	C		
				Estrogen	C		
				Tolterodine	C		
				Diphenhydramine Hcl	C		
				Potassium Salt	C		
				Arthrotec	C		

Date:03/01/99ISR Number: 3432818-6Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #A0078832

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Insulin	C		

Date:03/01/99ISR Number: 3432821-6Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #A0078834

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							
DAY / ORAL							
				Conjugated Estrogens	C		
				Medroxyprogesterone			
				Ace.	C		

Calcium Citrate C  
Alprazolam C

Date:03/01/99ISR Number: 3432825-3Report Type:Periodic Company Report #A0078835  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Aspirin C

Date:03/01/99ISR Number: 3432841-1Report Type:Periodic Company Report #A0078836  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Hypoaesthesia					
TWICE PER DAY		Vasodilatation					
/ ORAL							

Salmeterol Xinafoate C  
Salbutamol Sulphate C  
Famotidine C  
Beclomethasone  
Dipropion C



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3432845-9Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0078837

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Menorrhagia					
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3432846-0Report Type:Periodic  
Age:62 YR Gender:Male I/FU:I

Company Report #A0078838

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Nausea					
TWICE PER DAY							
/ ORAL							

Sinemet C

Date:03/01/99ISR Number: 3432851-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0078839

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Elevated Mood	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Insomnia					
TWICE PERDAY							
/ ORAL							

Date:03/01/99ISR Number: 3432852-6Report Type:Periodic  
Age:72 YR Gender:Male I/FU:I

Company Report #A0078840

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Nervousness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Tremor					
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3432869-1Report Type:Periodic Company Report #A0078841  
 Age:32 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							

Ibuprofen C

Date:03/01/99ISR Number: 3432871-XReport Type:Periodic Company Report #A0078842  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Nausea	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Vomiting	Professional				
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3432875-7Report Type:Periodic Company Report #A0078844  
 Age:55 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Flushing					
TWICE PER							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

DAY/ ORAL

Enalapril Maleate C  
 Imipramine C  
 Triamterene C  
 Alprazolam C  
 Prempro C

Date:03/01/99ISR Number: 3432877-0Report Type:Periodic Company Report #A0078861  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Influenza Like Illness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Pain					
TWICE PER							

DAY/ ORAL

Thyroid C

Date:03/01/99ISR Number: 3432880-0Report Type:Periodic Company Report #A0078880  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Menorrhagia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL							

Date:03/01/99ISR Number: 3432883-6Report Type:Periodic Company Report #A0078917  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Urticaria					
DAY / ORAL							

Date:03/01/99ISR Number: 3432935-0Report Type:Periodic Company Report #A0078918  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Nausea	Professional				
TWICE PER DAY							
/ ORAL	2	WK					

Date:03/01/99ISR Number: 3432938-6Report Type:Periodic Company Report #A0078919  
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Drug Ineffective					
TWICE PER DAY		Dry Mouth					
/ ORAL		Weight Increased		Frusemide	C		
				Colestipol			
				Hydrochloride	C		
				Lovastatin	C		
				Oxaprozin	C		
				Librium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3432940-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0078920

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3432943-XReport Type:Periodic  
 Age:54 YR Gender:Male I/FU:I

Company Report #A0078921

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3432961-1Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #A0078922

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Blood Pressure Increased	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL							
Dizziness							
Nausea							
Palpitations							

Date:03/01/99ISR Number: 3432962-3Report Type:Periodic  
 Age:31 YR Gender:Female I/FU:I

Company Report #A0078923

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Headache					
DAY / ORAL		Insomnia					

Date:03/01/99ISR Number: 3432964-7Report Type:Periodic Company Report #A0078942  
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Neuropathy Peripheral					
TWICE PER DAY		Paraesthesia					
/ ORAL				Metoprolol Tartrate	C		
				Isosorbide			
				Mononitrate	C		
				Aspirin	C		

Date:03/01/99ISR Number: 3432966-0Report Type:Periodic Company Report #A0078943  
 Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Aspirin	C		
				Metoprolol Tartrate	C		
				Diltiazem			
				Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3432968-4Report Type:Periodic  
 Age:51 YR Gender:Male I/FU:I

Company Report #A0078944

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disturbance In Attention	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3432970-2Report Type:Periodic  
 Age:33 YR Gender:Male I/FU:I

Company Report #A0078945

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Urticaria					
TWICE PER DAY							
/ ORAL				Buspirone Hydrochloride	C		

Date:03/01/99ISR Number: 3432971-4Report Type:Periodic  
 Age:28 YR Gender:Male I/FU:I

Company Report #A0078946

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Anxiety					
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3433096-4Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #1998013484-1

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Twitching	Health Professional	Paxil	PS	Smithkline Beecham Pharmaceuticals	ORAL
ORAL				Buprpion Glaxo Wellcome	SS	Glaxo Wellcome	

Date:03/01/99ISR Number: 3433106-4Report Type:Periodic Company Report #A0081596  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER DAY/ ORAL							

Date:03/01/99ISR Number: 3433108-8Report Type:Periodic Company Report #A0081599  
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Dysgeusia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE PER DAY / ORAL		Nervousness		Conjugated Estrogens	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433109-XReport Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0081641

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MJG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3433111-8Report Type:Periodic  
 Age: Gender:Not SpecifiI/FU:I

Company Report #A0081642

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Raynaud'S Phenomenon	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							
			Company Representative				

Date:03/01/99ISR Number: 3433113-1Report Type:Periodic  
 Age:32 YR Gender:Female I/FU:I

Company Report #A0081643

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Face Oedema	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER							
DAY/ ORAL							
			Urticaria				

Date:03/01/99ISR Number: 3433114-3Report Type:Periodic  
 Age:38 YR Gender:Male I/FU:I

Company Report #A0081644

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/	TWICE						
PER DAY /							
ORAL							

Date:03/01/99ISR Number: 3433116-7Report Type:Periodic Company Report #A0081645  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain Hypoaesthesia		Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /	ORAL	Migraine Tremor					

Date:03/01/99ISR Number: 3433117-9Report Type:Periodic Company Report #A0081646  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Urticaria					
TWICE PER DAY							
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433118-0Report Type:Periodic  
Age:57 YR Gender:Male I/FU:I

Company Report #A0081647

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER DAY / ORAL				Salbutamol Sulphate	C		
				Prednisone	C		

Date:03/01/99ISR Number: 3433121-0Report Type:Periodic  
Age:68 YR Gender:Female I/FU:I

Company Report #A0081648

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Nervousness	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE PER DAY/ORAL		Tachycardia		Quinine	C		
				Blood Pressure Medication	C		
				Prednisone	C		
				Diclofenac Sodium	C		
				Hydrochlorothiazide	C		

Date:03/01/99ISR Number: 3433123-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0081654

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Anxiety	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY/ ORAL		Drug Effect Decreased					
		Dyspnoea					
		Nervousness					

Tremor

Date:03/01/99ISR Number: 3433126-XReport Type:Periodic Company Report #A0081655  
Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Bupropion			
150 MG /		Nervousness		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL				Paroxetine			
				Hydrochloride	C		

Date:03/01/99ISR Number: 3433127-1Report Type:Periodic Company Report #A0081657  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Bupropion			
150 MG /				Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433130-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0081658

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eczema Tachycardia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER DAY/ ORAL							

Date:03/01/99ISR Number: 3433131-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0081659

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State Dizziness Nausea	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /ORAL							

Date:03/01/99ISR Number: 3433134-9Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #A0081913

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY / ORAL							

Date:03/01/99ISR Number: 3433135-0Report Type:Periodic  
 Age:44 YR Gender:Female I/FU:I

Company Report #A0081915

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Bupropion			

150 MG /  
TWICE PER DAY  
/ ORAL  
Oedema Peripheral  
Urticaria  
Hydrochloride  
PS  
Glaxo Wellcome Inc  
ORAL

Date:03/01/99ISR Number: 3433137-4Report Type:Periodic Company Report #A0081916  
Age:39 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Insomnia Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

TWICE PER DAY  
/ ORAL

Date:03/01/99ISR Number: 3433138-6Report Type:Periodic Company Report #A0081918  
Age:44 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY/ ORAL		Disorientation Dizziness Nausea	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433141-6Report Type:Periodic  
 Age:55 YR Gender:Female I/FU:I

Company Report #A0081919

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Agitation Cough  Increased Appetite  Weight Increased	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Nabumetone	C		
				Multivitamin	C		
				Estrogen	C		

Date:03/01/99ISR Number: 3433143-XReport Type:Periodic  
 Age:50 YR Gender:Female I/FU:I

Company Report #A0078424

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ORAL		Dry Mouth Groin Pain  Insomnia  Renal Colic	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433145-3Report Type:Periodic  
 Age:61 YR Gender:Female I/FU:I

Company Report #A0078425

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Sedation		Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433147-7Report Type:Periodic Company Report #A0078426  
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Euphoric Mood Insomnia		Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
PER DAY/ORAL				Azithromycin	C		

Date:03/01/99ISR Number: 3433148-9Report Type:Periodic Company Report #A0078427  
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3433150-7Report Type:Periodic Company Report #A0078428  
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Amnesia Condition Aggravated	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Cough					
TWICE PER DAY		Thinking Abnormal					
/ ORAL				Verapamil			



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride C  
 Enalapril Maleate C  
 Alprazolam C

Date:03/01/99ISR Number: 3433151-9Report Type:Periodic Company Report #A0078429  
 Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urticaria	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3433153-2Report Type:Periodic Company Report #A0078430  
 Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Haematuria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE							
PER DAY /							
ORAL							

Date:03/01/99ISR Number: 3433156-8Report Type:Periodic Company Report #A0078431  
 Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Apathy Lethargy	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3433158-1Report Type:Periodic Company Report #A0078432  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Burning Sensation Pruritus	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Urticaria					
TWICE PER							
DAY/ ORAL							

Date:03/01/99ISR Number: 3433160-XReport Type:Periodic Company Report #A0078433  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mental Impairment	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER							
DAY/ORAL				Etodolac	C		

Date:03/01/99ISR Number: 3433163-5Report Type:Periodic Company Report #A0078434  
Age:52 YR Gender:Male I/FU:I

Outcome	PT
	Insomnia Paranoia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Psychotic Disorder Sleep Disorder	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER							
DAY/ ORAL							

Date:03/01/99ISR Number: 3433172-6Report Type:Periodic Company Report #A0077570  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL							

Date:03/01/99ISR Number: 3433174-XReport Type:Periodic Company Report #A0077573  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3433175-1Report Type:Periodic Company Report #A0077574  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL				Ortho Tri-Cyclen	C		

Date:03/01/99ISR Number: 3433203-3Report Type:Periodic Company Report #A0077575  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Sedation					
PER DAY/ ORAL		Skin Warm					

Date:03/01/99ISR Number: 3433206-9Report Type:Periodic Company Report #A0077577  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Diarrhoea					
PER DAY/ ORAL				Azithromycin (Formulation Unknown)	SS		

Date:03/01/99ISR Number: 3433207-0Report Type:Periodic Company Report #A0077578  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Migraine	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Nausea					
DAY/ ORAL		Vision Blurred					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433210-0Report Type:Periodic  
 Age:23 YR Gender:Female I/FU:I

Company Report #A0077580

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY/ ORAL		Dermatitis Hypersensitivity Pruritus Tremor	Consumer	Zyban  Birth Control Omeprazole	PS  C C	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433212-4Report Type:Periodic  
 Age:64 YR Gender:Female I/FU:I

Company Report #A0077581

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Estratest Medroxyprogesterone Ace. Cimetidine Oxybutynin Hydrochloride	C C C C		

Date:03/01/99ISR Number: 3433213-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0077583

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ SINGLE DOSE/ ORAL		Alcoholic Hangover Anxiety Dizziness Feeling Drunk Palpitations Panic Reaction Tachycardia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433216-1Report Type:Periodic Company Report #A0077584  
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Dry Mouth					
DAY/ ORAL		Headache		Omeprazole	C		
		Tremor		Methocarbamol	C		
				Vicodin	C		

Date:03/01/99ISR Number: 3433218-5Report Type:Periodic Company Report #A0077585  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Face Oedema					
PER DAY/ ORAL		Pruritus		Lorazepam	C		

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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433221-5Report Type:Periodic Company Report #A0077586  
 Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Herpes Virus Infection					
PER DAY/ ORAL		Panic Reaction					

Date:03/01/99ISR Number: 3433223-9Report Type:Periodic Company Report #A0077588  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Discomfort	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Insomnia					
DAY/ ORAL		Mental Impairment					
		Restlessness					
		Tremor					

Date:03/01/99ISR Number: 3433226-4Report Type:Periodic Company Report #A0077617  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea Exertional	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Emphysema					
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3433228-8Report Type:Periodic Company Report #A0077659  
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL							

Swelling  
Urticaria

Professional  
Company  
Representative

Date:03/01/99ISR Number: 3433230-6Report Type:Periodic Company Report #A0077661  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Swelling	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL			Professional Company Representative				

Date:03/01/99ISR Number: 3433233-1Report Type:Periodic Company Report #A0077672  
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decreased Appetite	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Extrapyramidal Disorder					
PER DAY/ ORAL		Feeding Problem In Child					
		Hallucination					
		Lethargy					
		Memory Impairment					
		Petit Mal Epilepsy					



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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433236-7Report Type:Periodic Company Report #A0077673  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ ORAL	2 WK	Chest Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Tachycardia					

Date:03/01/99ISR Number: 3433238-0Report Type:Periodic Company Report #A0077686  
 Age:74 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ ORAL		Unevaluable Event	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433240-9Report Type:Periodic Company Report #A0077687  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ ORAL	11 DAY	Anxiety	Other	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Insomnia					
		Pain					

Date:03/01/99ISR Number: 3433255-0Report Type:Periodic Company Report #A0082002  
 Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ PER		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY/ ORAL							

Date:03/01/99ISR Number: 3433256-2Report Type:Periodic Company Report #A0082003  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lethargy	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL				Conjugated Estrogens	C		
				Thyroxine Sodium	C		

Date:03/01/99ISR Number: 3433259-8Report Type:Periodic Company Report #A0082004  
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Dry Mouth					
PER DAY/ ORAL		Headache		Lorazepam	C		

Date:03/01/99ISR Number: 3433260-4Report Type:Periodic Company Report #A0082005  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Breast Tenderness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Headache					
DAY/ ORAL		Insomnia		Ibuprofen	C		
		Tenderness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433262-8Report Type:Periodic Company Report #A0082006  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/		Pollakiuria					
TWICE PER							
DAY/ ORAL							

Date:03/01/99ISR Number: 3433263-XReport Type:Periodic Company Report #A0082022  
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Hypersensitivity					
PER DAY/ ORAL		Oedema Peripheral					

Date:03/01/99ISR Number: 3433264-1Report Type:Periodic Company Report #A0082023  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3433265-3Report Type:Periodic Company Report #A0082087  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							

PER DAY/ ORAL

Date:03/01/99ISR Number: 3433266-5Report Type:Periodic Company Report #A0082267  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							

DAY/ ORAL

Date:03/01/99ISR Number: 3433268-9Report Type:Periodic Company Report #A0082272  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Psoriasis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							

DAILY/ ORAL

Date:03/01/99ISR Number: 3433269-0Report Type:Periodic Company Report #A0079578  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menstrual Disorder	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							

PER DAY/ ORAL

Thyroxine Sodium C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433271-9Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #A0079579

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL		Agitation Tremor	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433272-0Report Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #A0079580

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL		Dry Mouth Feeling Hot Insomnia Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433274-4Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0079581

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ THREE TIMES PER DAY/ ORAL		Vomiting	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Cimetidine	C
Tetracycline	C
Metronidazole	C
Conjugated Estrogens	C

Date:03/01/99ISR Number: 3433276-8Report Type:Periodic Company Report #A0079582  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							
DAY/ ORAL							

Date:03/01/99ISR Number: 3433278-1Report Type:Periodic Company Report #A0079583  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Palpitations	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							
Tachycardia							
Hrt							
Multivitamin							
C							
C							

Date:03/01/99ISR Number: 3433279-3Report Type:Periodic Company Report #A0079584  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							
Nausea							
Salbutamol Sulphate							
C							
Fluticasone							
Propionate							
C							
Hydroxyzine							
Hydrochloride							
C							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433281-1Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0079585

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL		Rash Papular	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Quinapril Hydrochloride	C		
				Glipizide	C		
				Dicloxacillin	C		

Date:03/01/99ISR Number: 3433282-3Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0079586

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Diazepam	C		
				Meclozine	C		

Date:03/01/99ISR Number: 3433284-7Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #A0079587

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433436-6Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0074714

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/PER			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY/ORAL							
Date:03/01/99ISR Number: 3433437-8Report Type:Periodic Company Report #A0074885							
Age:50 YR Gender:Female I/FU:I							
150 MG/TWICE		Disturbance In Attention	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ORAL		Dizziness					
		Headache					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ORAL		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Micturition Urgency					
				Multivitamin	C		
				Guaiphenesin	C		
				Combivent	C		



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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433439-1Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0074890

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
PER DAY/ORAL							
				Atenolol	C		
				Hydrochlorothiazide	C		

Date:03/01/99ISR Number: 3433440-8Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #A0074942

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ORAL							
		Disorientation		Alprazolam	C		
		Fear					
		Paraesthesia					

Date:03/01/99ISR Number: 3433441-XReport Type:Periodic  
Age:22 YR Gender:Female I/FU:I

Company Report #A0074943

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
PER DAY/ORAL							
				Ethanol	SS		
				Ortho Cyclen	C		

Date:03/01/99ISR Number: 3433442-1Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0074956

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150 MG/TWICE		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Urticaria					
	PER DAY/ORAL				Isoniazid	C		
Date:03/01/99ISR Number: 3433443-3Report Type:Periodic				Company Report #A0074978				
Age:48 YR Gender:Female I/FU:I								
	150 MG/ORAL		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Urticaria	Health Professional Other	Nicotine	C		
Date:03/01/99ISR Number: 3433444-5Report Type:Periodic				Company Report #A0074980				
Age:33 YR Gender:Female I/FU:I								
	150 MG/TWICE		Dermatitis	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
					Loestrin	C		
	PER DAY/ORAL							

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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433445-7Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0074982

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER		Nervousness					
DAY/ORAL				Prednisone	C		
				Cimetidine	C		
				Acetazolamide	C		
				Doxepin	C		

Date:03/01/99ISR Number: 3433446-9Report Type:Periodic  
Age:58 YR Gender:Male I/FU:I

Company Report #A0074983

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Psychomotor Hyperactivity					
PER DAY/ORAL		Sedation		Fosinopril Sodium	C		

Date:03/01/99ISR Number: 3433447-0Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0074984

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER		Dissociation					
DAY/ORAL		Insomnia		Clarithromycin Tablet	SS		ORAL
ORAL				Multivitamin	C		
				Percodan	C		
				Nitroglycerin	C		

Date:03/01/99ISR Number: 3433448-2Report Type:Periodic Company Report #A0074985  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Erythema	Professional				
PER DAY/ORAL		Oedema Peripheral Skin Irritation					

Date:03/01/99ISR Number: 3433449-4Report Type:Periodic Company Report #A0074986  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sensory Disturbance	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Tinnitus					
PER DAY/ORAL		Tremor		Sertraline Hydrochloride Ibuprofen	C C		

Date:03/01/99ISR Number: 3433450-0Report Type:Periodic Company Report #A0074987  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depressed Level Of Consciousness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Dysgeusia		Enalapril Maleate	C		
PER DAY/ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Conjugated Estrogens C

Date:03/01/99ISR Number: 3433451-2Report Type:Periodic Company Report #A0074988  
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Urticaria					
PER DAY/ORAL							

Date:03/01/99ISR Number: 3433452-4Report Type:Periodic Company Report #A0074989  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Disturbance In Attention	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ORAL		Drug Ineffective					
		Headache					
		Nausea					

Date:03/01/99ISR Number: 3433453-6Report Type:Periodic Company Report #A0074991  
 Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
PER DAY/ORAL							

Date:03/01/99ISR Number: 3433454-8Report Type:Periodic Company Report #A0074992  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG/SEE	Chest Pain	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
TEXT/ORAL	Dizziness	Professional				
	Nausea		Paracetamol	C		
	Visual Disturbance					
	Vomiting					

Date:03/01/99ISR Number: 3433456-1Report Type:Periodic Company Report #A0074993  
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER		Irritability					
DAY/ORAL				Salbutamol Sulphate	C		
				Theophylline	C		

Date:03/01/99ISR Number: 3433457-3Report Type:Periodic Company Report #A0058433  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL	2 WK	Ill-Defined Disorder					

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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433458-5Report Type:Periodic Company Report #A0059740  
 Age:52 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Abdominal Pain Upper	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Estrogen	C		
				Naproxen Sodium	C		

Date:03/01/99ISR Number: 3433460-3Report Type:Periodic Company Report #A0061525  
 Age:27 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Arrhythmia Chest Pain Disorientation Dizziness Hyperhidrosis Hypoaesthesia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433461-5Report Type:Periodic Company Report #A0062656  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433462-7Report Type:Periodic Company Report #A0062756  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG SEE TEXT ORAL		Malaise Nervousness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Tremor  
Vomiting

Date:03/01/99ISR Number: 3433463-9Report Type:Periodic Company Report #A0062799  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL	3 WK						

Date:03/01/99ISR Number: 3433464-0Report Type:Periodic Company Report #A0063424  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3433466-4Report Type:Periodic Company Report #A0066887  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL	2 WK		Professional				

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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433467-6Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #A0066916

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
				Estrogen	C		
				Alprazolam	C		

Date:03/01/99ISR Number: 3433468-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0069007

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL	8 WK						

Date:03/01/99ISR Number: 3433469-XReport Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #A0081373

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Excitability	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							

Date:03/01/99ISR Number: 3433470-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0081374

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							
		Dry Throat					

Date:03/01/99ISR Number: 3433471-8Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #A0081387

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL		Oedema Peripheral	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Triamterene	C		

Date:03/01/99ISR Number: 3433474-3Report Type:Periodic Company Report #A0081394  
 Age:45 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Paraesthesia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL				Premphase	C		

Date:03/01/99ISR Number: 3433475-5Report Type:Periodic Company Report #A0081398  
 Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER		Accommodation Disorder	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY ORAL		Diarrhoea					
		Disorientation		Fluticasone			
		Dyspepsia		Propionate	C		
		Headache					
		Nausea					
		Tremor					

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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433478-0Report Type:Periodic Company Report #A0081402  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Migraine	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3433479-2Report Type:Periodic Company Report #A0081494  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
				Estrogen	C		

Date:03/01/99ISR Number: 3433480-9Report Type:Periodic Company Report #A0081516  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
		Insomnia					
		Psychomotor Hyperactivity		Thyroxine Sodium	C		

Date:03/01/99ISR Number: 3433483-4Report Type:Periodic Company Report #A0081521  
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3433486-XReport Type:Periodic Company Report #A0081585  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Irritability	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3433488-3Report Type:Periodic Company Report #A0074994  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disturbance In Attention	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE		Dysarthria					
PER DAY ORAL		Dyspraxia		Clonazepam	C		
		Headache		Lorazepam	C		
		Insomnia		Gabapentin	C		
		Speech Disorder					

Date:03/01/99ISR Number: 3433489-5Report Type:Periodic Company Report #A0080347  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoaesthesia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Hypoaesthesia Oral					
PER DAY ORAL							



150 MG TWICE	Crying	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL	Depression					
	Insomnia		Hyzaar	C		
	Suicidal Ideation		Alpha-Anilinophenyl Acet.	C		
			Vitamin	C		
			Buspirone			
			Hydrochloride	C		

Date:03/01/99ISR Number: 3433501-3Report Type:Periodic Company Report #A0075016  
 Age:41 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
150 MG TWICE	Depression	Consumer	Zyban Tablet - Zyban	PS		ORAL
PER DAY ORAL	Dermatitis					
	Disturbance In Attention					

Date:03/01/99ISR Number: 3433502-5Report Type:Periodic Company Report #A0080356  
 Age:54 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
150 MG TWICE	Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL	Urticaria					
			Dyazide	C		

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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433503-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0080357

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG ORAL	5 DAY	Dysgeusia Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433506-2Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0080358

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG ORAL		Diarrhoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433508-6Report Type:Periodic  
 Age:62 YR Gender:Female I/FU:I

Company Report #A0080359

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG TWICE PER DAY ORAL		Cough Disturbance In Attention Dry Mouth Dry Skin Ear Disorder Ear Haemorrhage Haemorrhage Keratoconjunctivitis Sicca Tongue Disorder	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433510-4Report Type:Periodic  
 Age:28 YR Gender:Female I/FU:I

Company Report #A0080360

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG TWICE  
PER DAY ORAL  
Dermatitis  
Urticaria  
Consumer  
Zyban  
PS  
Glaxo Wellcome Inc  
ORAL

Date:03/01/99ISR Number: 3433512-8Report Type:Periodic Company Report #A0080361  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphagia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL		Foreign Body Trauma					

Date:03/01/99ISR Number: 3433514-1Report Type:Periodic Company Report #A0075017  
Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE		Insomnia					
PER DAY ORAL		Productive Cough					

Date:03/01/99ISR Number: 3433516-5Report Type:Periodic Company Report #A0080362  
Age:59 YR Gender:Female I/FU:I

Outcome	PT
	Cough
	Dry Mouth



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Fatigue Insomnia Nervousness Pruritus	Health Professional	Zyban Prempro	PS C	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433517-7Report Type:Periodic Company Report #A0075019  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Dysgeusia	Consumer	Zyban Tablet - Zyban	PS		ORAL
				Glucophage	C		
				Conjugated Estrogens	C		
				Medroxyprogesterone			
				Ace.	C		
				Vitamin	C		
				Hypericum	C		
				Ginkgo Biloba	C		

Date:03/01/99ISR Number: 3433519-0Report Type:Periodic Company Report #A0080363  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Constipation Diarrhoea Insomnia Mood Altered	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433520-7Report Type:Periodic Company Report #A0075024  
 Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3433521-9Report Type:Periodic Company Report #A0080364  
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mydriasis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER							
DAY ORAL				Theraflu	C		

Date:03/01/99ISR Number: 3433524-4Report Type:Periodic Company Report #A0075028  
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral	Health Professional	Zyban Tablet - Zyban	PS		ORAL
150 MG ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433525-6Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #A0080365

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Agitation Disturbance In Attention Dizziness Eating Disorder Eructation Intestinal Functional Disorder Nausea Sedation Vertigo	Consumer	Zyban  Loratadine Fluticasone Propionate	PS  C C	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433527-XReport Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #A0075030

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Dysuria Vomiting	Consumer	Zyban Tablet - Zyban  Frusemide Simvastatin	PS  C C		ORAL

Date:03/01/99ISR Number: 3433528-1Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #A0080366

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433531-1Report Type:Periodic Company Report #A0075032  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Generalised	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3433532-3Report Type:Periodic Company Report #A0080367  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL		Flatulence					

Date:03/01/99ISR Number: 3433534-7Report Type:Periodic Company Report #A0080368  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Feeling Abnormal					
PER DAY ORAL		Insomnia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433535-9Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #A0077340

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dissociation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TOTAL		Headache					
TWICE PER DAY		Sedation					
ORAL				Birth Control	C		

Date:03/01/99ISR Number: 3433538-4Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #A0077342

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE		Nausea					
PER DAY ORAL		Tremor		Birth Control	C		

Date:03/01/99ISR Number: 3433539-6Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #A0080369

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anaphylactoid Reaction	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Bronchospasm					
PER DAY ORAL		Dyspnoea		Diphenhydramine Hcl	C		
		Rash Erythematous		Multivitamin	C		
		Urticaria		Allegra-D	C		

Date:03/01/99ISR Number: 3433541-4Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #A0077343

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cough	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE		Dry Throat					
PER DAY ORAL							

Insulin	C
Asmacort	C
Ipratropium Bromide	C
Loratadine	C
Loratadine	C
Tramadol	
Hydrochloride	C
Amitriptyline Hcl	C
Orciprenaline	
Sulphate	C

Date:03/01/99ISR Number: 3433542-6Report Type:Periodic Company Report #A0080370  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Pruritus					
PER DAY ORAL							

Date:03/01/99ISR Number: 3433544-XReport Type:Periodic Company Report #A0080371  
Age:39 YR Gender:Female I/FU:I

Outcome	PT
	Drug Dependence
	Insomnia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Irritability Nervousness	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3433545-1Report Type:Periodic Company Report #A0077344  
 Age:54 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration		Consumer	Zyban Tablet - Zyban	PS		ORAL
Dose		Elevated Mood					
150 MG TWICE		Malaise					
PER DAY		Sedation					

Date:03/01/99ISR Number: 3433548-7Report Type:Periodic Company Report #A0080372  
 Age:44 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Dose		Tremor					
150 MG PER							
DAY ORAL							

Date:03/01/99ISR Number: 3433550-5Report Type:Periodic Company Report #A0077346  
 Age:67 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration		Consumer	Zyban Tablet - Zyban	PS		ORAL
Dose		Condition Aggravated					
150 MG TWICE		Constipation					
PER DAY ORAL		Dermatitis					
		Ill-Defined Disorder					

Insomnia  
Nervousness  
Tobacco Abuse

Date:03/01/99ISR Number: 3433552-9Report Type:Periodic Company Report #A0077348  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3433556-6Report Type:Periodic Company Report #A0077439  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bronchospasm	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG PER							
DAY ORAL							

Date:03/01/99ISR Number: 3433559-1Report Type:Periodic Company Report #A0077565  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE							
PER DAY ORAL							
Insomnia							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433562-1Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #A0077566

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE		Depression					
PER DAY ORAL							

Date:03/01/99ISR Number: 3433563-3Report Type:Periodic  
Age:60 YR Gender:Male I/FU:I

Company Report #A0077567

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE		Anxiety					
PER DAY ORAL		Insomnia					
		Tremor					

Date:03/01/99ISR Number: 3433566-9Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #A0079000

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3433570-0Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0079001

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG PER							
DAY ORAL							

Cetirizine  
Hydrochloride C  
Nasonex C  
Duratuss C

Date:03/01/99ISR Number: 3433573-6Report Type:Periodic Company Report #A0079003  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 NG THREE							
TIMES PER DAY							
ORAL							

Librax C

Date:03/01/99ISR Number: 3433574-8Report Type:Periodic Company Report #A0079006  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE		Insomnia					
PER DAY ORAL		Panic Disorder					
		Tachycardia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433578-5Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0079088

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE		Emotional Disorder					
PER DAY ORAL		Motion Sickness		Lo/Ovral	C		
		Nausea		Multivitamin	C		
		Tremor					

Date:03/01/99ISR Number: 3433580-3Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #A0079089

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG PER		Tension					
DAY ORAL				Conjugated Estrogens	C		

Date:03/01/99ISR Number: 3433583-9Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #A0079091

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE		Dyspnoea					
PER DAY ORAL		Thinking Abnormal		Medroxyprogesterone			
				Ace.	C		
				Estrogen	C		
				Salbutamol Sulphate	C		

Date:03/01/99ISR Number: 3433584-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0079099

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
150 MG ORAL	3 WK	Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Hypersensitivity					
		Pruritus					
		Swelling					

Date:03/01/99ISR Number: 3433585-2Report Type:Periodic Company Report #A0079100  
Age: Gender: I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
150 MG ORAL		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Hypersensitivity					
		Pruritus					
		Swelling					

Date:03/01/99ISR Number: 3433587-6Report Type:Periodic Company Report #A0079135  
Age: Gender: I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
150 MG ORAL		Hypersensitivity	Consumer	Zyban Tablet - Zyban	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433588-8Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0079181

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG TWICE		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
PER DAY ORAL		Urticaria					

Date:03/01/99ISR Number: 3433590-6Report Type:Periodic  
Age:65 YR Gender:Female I/FU:I

Company Report #A0079182

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG ORAL		Tremor	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:03/01/99ISR Number: 3433591-8Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #A0080994

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG ORAL		Angioneurotic Oedema	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Arthralgia	Professional				
		Erythema	Company				
		Oedema Peripheral	Representative				
		Pain In Extremity					
		Pruritus					
		Urticaria					

Date:03/01/99ISR Number: 3433592-XReport Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #A0079183

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG TWICE		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
PER DAY ORAL							

Atorvastyatin  
 Calcium C  
 Maxzide C  
 Conjugated Estrogens C  
 Cimetidine C

Date:03/01/99ISR Number: 3433593-1Report Type:Periodic Company Report #A0079184  
 Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG PER		Irritability					
DAY ORAL							

Date:03/01/99ISR Number: 3433595-5Report Type:Periodic Company Report #A0081079  
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Extrasystoles					
DAY ORAL		Heart Rate Increased		Thyroxine Sodium	C		
		Insomnia		Clonazepam	C		
		Palpitations		Fluoxetine			
		Polydipsia		Hydrchloride	C		
		Tachycardia		Nefazodone			
		Tremor		Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433596-7Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0079185

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG PER							
DAY ORAL							

Omeprazole C

Date:03/01/99ISR Number: 3433598-0Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #A0079186

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE							
PER DAY ORAL							

Atorvastatin Calcium C  
Ticlopidine C  
Metoprolol Succinate C

Date:03/01/99ISR Number: 3433599-2Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #A0081085

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disturbance In Attention	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Memory Impairment					
DAY ORAL		Tremor					

Glibenclamide C  
Quinine Sulphate C  
Conjugated Estrogens C  
Aspirin C

Date:03/01/99ISR Number: 3433600-6Report Type:Periodic  
Age:31 YR Gender:Male I/FU:I

Company Report #A0079187

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE							
PER DAY ORAL				Gemfibrozil	C		

Date:03/01/99ISR Number: 3433601-8Report Type:Periodic Company Report #A0081090  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Dry Mouth					
PER DAY ORAL		Insomnia					
		Tremor					

Date:03/01/99ISR Number: 3433602-XReport Type:Periodic Company Report #A0079188  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE		Jealous Delusion					
PER DAY ORAL		Nervousness		Ibuprofen	C		
		Paranoia		Anacin	C		
		Suicidal Ideation					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433603-1Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0081096

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Myalgia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3433604-3Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #A0081097

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER							
DAY ORAL							
Syncope							
Vision Blurred							

Date:03/01/99ISR Number: 3433605-5Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0079213

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE							
PER DAY ORAL							
Constipation							
Libido Decreased							
Nicotine							
Trifluoperazine Hcl							
C							
C							

Date:03/01/99ISR Number: 3433606-7Report Type:Periodic  
Age:34 YR Gender:Male I/FU:I

Company Report #A0081099

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							

Pruritus

PER DAY ORAL

Date:03/01/99ISR Number: 3433607-9Report Type:Periodic Company Report #A0079214  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE		Medication Error					

PER DAY ORAL

Date:03/01/99ISR Number: 3433608-0Report Type:Periodic Company Report #A0081101  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Anxiety					
DAY ORAL		Tremor		Fenofibrate	C		
				Amlodipine Besylate	C		
				Alprazolam	C		
				Aspirin	C		

Date:03/01/99ISR Number: 3433610-9Report Type:Periodic Company Report #A0081115  
Age:46 YR Gender:Female I/FU:I

Outcome	PT
	Eye Irritation
	Keratoconjunctivitis
	Sicca

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vision Blurred

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG PER		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY ORAL			Wellbutrin Tablet	SS		ORAL
100 MG PER			Multivitamin	C		
DAY ORAL			Kelp	C		
			Alfalfa	C		

Date:03/01/99ISR Number: 3433611-0Report Type:Periodic Company Report #A0079252  
 Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Vomiting		Famotidine	C		
DAY / ORAL							

Date:03/01/99ISR Number: 3433612-2Report Type:Periodic Company Report #A0081148  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nightmare	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL			Professional Company Representative				

Date:03/01/99ISR Number: 3433614-6Report Type:Periodic Company Report #A0079253  
 Age:67 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Balance Disorder	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Dizziness					
/ ORAL		Insomnia		Montelukast Sodium	C		

Date:03/01/99ISR Number: 3433616-XReport Type:Periodic Company Report #A0079256  
 Age:32 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Headache	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Nausea					
/ ORAL		Vision Blurred					

Date:03/01/99ISR Number: 3433620-1Report Type:Periodic Company Report #A0079300  
 Age:40 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Constipation	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Dizziness					
/ ORAL		Dry Mouth					
		Dysgeusia		Darvocet-N	C		
		Nausea					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433626-2Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0079301

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3433634-1Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0079302

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Hypertension					
TWICE PER DAY		Tachycardia					
/ ORAL							
				Trazodone	C		
				Amlodipine	C		
				Prempro	C		
				Fluphenazine Hcl	C		

Date:03/01/99ISR Number: 3433637-7Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0079303

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Medroxyprogesterone			
				Ace	C		

Date:03/01/99ISR Number: 3433641-9Report Type:Periodic Company Report #A0079304  
Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pruritus					
TWICE PER DAY							
/ ORAL							
				Lisinopril	C		
				Atorvastatin Calcium	C		

Date:03/01/99ISR Number: 3433643-2Report Type:Periodic Company Report #A0079305  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Other	Zyban Tablet - Zyban	PS		ORAL
150 MG / UNK							
/ ORAL							

Date:03/01/99ISR Number: 3433646-8Report Type:Periodic Company Report #A0079306  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / UNK			Professional				
/ ORAL			Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433648-1Report Type:Periodic  
Age:60 YR Gender:Male I/FU:I

Company Report #A0079307

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Cough Dry Mouth Feeling Hot Oral Soft Tissue Disorder	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:03/01/99ISR Number: 3433650-XReport Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #A0079308

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Disorientation Disturbance In Attention Dizziness Photosensitivity Reaction Tremor Vision Blurred	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:03/01/99ISR Number: 3433668-7Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #A0079450

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Anxiety Chest Pain Dizziness Emotional Disorder Insomnia	Consumer	Zyban Tablet - Zyban Maxzide	PS C		ORAL

Date:03/01/99ISR Number: 3433672-9Report Type:Periodic Company Report #A0079451  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Difficulty In Walking	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Face Oedema					
TWICE PER DAY		Oedema					
/ ORAL		Urticaria					

Date:03/01/99ISR Number: 3433676-6Report Type:Periodic Company Report #A0081149  
Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Dizziness					
PER DAY ORAL		Hyperhidrosis					
		Nausea					

Date:03/01/99ISR Number: 3433677-8Report Type:Periodic Company Report #A0079452  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Other	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Nervousness					
DAY / ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433681-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0079572

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depressed Mood	Consumer	Zyban Tablet - Zyban	PS		ORAL
1450 MG /		Depression					
UNK/ ORAL		Mood Altered					

Date:03/01/99ISR Number: 3433683-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0079573

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / UNK		Nervousness					
/ ORAL	4 DAY	Tremor					

Date:03/01/99ISR Number: 3433686-9Report Type:Periodic  
 Age:47 YR Gender:Female I/FU:I

Company Report #A0079574

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fibromyalgia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3433689-4Report Type:Periodic  
 Age:42 YR Gender:Female I/FU:I

Company Report #A0079576

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							

TWICE PER DAY	Influenza Like Illness					
/ ORAL	Nausea					
	Tachycardia			Fluoxetine		
	Tremor			Hydrochloride		C
	Vomiting			Conjugated Estrogens		C

Date:03/01/99ISR Number: 3433692-4Report Type:Periodic Company Report #A0079577  
 Age:56 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Burning Sensation	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Condition Aggravated					
TWICE PER DAY		Dysgeusia					
/ ORAL		Tobacco Abuse		Prempro			C

Date:03/01/99ISR Number: 3433694-8Report Type:Periodic Company Report #A0081150  
 Age:38 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL		Anxiety					
		Headache					
		Insomnia					
		Nausea					
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433697-3Report Type:Periodic Company Report #A0081151  
 Age:44 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Nausea	Health				
			Professional Other	Nicotine Patch	SS		

Date:03/01/99ISR Number: 3433699-7Report Type:Periodic Company Report #A0081152  
 Age:36 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Flatulence	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433702-4Report Type:Periodic Company Report #A0081161  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL		Toxicologic Test Abnormal	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433705-XReport Type:Periodic Company Report #A0081177  
 Age:35 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Pruritus	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Rash Macular	Professional				
		Swelling	Company Representative				

Date:03/01/99ISR Number: 3433708-5Report Type:Periodic Company Report #A0081193  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ageusia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG SINGLE		Elevated Mood					
DOSE ORAL		Tongue Coated					

Date:03/01/99ISR Number: 3433732-2Report Type:Periodic Company Report #A0081198  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL			Professional Company Representative				

Date:03/01/99ISR Number: 3433734-6Report Type:Periodic Company Report #A0081199  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL			Professional Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433735-8Report Type:Periodic Company Report #A0081200  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL			Professional Company Representative				

Date:03/01/99ISR Number: 3433749-8Report Type:Periodic Company Report #A0081201  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL			Professional Company Representative				

Date:03/01/99ISR Number: 3433763-2Report Type:Periodic Company Report #A0081202  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL			Professional Company Representative				

Date:03/01/99ISR Number: 3433768-1Report Type:Periodic Company Report #A0081203  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Health	Zyban	PS	Glaxo Wellcome Inc	
			Professional Company				

Representative

Date:03/01/99ISR Number: 3433771-1Report Type:Periodic Company Report #A0081319  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Back Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICER							
PER DAY ORAL							

Date:03/01/99ISR Number: 3433776-0Report Type:Periodic Company Report #A0081323  
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
		Insomnia					
		Nervousness		Thyroxine Sodium	C		
				Cortisone	C		
				Testosterone	C		
				Hydrocodone	C		
				Calcium Salt	C		
				Nicotine	C		



Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL			Dizziness Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
					Orphenadrine	C		
Date:03/01/99ISR Number: 3433790-5Report Type:Periodic				Company Report #A0081331				
Age:46 YR Gender:Male I/FU:I								
150 MG TWICE PER DAY ORAL			Accommodation Disorder Anxiety Insomnia Malaise Mental Impairment Tremor	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
					Pravastatin Sodium	C		
Date:03/01/99ISR Number: 3433793-0Report Type:Periodic				Company Report #A0081332				
Age:41 YR Gender:Female I/FU:I								
150 MG TWICE PER DAY ORAL			Drug Interaction Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
					Nicotine (Formulation Unknown)	SS		
TRANSDERMAL		TRANSDERMAL						

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433797-8Report Type:Periodic Company Report #A0081333  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3433799-1Report Type:Periodic Company Report #A0081334  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bronchospasm	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Chest Discomfort					
PER DAY ORAL							

Date:03/01/99ISR Number: 3433801-7Report Type:Periodic Company Report #A0081335  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Urticaria					
PER DAY ORAL							

Date:03/01/99ISR Number: 3433818-2Report Type:Periodic Company Report #A0081336  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							

Date:03/01/99ISR Number: 3433819-4Report Type:Periodic Company Report #A0081337  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL	3	MON					

Glipizide	C
Glucophage	C
Fexofenadine	
Hydrochlorid	C

Date:03/01/99ISR Number: 3433823-6Report Type:Periodic Company Report #A0081338  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nervousness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Tremor					
PER DAY ORAL							

Birth Control	C
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Date:03/01/99ISR Number: 3433826-1Report Type:Periodic Company Report #A0081339  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433829-7Report Type:Periodic  
Age:39 YR Gender: I/FU:I

Company Report #A0081340

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL	2 MON	Libido Decreased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3433837-6Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #A0074417

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Face Oedema Photosensitivity Reaction Rash Erythematous	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Tenorectic Sertraline Hydrochloridge Buspirone Hydrochloride Promethazine Hcl Simastatin	C C C C C C		

Date:03/01/99ISR Number: 3433839-XReport Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0074418

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /TWICE PER DAY/ORAL		Confusional State Dissociation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				No Concurrent Medication Ethanol	C		

Date:03/01/99ISR Number: 3433843-1Report Type:Periodic Company Report #A0074419  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /			Professional				
TWICE PER DAY							
/ORAL	2	WK					

Date:03/01/99ISR Number: 3433847-9Report Type:Periodic Company Report #A0074420  
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Muscular Weakness	Professional				
TWICE PER DAY							
/ORAL							

Simvastatin C  
Multivitamin C  
Enalapril Maleste C

Date:03/01/99ISR Number: 3433850-9Report Type:Periodic Company Report #A0074421  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source
Headache		Consumer
		Health
		Professional

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Other

Dose	Duration	Product	Role	Manufacturer	Route
150 MG / UNK		Zyban	PS	Glaxo Wellcome Inc	ORAL
/ORAL	1 DAY	Nicotine	C		

Date:03/01/99ISR Number: 3433853-4Report Type:Periodic Company Report #A0074423  
 Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3433855-8Report Type:Periodic Company Report #A0074424  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction	Consumer	Zyban	PS	Glaxo Wellcome Inc	
150 MG /				Ranitidine Hydrochloride	C		
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3433857-1Report Type:Periodic Company Report #A0074425  
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Urticaria Consumer Zyban PS Glaxo Wellcome Inc ORAL  
 150 MG /  
 TWICE PER DAY  
 / ORAL  
 Metoprolol Succinate C  
 Nicotinic Acid C

Date:03/01/99ISR Number: 3433859-5Report Type:Periodic Company Report #A0074426  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY /							
ORAL							

Date:03/01/99ISR Number: 3433863-7Report Type:Periodic Company Report #A0074427  
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							
DAY / ORAL		Vitiligo					
				Diazepam	C		
				Co-Trimoxazole	C		
				Zyban	C	Glaxo Wellcome Inc	
				Bethanidine	C		
				Triamcinolone			
				Acetonide	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433867-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0074428

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / UNK		Constipation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
/ ORAL		Dysgeusia					
		Insomnia					
		Vision Blurred					

Date:03/01/99ISR Number: 3433871-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0074429

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / UNK		Drug Effect Decreased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
/ ORAL		Hypersensitivity					

Date:03/01/99ISR Number: 3433874-1Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #A0074430

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /TWICE		Hyperglycaemia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY /ORAL		Vision Blurred	Professional				
				Hydroxyzine Hydrochloride	C		
				Digoxic	C		
				Omeprazole	C		
				Trimethoprim	C		

Date:03/01/99ISR Number: 3433877-7Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0074431

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3433880-7Report Type:Periodic Company Report #A0074432  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY /ORAL							

Date:03/01/99ISR Number: 3433882-0Report Type:Periodic Company Report #A0074481  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG / TWICE							
PER DAY /							
ORAL	2	WK					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3433883-2Report Type:Periodic Company Report #A0074482  
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER							
DAY/ ORAL	2	WK					

Date:03/01/99ISR Number: 3433886-8Report Type:Periodic Company Report #A0074547  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / UNK							
/ ORAL							

Date:03/01/99ISR Number: 3433889-3Report Type:Periodic Company Report #A0074641  
 Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG / TWICE		Dermatitis					
PER DAY /		Gastritis					
ORAL		Hypersensitivity					
		Pain					
		Pruritus					
		Rash Erythematous					
		Swelling					

Date:03/01/99ISR Number: 3433951-5Report Type:Periodic Company Report #A0074713  
 Age:41 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Constipation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Haematochezia					
/ ORAL		Headache					
		Insomnia					
		Nausea					
		Tremor					

Date:03/01/99ISR Number: 3434001-7Report Type:Periodic Company Report #A0077688  
Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ ORAL		Drug Ineffective	Health Professional	Zyban Tablet - Zyban	PS		ORAL

Date:03/01/99ISR Number: 3434003-0Report Type:Periodic Company Report #A0077689  
Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ ORAL		Nausea	Consumer	Zyban Tablet - Zyban	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3434006-6Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #A0077690

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ PER							
DAY/ ORAL							

Thyroxine Sodium C

Date:03/01/99ISR Number: 3434009-1Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0077691

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Oedema Peripheral					
PER DAY/ ORAL		Urticaria					

Date:03/01/99ISR Number: 3434012-1Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #A0077692

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Pruritus					
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3434016-9Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #A0077693

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Atorvastatin Calcium C  
 Calcium Carbonate C  
 Centrum C  
 Loratadine C

Date:03/01/99ISR Number: 3434018-2Report Type:Periodic Company Report #A0077694  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
75 MG / SEE		Pain					
TEXT/ ORAL		Pruritus		Darvocet-N	C		
				Estrogen	C		
				Nortriptyline	C		
				Zolpidem Tartrate	C		
				Amlodipine	C		
				Cisapride	C		
				Metolazone	C		
				Multivitamin	C		
				Calcium Salt	C		
				Mirtazapine	C		
				Clonazepam	C		
				Simvastatin	C		
				Aspirin	C		
				Diltiazem			
				Hydrochloride	C		
				Cyproheptadine	C		
				Desocort	C		
				Decongestant	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lansoprazole C

Date:03/01/99ISR Number: 3434021-2Report Type:Periodic Company Report #A0077695  
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Dizziness	Professional				
DAY/ ORAL		Hypoaesthesia		Oxaprozin	C		
		Nausea					
		Sedation					

Date:03/01/99ISR Number: 3434025-XReport Type:Periodic Company Report #A0077696  
 Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /TWICE		Urticaria					
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3434027-3Report Type:Periodic Company Report #A0077697  
 Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gastrointestinal Disorder	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /TWICE		Vomiting					
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3434030-3Report Type:Periodic Company Report #A0077698  
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /TWICE							
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3434033-9Report Type:Periodic Company Report #A0077709  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Health	Zyban Tablet - Zyban	PS		ORAL
ORAL							
			Professional Company Representative				

Date:03/01/99ISR Number: 3434036-4Report Type:Periodic Company Report #A0077837  
 Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
/ SEE TEXT/							
		Ventricular Extrasystoles					
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3434040-6Report Type:Periodic  
Age:59 YR Gender:Male I/FU:I

Company Report #A0077847

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		3 WK	Convulsion Head Injury	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:03/01/99ISR Number: 3434041-8Report Type:Periodic  
Age:33 YR Gender:Male I/FU:I

Company Report #A0077869

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150 MG / PER		Chest Pain Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
	DAY / ORAL		Insomnia Tachycardia		Multivitamin Chromium Picolinate	C C		

Date:03/01/99ISR Number: 3434044-3Report Type:Periodic  
Age:55 YR Gender:Male I/FU:I

Company Report #A0077871

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150 MG/ PER		Amnesia Chest Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL
	DAY / ORAL		Confusional State Disorientation Dizziness Eye Disorder Headache Nausea		Nicotine	C		

Date:03/01/99ISR Number: 3434047-9Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0077952

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphagia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Hyperhidrosis					
TWICE PER DAY		Insomnia					
/ ORAL		Tachycardia		Hydroxyzine	C		

Date:03/01/99ISR Number: 3434049-2Report Type:Periodic Company Report #A0077953  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Major Depression					
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3434051-0Report Type:Periodic Company Report #A0077954  
 Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Urinary Retention	Professional				
TWICE PER							
DAY/ ORAL							
				Fluoxetine			
				Hydrochloride	C		
				Cyanocobalamin	C		
				Salbutamol Sulphate	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Salbutamol Sulphate C  
 Salmeterol Xinafoate C  
 Multivitamin C

Date:03/01/99ISR Number: 3434055-8Report Type:Periodic Company Report #A0077955  
 Age:38 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Headache	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Myalgia					
/ ORAL				Tri-Levlen	C		

Date:03/01/99ISR Number: 3434063-7Report Type:Periodic Company Report #A0081370  
 Age:46 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Disturbance In Attention	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY /							
ORAL							

Date:03/01/99ISR Number: 3434064-9Report Type:Periodic Company Report #A0077956  
 Age:43 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Confusional State	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY /		Insomnia					
ORAL		Nervousness					
				Nafarelin Acetate	C		

Date:03/01/99ISR Number: 3434065-0Report Type:Periodic Company Report #A0077957  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							
				Estrogen	C		
				Medroxyprogesterone	C		

Date:03/01/99ISR Number: 3434066-2Report Type:Periodic Company Report #A0077958  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL		Urticaria					
				Nicotine	C		

Date:03/01/99ISR Number: 3434067-4Report Type:Periodic Company Report #A0077959  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Depressed Mood	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY /							
ORAL							

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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3434068-6Report Type:Periodic  
Age:31 YR Gender:Male I/FU:I

Company Report #A0077960

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY / ORAL		Anxiety Irritability	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3434069-8Report Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #A0077961

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL		Dizziness Tremor	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Amitriptyline	C		
				Aspirin	C		
				Multivitamin	C		

Date:03/01/99ISR Number: 3434070-4Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #A0077962

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ORAL		Aggression Disorientation Personality Change	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Sertraline Hydrochloride	C		
				Olanzapine	C		
				Propranolol Hydrochloride	C		

Date:03/01/99ISR Number: 3434071-6Report Type:Periodic Company Report #A0077963  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Indifference					
DAILY/ ORAL		Insomnia					

Date:03/01/99ISR Number: 3434072-8Report Type:Periodic Company Report #A0077974  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
DAILY/ ORAL							

Date:03/01/99ISR Number: 3434073-XReport Type:Periodic Company Report #A0078008  
Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disturbance In Attention	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Dizziness					
PER DAY/ ORAL		Hallucination					
		Tremor					
		Visual Disturbance					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3434074-1Report Type:Periodic Company Report #A0078014  
 Age:37 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ YWICE PER DAY/ ORAL		Burning Sensation Drug Ineffective Joint Swelling Lymphocytosis Nausea Pain Paraesthesia Rash Pruritic Red Blood Cell Sedimentation Rate Increased White Blood Cell Count Increased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3434075-3Report Type:Periodic Company Report #A0078054  
 Age:45 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY / ORAL		Pruritus Urticaria	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3434076-5Report Type:Periodic Company Report #A0078056  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ ORAL		Nightmare	Health Professional Company Representative	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3434077-7Report Type:Periodic Company Report #A0078182  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ TWICE		Abdominal Distension	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL		Abdominal Pain					
		Abdominal Pain Upper					

Date:03/01/99ISR Number: 3434078-9Report Type:Periodic Company Report #A0078183  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ TWICE		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY /							
ORAL							

Date:03/01/99ISR Number: 3434079-0Report Type:Periodic Company Report #A0078184  
Age:32 YR Gender:Female I/FU:I

Outcome	PT
	Constipation
	Dizziness

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Freedom Of Information (FOI) Report

		Headache Tachycardia Vision Blurred	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER DAY/ ORAL							

Date:03/01/99ISR Number: 3434080-7Report Type:Periodic Company Report #A0078185  
Age:26 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE PER DAY/ ORAL		Insomnia					
		Therapeutic Response Unexpected		Percodan	C		

Date:03/01/99ISR Number: 3434081-9Report Type:Periodic Company Report #A0078187  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Conversion Disorder	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE DAILY/ ORAL		Crying					
		Dysgeusia Fear		Paracetamol Ibuprofen	C C		

Date:03/01/99ISR Number: 3434082-0Report Type:Periodic Company Report #A0078188  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							

Psoriasis

PER DAY /

ORAL

Estrogen

C

Date:03/01/99ISR Number: 3434083-2Report Type:Periodic

Company Report #A0078189

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Choking	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL		Retching					

Date:03/01/99ISR Number: 3434084-4Report Type:Periodic

Company Report #A0078190

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Dizziness					
PER DAY/ORAL		Fatigue					
		Feeling Abnormal					
		Nausea					
		Vertigo					
		Vomiting					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3434085-6Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #A0078191

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
PER DAY/ORAL							
				Semisodium Valproate	C		
				Clonazepam	C		
				Zolpidem Tartrate	C		

Date:03/01/99ISR Number: 3434086-8Report Type:Periodic  
 Age:40 YR Gender:Male I/FU:I

Company Report #A0078192

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG / TWICE		Feeling Abnormal					
PER DAY/ORAL		Palpitations					
		Syncope					
		Tachycardia					

Date:03/01/99ISR Number: 3434087-XReport Type:Periodic  
 Age:51 YR Gender:Female I/FU:I

Company Report #A0078193

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150		Vomiting					
N=MG/TWICE							
PER DAY/ORAL							
				Amlodipine	C		
				Atorvastatin Calcium	C		
				Nicotinic Acid	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Dermatitis Exfoliative	Professional				
PER DAY/ ORAL		Face Oedema		Waarfarin Sodium	C		
				Salbutamol Sulphate	C		
				Beclomethasone			
				Dipropion	C		
				Ipratropium Bromide	C		
				Oxygen	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Eczema					
TWICE PER DAY		Insomnia					
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3434090-XReport Type:Periodic Company Report #A0078196  
 Age:40 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL 3 WK							

Date:03/01/99ISR Number: 3434091-1Report Type:Periodic Company Report #A0078270  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Unevaluable Event	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3434092-3Report Type:Periodic Company Report #A0078413  
 Age:53 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /TWICE		Fluid Retention	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY /							
ORAL							

Date:03/01/99ISR Number: 3434093-5Report Type:Periodic Company Report #A0078414  
 Age:40 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /TWICE		Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3434094-7Report Type:Periodic Company Report #A0078415  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Disorientation Dizziness Dry Mouth Hyperacusis Mental Disorder Sedation Tachycardia Tremor	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3434095-9Report Type:Periodic Company Report #A0078416  
Age:47 YR Gender:Female I/FU:I

Outcome	PT
	Arthralgia Cough Dermatitis Difficulty In Walking Dry Mouth Headache Hypoaesthesia Logorrhoea Nausea Oedema Peripheral Paraesthesia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pruritus

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG /TWICE PER DAY / ORAL		Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3434096-0Report Type:Periodic Company Report #A0078417  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /TWICE PER DAY / ORAL		Condition Aggravated Headache Nervousness Smoker	Consumer	Zyban Fexofenadine Hydrochlorid Multivitamin	PS C C	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3434097-2Report Type:Periodic Company Report #A0078418  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Dermatitis Eyelid Oedema Urticaria	Consumer	Zyban Birth Control	PS C	Glaxo Wellcome Inc	ORAL

Date:03/01/99ISR Number: 3434098-4Report Type:Periodic Company Report #A0078419  
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /PER							
DAY / ORAL				Maxzide	C		
				Lisinopril	C		

Date:03/01/99ISR Number: 3434099-6Report Type:Periodic Company Report #A0078420  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							
DAY / ORAL							

Date:03/01/99ISR Number: 3434100-XReport Type:Periodic Company Report #A0078421  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							
DAY / ORAL		Confusional State					
		Disturbance In Attention					
		Influenza Like Illness					
		Irritability					
		Myalgia					
		Pain					
		Skin Irritation					



150 MG /  
TWICE PER DAY  
/ ORAL

Dizziness  
Nausea  
Tinnitus

Consumer

Zyban Tablet - Zyban PS

ORAL

Date:03/01/99ISR Number: 3434407-6Report Type:Periodic Company Report #A0078952  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
1 TABLET /							
ORAL							

Diphenhydramine Hcl C  
Esgic-Plus C

Date:03/01/99ISR Number: 3434408-8Report Type:Periodic Company Report #A0078953  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Excitability	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY		Mood Swings					
/ ORAL							

Birth Control C  
Multivitamin C



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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3434411-8Report Type:Periodic  
 Age:28 YR Gender:Female I/FU:I

Company Report #A0078954

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3434413-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0078955

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL							

Date:03/01/99ISR Number: 3434414-3Report Type:Periodic  
 Age:32 YR Gender:Female I/FU:I

Company Report #A0078956

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL							

Date:03/01/99ISR Number: 3434418-0Report Type:Periodic  
 Age:56 YR Gender:Female I/FU:I

Company Report #A0078957

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Psychomotor Hyperactivity	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Clarithromycin	C
Vitamin E	C

Librium C  
Doxylamine Succinate C  
Pyridoxine  
Hydrochloride C

Date:03/01/99ISR Number: 3434421-0Report Type:Periodic Company Report #A0078958  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Chest Discomfort					
TWICE PER		Dyspepsia					
DAY/ ORAL				Doxylamine Succinate	C		

Date:03/01/99ISR Number: 3434422-2Report Type:Periodic Company Report #A0078960  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthma	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL			Professional Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3434425-8Report Type:Periodic  
 Age:55 YR Gender:Female I/FU:I

Company Report #A0078963

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dysgeusia			Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL							
				Verapamil	C		
				Atenolol	C		
				Medrol	C		
				Potassium Salt	C		
				Conjugates Estrogens	C		
				Triamter.			
				+Hclthiazide	C		
				Pravastatin Sodium	C		

Date:03/01/99ISR Number: 3434427-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0078967

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Anxiety			Consumer	Zyban Tablet-Zyban	PS		ORAL
150MG/ THREE							
TIMES PER							
DAY/ORAL							
		Insomnia					
		Nausea					
		Overdose					
		Tremor					

Date:03/01/99ISR Number: 3434429-5Report Type:Periodic  
 Age:45 YR Gender:Female I/FU:I

Company Report #A0078968

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Anxiety			Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY/ORAL							
		Tremor					
				Oestradiol	C		
				Simvastatin	C		

Date:03/01/99ISR Number: 3434430-1Report Type:Periodic Company Report #A0079588  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Consumer	Zyban	PS		ORAL
150 MG /TWICE		Drug Ineffective					
PER DAY /		Insomnia					
ORAL				Pravastatin Sodium	C		

Date:03/01/99ISR Number: 3434432-5Report Type:Periodic Company Report #A0079589  
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS		ORAL
150 MG /		Feeling Abnormal					
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3434434-9Report Type:Periodic Company Report #A0079590  
Age:67 YR Gender:Male I/FU:I

Outcome PT  
Constipation  
Dizziness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Insomnia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY / ORAL		Consumer	Zyban	PS		ORAL
			Ibuprofen	C		

Date:03/01/99ISR Number: 3434436-2Report Type:Periodic Company Report #A0079591  
Age:56 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150 MG / TWICE PER DAY / ORAL	14 DAY	Nausea Retching	Consumer	Zyban	PS		ORAL
					Allopurinol Terazosin Hydrochloride Chlorthalidone	C C C		

Date:03/01/99ISR Number: 3434438-6Report Type:Periodic Company Report #A0079592  
Age: Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150 MG / ORAL	7 DAY	Pollakiuria	Consumer	Zyban	PS		ORAL
					Amoxicillin	C		

Date:03/01/99ISR Number: 3434439-8Report Type:Periodic Company Report #A0079617  
Age:30 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150 MG / PER		Chest Discomfort	Consumer	Zyban	PS		ORAL
	DAY / ORAL		Dizziness					
			Dyspnoea		Birth Control	C		
			Feeling Abnormal					
			Hyperventilation					
			Nausea					
			Pruritus					
			Syncope					
			Tachycardia					
			Tremor					
			Urticaria					

Date:03/01/99ISR Number: 3434440-4Report Type:Periodic Company Report #A0079764

Age: Gender:Unknown I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		3 DAY	Oedema Peripheral	Health	Zyban Tablet - Zyban	PS		ORAL
			Pruritus	Professional	Wellbutrin			
			Urticaria		Tablet-Controlled			
150 MG / ORAL		3 WK			Release	SS		ORAL

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Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3434441-6Report Type:Periodic  
Age:73 YR Gender:Male I/FU:I

Company Report #A0079951

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /TWICE		Anxiety					
PER DAY /		Headache					
ORAL		Nausea		Cortisone	C		
		Nervousness		Prednisone	C		

Date:03/01/99ISR Number: 3434442-8Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #A0079953

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /TWICE		Insomnia					
PER DAY /							
ORAL				Tylenol W/ Codeine			
				No. 4	C		
				Cyclobenzaprine Hcl	C		
				Clorazepic Acid	C		

Date:03/01/99ISR Number: 3434443-XReport Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #A0079954

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3434444-1Report Type:Periodic Company Report #A0079955  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Hypertension					
PER DAY /		Pruritus					
ORAL		Vomiting					

Date:03/01/99ISR Number: 3434445-3Report Type:Periodic Company Report #A0079956  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL 1 WK		Crying					
		Dermatitis					

Date:03/01/99ISR Number: 3434447-7Report Type:Periodic Company Report #A0079957  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dissociation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dyspraxia					
TWICE PER DAY		Nervousness					
/ ORAL		Sedation		Nortriptyline Hcl	C		
				Diazepam	C		
				Carisoprodol	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Oxycodone  
Hydrochloride C

Date:03/01/99ISR Number: 3434448-9Report Type:Periodic Company Report #A0079958  
Age:29 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Abnormal Behaviour	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Fatigue Feeling Abnormal		Dilantin	C		

Date:03/01/99ISR Number: 3434454-4Report Type:Periodic Company Report #A0080013  
Age:49 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY /ORAL		Chills	Health	Zyban Tablet - Zyban	PS		ORAL
		Convulsion	Professional				
		Influenza Like Illness Lacrimation Increased Pyrexia Rhinorrhoea Sneezing Tremor					

Date:03/01/99ISR Number: 3434457-XReport Type:Periodic Company Report #A0080099  
Age:20 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Dyspnoea	Health	Zyban Tablet-Zyban	PS		ORAL
			Professional				

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Paraesthesia					
DAY / ORAL				Prempro	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL							

Isosorbide  
Mononitrate C  
Conjugated Estrogens C  
Atorvastatin Calcium C  
Ranitidine  
Hydrochloride C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3434466-0Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #A0080052

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Paraesthesia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Tachycardia					
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3434467-2Report Type:Periodic  
Age:72 YR Gender:Female I/FU:I

Company Report #A0080053

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Dizziness					
DAY / ORAL		Nausea		Amlodipine	C		
				Diazepam	C		
				Insulin	C		
				Guanfacine			
				Hydrochloride	C		
				Glipizide	C		

Date:03/01/99ISR Number: 3434470-2Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0080054

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY/ORAL				Glipizide	C		

Date:03/01/99ISR Number: 3434472-6Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0080055

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Consumer	Zyban Tablet-Zyban	PS		ORAL
150M MG/PER		Hypoglycaemia					
DAY/ORAL				Insulin	C		
				Captopril	C		
				Frusemide	C		
				Digoxin	C		
				Isotretinoin	C		
				Warfarin Sodium	C		
				Amlodipine	C		
				Isosorbide	C		

Date:03/01/99ISR Number: 3434474-XReport Type:Periodic Company Report #A0080056  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TIWCE							
PER DAY/							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3434476-3Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0080057

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /TWICE		Dermatitis	Consumer	Zyban Tablet-Zyban	PS		ORAL
PER DAY/ ORAL		Pruritus					
		Urticaria					

Date:03/01/99ISR Number: 3434478-7Report Type:Periodic  
Age:25 YR Gender:Female I/FU:I

Company Report #A0080058

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ PER		Aggression	Consumer	Zyban Tablet-Zyban	PS		ORAL
DAY/ ORAL		Agitation					
		Disorientation					
		Headache					
		Hyperhidrosis					

Date:03/01/99ISR Number: 3434479-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0080059

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ ORAL		Hypersensitivity	Consumer	Zyban Tablet- Zyban	PS		ORAL
				Famotidine	I		

Date:03/01/99ISR Number: 3434482-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0080060

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ ORAL		Dermatitis	Consumer	Zyban Tablet-Zyban	PS		ORAL
		Pruritus					

Date:03/01/99ISR Number: 3434485-4Report Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #A0080061

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ORAL		Anorexia					

Blood Pressure Increased  
Chest Pain  
Depression  
Disturbance In Attention  
Heart Rate Increased  
Insomnia  
Mental Disorder  
Paranoia  
Tremor

Date:03/01/99ISR Number: 3434490-8Report Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #A0080062

Outcome	PT
	Dry Mouth
	Dry Skin
	Hypersensitivity
	Insomnia
	Lichen Planus

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Freedom Of Information (FOI) Report

		Milia				
		Pharyngolaryngeal Pain				
		Pruritus	Report Source	Product	Role	Manufacturer
Dose	Duration		Consumer	Zyban Tablet-Zyban	PS	Route
150 MG /						ORAL
TWICE PER						
DAY/ ORAL				Multivitamin	C	

Date:03/01/99ISR Number: 3434492-1Report Type:Periodic Company Report #A0080064  
 Age:69 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Arthralgia	Consumer	Zyban Tablet-Zyban	PS		ORAL
Dose		Dry Mouth					
150 MG/ TWICE		Insomnia					
PER DAY/ORAL							

Date:03/01/99ISR Number: 3434494-5Report Type:Periodic Company Report #A0080065  
 Age:58 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Drug Ineffective	Consumer	Zyban Tablet-Zyban	PS		ORAL
Dose							
150 MG/ TWICE				Natural Medication	C		
PER DAY/ORAL							

Date:03/01/99ISR Number: 3434497-0Report Type:Periodic Company Report #A0080066  
 Age:56 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Abnormal Dreams	Consumer	Zyban Tablet-Zyban	PS		ORAL
Dose							
150 MG/ TWICE							

Dermatitis

PER DAY/ ORAL

Date:03/01/99ISR Number: 3434499-4Report Type:Periodic Company Report #A0080079  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG /							
TIWCE PER							
DAY/ ORAL							

Date:03/01/99ISR Number: 3434502-1Report Type:Periodic Company Report #A0080093  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ ORAL				Thiothixene	C		
				Benztropine Mesylate	C		
				Clonidine			
				Hydrochloride	C		

Date:03/01/99ISR Number: 3434505-7Report Type:Periodic Company Report #A0080094  
Age:31 YR Gender:Female I/FU:I

Outcome	PT
	Burning Sensation
	Pruritus



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Rash Erythematous

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Consumer	Zyban Tablet-Zyban	PS		ORAL
PER DAY/ORAL			Naproxen Sodium	C		

Date:03/01/99ISR Number: 3434507-0Report Type:Periodic Company Report #A0080095  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY/ORAL							

Date:03/01/99ISR Number: 3434509-4Report Type:Periodic Company Report #A0080096  
 Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Health	Zyban Tablet-Zyban	PS		ORAL
150 MG/ ORAL	2 WK		Professional				

Date:03/01/99ISR Number: 3434512-4Report Type:Periodic Company Report #A0080097  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Suicidal Ideation	Health	Zyban Tablet-Zyban	PS		ORAL
150 MG/ ORAL			Professional				

Date:03/01/99ISR Number: 3434514-8Report Type:Periodic Company Report #A0080098  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TWICE		Bruxism					
PER DAY/ORAL		Dermatitis		Birth Control	C		
		Headache					
		Pruritus					
		Toothache					
		Urticaria					

Date:03/01/99ISR Number: 3434521-5Report Type:Periodic Company Report #A0080106  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Blood Pressure Increased					
TWICE PER DAY		Cold Sweat					
/ ORAL		Dermatitis					
		Dizziness					
		Headache					
		Insomnia					
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3434523-9Report Type:Periodic  
 Age:62 YR Gender:Female I/FU:I

Company Report #A0080108

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / VARIABLE DOSE / ORAL		Condition Aggravated Insomnia Ulcer	Consumer	Zyban Tablet - Zyban  Ranitidine Hydrochloride	PS  C		ORAL

Date:03/01/99ISR Number: 3434524-0Report Type:Periodic  
 Age:77 YR Gender:Female I/FU:I

Company Report #A0080171

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Insomnia	Consumer	Zyban Tablet - Zyban  Atenolol Thyroxine Sodium Aspirin Conjugated Estrogens Medroxyprogesterone Ace. Trazodone	PS  C C C C C C		ORAL

Date:03/01/99ISR Number: 3434528-8Report Type:Periodic  
 Age:29 YR Gender:Female I/FU:I

Company Report #A0080172

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY		Tinnitus	Consumer	Zyban Tablet - Zyban	PS		ORAL

/ ORAL

Date:03/01/99ISR Number: 3434530-6Report Type:Periodic Company Report #A0080173  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Nausea					
TWICE PER DAY							

/ ORAL

Fexofenadine  
Hydrochlorid C  
Desogen C

Date:03/01/99ISR Number: 3434532-XReport Type:Periodic Company Report #A0080174  
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eating Disorder	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 /TWICE		Hyperhidrosis					
PER DAY /		Nausea					
ORAL		Throat Tightness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3434535-5Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0080175

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Drug Ineffective		Aspirin	C		
				Cetirizine			
				Hydrochloride	C		
				Lisinopril	C		
				Diltiazem			
				Hydrochloride	C		
				Pravastatin Sodium	C		

Date:03/01/99ISR Number: 3434539-2Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0080326

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL			Professional				

Date:03/01/99ISR Number: 3434551-3Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0080329

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY		Pollakiuria					
/ ORAL				Conjugated Estrogens	C		
				Gemfibrozil	C		
				Famotidine	C		

Date:03/01/99ISR Number: 3434559-8Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0080346

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Chest Pain		Zyban Tablet - Zyban	PS		ORAL
		Rash Pruritic					
TWICE PER DAY							
/ ORAL							

Date:03/01/99ISR Number: 3434562-8Report Type:Periodic Company Report #A0081341  
 Age:55 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Pyrexia	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL							

Vicodin	C
Ciprofloxacin Hcl	C
Prempro	C
Hydrochlorothiazide	C

Date:03/01/99ISR Number: 3434568-9Report Type:Periodic Company Report #A0081342  
 Age: Gender:Female I/FU:I

Outcome	PT
	Dysphagia
	Glossitis

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tongue Disorder

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL	13 DAY	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:03/01/99ISR Number: 3434573-2Report Type:Periodic Company Report #A0081343  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Dry Mouth					
/ ORAL		Excitability					
		Flushing		Tramadol Hydrochloride	C		

Date:03/01/99ISR Number: 3434575-6Report Type:Periodic Company Report #A0081344  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER		Confusional State	Consumer	Zyban Tablet - Zyban	PS		ORAL
DAY / ORAL		Dermatitis					
		Diarrhoea					
		Dizziness					
		Dysarthria					
		Insomnia					
		Nausea					
		Tremor					

Date:03/01/99ISR Number: 3434579-3Report Type:Periodic Company Report #A0081345  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Monarthritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Levlen	C		

Date:03/01/99ISR Number: 3434581-1Report Type:Periodic Company Report #A0081346  
 Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Zyban Tablet - Zyban	PS		ORAL
150 MG /		Urticaria					
TWICE PER DAY							
/ ORAL				Enalapril Maleate	C		
				Atorvastatin Calcium	C		

Date:03/01/99ISR Number: 3434584-7Report Type:Periodic Company Report #A0081347  
 Age: Gender: I/FU:I

Outcome PT  
 Abnormal Dreams  
 Confusional State  
 Constipation  
 Dry Mouth  
 Eye Disorder



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Hallucination, Auditory Ill-Defined Disorder Palpitations	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL	8 DAY	Tremor	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:03/01/99ISR Number: 3434589-6Report Type:Periodic Company Report #A0081348  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY				Thyroxine Sodium	C		
/ ORAL							

Date:03/01/99ISR Number: 3434590-2Report Type:Periodic Company Report #A0081349  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY				Propranolol Hydrochloride	C		
/ ORAL							

Date:03/01/99ISR Number: 3434592-6Report Type:Periodic Company Report #A0081369  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL							

Date:03/01/99ISR Number: 3434602-6Report Type:Periodic Company Report #A0080531  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Blood Pressure Increased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Chest Pain					
PER DAY/ ORAL							
				Moexipril			
				Hydrochloride	C		
				Clonidine	C		
				Percocet	C		

Date:03/01/99ISR Number: 3434605-1Report Type:Periodic Company Report #A0080532  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Flatulence					
PER DAY/ ORAL		Insomnia					
				Glibenclamide	C		
				Verapamil			
				Hydrochloride	C		
				Conjugated Estrogens	C		
				Triamterene	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3434606-3Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #A0080667

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
PER DAY/ORAL							

Conjugated Estrogens C

Date:03/01/99ISR Number: 3434607-5Report Type:Periodic  
Age:63 YR Gender:Male I/FU:I

Company Report #A0080690

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
PER DAY/ORAL		Oral Mucosal Blistering					

Simvastatin C

Date:03/01/99ISR Number: 3434608-7Report Type:Periodic  
Age:67 YR Gender:Male I/FU:I

Company Report #A0080691

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/TWICE							
PER DAY/ORAL		Eating Disorder					

Quinapril Hydrochloride C

Date:03/01/99ISR Number: 3434611-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0080693

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hyperglycaemia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/UNKNOWN							

/ ORAL

Date:03/01/99ISR Number: 3435069-4Report Type:Periodic Company Report #A0080025  
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity	Consumer	Zyban Tablet -Zyban	PS		ORAL
150 MG /TWICE		Urticaria					
PER DAY /							
ORAL							

Date:03/01/99ISR Number: 3436404-3Report Type:Periodic Company Report #A0080694  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/TWICE		Mania					
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3436406-7Report Type:Periodic Company Report #A0080695  
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/TWICE		Dry Mouth					
PER DAY/ ORAL		Insomnia		Alprazolam	C		
		Tobacco Abuse					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3436408-0Report Type:Periodic Company Report #A0080696  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Paraesthesia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/TWICE							
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3436411-0Report Type:Periodic Company Report #A0078820  
 Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/TWICE							
PER DAY/ ORAL							

Frusemide	C
Warfarin Sodium	C
Human Insulin	C
Nitroglycerin	C
Diazepam	C
Fluoxetine	
Hydrochloride	C
Thyroxine Sodium	C
Meclozine	
Hydrochloride	C
Metoprolol Tartrate	C
Isosorbide	
Mononitrate	C
Tolterodine	C

Date:03/01/99ISR Number: 3436415-8Report Type:Periodic Company Report #A0078821  
 Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/TWICE							
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3436418-3Report Type:Periodic Company Report #A0078822  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/TWICE							
PER DAY/ ORAL							

Date:03/01/99ISR Number: 3436421-3Report Type:Periodic Company Report #A0078823  
Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/TWICE							
PER DAY/ ORAL							
				Birth Control	C		

Date:03/01/99ISR Number: 3436425-0Report Type:Periodic Company Report #A0078824  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/TWICE							
PER DAY/ ORAL							
				No Concurrent			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Medication C

Date:03/01/99ISR Number: 3436427-4Report Type:Periodic Company Report #A0078825  
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nervousness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/TWICE							
PER DAY/	ORAL						

Date:03/01/99ISR Number: 3436430-4Report Type:Periodic Company Report #A0078826  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/TWICE		Disturbance In Attention					
PER DAY/	ORAL	Insomnia					

Date:03/01/99ISR Number: 3436433-XReport Type:Periodic Company Report #A0078827  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/TWICE							
PER DAY/	ORAL						

Date:03/01/99ISR Number: 3436437-7Report Type:Periodic Company Report #A0078828  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/TWICE							

Joint Swelling

PER DAY/ ORAL

Date:03/01/99ISR Number: 3436440-7Report Type:Periodic Company Report #A078829

Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/TWICE		Disorientation					
PER DAY/ ORAL		Dyspnoea		Medroxyprogesterone	C		
		Hyperhidrosis		Ace			
		Insomnia					

Date:03/01/99ISR Number: 3436444-4Report Type:Periodic Company Report #A0080373

Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disturbance In Attention	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG PER		Irritability					
DAY ORAL							

Date:03/01/99ISR Number: 3436447-XReport Type:Periodic Company Report #A0080374

Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150MG TWICE		Malaise					
PER DAY ORAL		Tremor		Enalapril Maleate	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Diltiazem  
 Hydrochloride C  
 Amoxicillin  
 Trihydrate C  
 Sildenafil Citrate C

Date:03/01/99ISR Number: 3436451-1Report Type:Periodic Company Report #A0080375  
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE							
PER DAY ORAL				Omeprazole	C		

Date:03/01/99ISR Number: 3436456-0Report Type:Periodic Company Report #A0080376  
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3436458-4Report Type:Periodic Company Report #A0080377  
 Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE							
PER DAY ORAL				Amoxicillin	C		
				Atenolol	C		

Date:03/01/99ISR Number: 3436463-8Report Type:Periodic Company Report #A0080378  
Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE		Pruritus					
PER DAY ORAL							

Date:03/01/99ISR Number: 3436469-9Report Type:Periodic Company Report #A0080379  
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE		Bone Pain					
PER DAY ORAL		Dyspraxia		Diltiazem			
		Feeling Abnormal		Hydrochloride	C		
		Hyperhidrosis		Pravastatin Sodium	C		
		Insomnia		Aspirin	C		

Date:03/01/99ISR Number: 3436471-7Report Type:Periodic Company Report #A0080405  
Age:49 YR Gender:Female I/FU:I

Outcome	PT
	Diplopia
	Disorientation
	Dizziness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Headache Hypoaesthesia Vision Blurred	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Zyban Tablet - Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL				Thyroxine Sodium	C		

Date:03/01/99ISR Number: 3436474-2Report Type:Periodic Company Report #A0080406  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE							
PER DAY ORAL							

Date:03/01/99ISR Number: 3436480-8Report Type:Periodic Company Report #A0080411  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Consumer	Zyban Tablet - Zyban	PS		ORAL
Other							
150 MG ORAL							

Date:03/01/99ISR Number: 3436483-3Report Type:Periodic Company Report #A0080413  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decreased Appetite	Consumer	Zyban Tablet - Zyban	PS		ORAL
Hospitalization -		Disturbance In Attention					
150 MG TWICE							
Initial or Prolonged							
PER DAY ORAL		Dysgeusia Hallucination Headache					

Insomnia

Date:03/01/99ISR Number: 3436485-7Report Type:Periodic Company Report #A0080417  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE		Insomnia					
PER DAY ORAL		Restlessness Suicidal Ideation		Hydroxyzine Hydrochloride	C		

Date:03/01/99ISR Number: 3436487-0Report Type:Periodic Company Report #A0080422  
 Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE		Fatigue					
PER DAY ORAL				Aspirin	C		

Date:03/01/99ISR Number: 3436492-4Report Type:Periodic Company Report #A0080423  
 Age:55 YR Gender:Female I/FU:I

Outcome	PT
	Anxiety Disturbance In Attention

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nausea

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Consumer	Zyban Tablet - Zyban	PS		ORAL
PER DAY ORAL	3 DAY					

Date:03/01/99ISR Number: 3436495-XReport Type:Periodic Company Report #A0080424  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Urticaria	Consumer	Zyban Tablet - Zyban	PS		ORAL
PER DAY ORAL							

Date:03/01/99ISR Number: 3436498-5Report Type:Periodic Company Report #A0080442  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Headache	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Photopsia		Ibuprofen	C		

Date:03/01/99ISR Number: 3436502-4Report Type:Periodic Company Report #A0080444  
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Ill-Defined Disorder	Consumer	Zyban Tablet - Zyban	PS		ORAL
PER DAY ORAL		Nausea		Nifedipine	C		
				Thyroxine Sodium	C		

Date:03/01/99ISR Number: 3436504-8Report Type:Periodic Company Report #A0080446  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ill-Defined Disorder	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG ORAL		Nausea					

Date:03/01/99ISR Number: 3436509-7Report Type:Periodic Company Report #A0080529  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG ORAL							

Date:03/01/99ISR Number: 3436513-9Report Type:Periodic Company Report #A0080530  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG TWICE							
PER DAY ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/99ISR Number: 3443034-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0065843

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
SUBCUTANEOUS	AS	Drug Ineffective	Consumer	Imitrex Injection	PS		
REQUIRED/SUBC		Drug Interaction					
UTANEOUS	4 YR			Wellbutrin Tablet-Controlled Release	SS		ORAL
150 MG/TWICE							
PER DAY/ORAL							

Date:03/02/99ISR Number: 3210328-9Report Type:Expedited (15-DaCompany Report #A0079255  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150 MG TWICE		Arthralgia	Health	Zyban Tablet - Zyban	PS		ORAL
Initial or Prolonged PER DAY ORAL		Pain In Extremity	Professional				
Other				Buspirone Hydrochloride Ibuprofen	C C		

Date:03/02/99ISR Number: 3210330-7Report Type:Expedited (15-DaCompany Report #A0082565  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150 MG TWICE		Constipation	Foreign	Zyban Tablet - Zyban	PS		ORAL
Initial or Prolonged PER DAY ORAL		Faecaloma	Consumer				
				Antibiotics	C		

Date:03/02/99ISR Number: 3210332-0Report Type:Expedited (15-DaCompany Report #A0082079  
Age:69 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG PER Initial or Prolonged DAY ORAL		Balance Disorder Confusional State	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Feeling Abnormal Nausea Vision Blurred		Vitamin E Calcium Salt	C C		

Date:03/02/99ISR Number: 3433775-9Report Type:Periodic Company Report #A0081320  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Diphenhydramine Hcl	C		

Date:03/04/99ISR Number: 3212945-9Report Type:Expedited (15-DaCompany Report #A0082408  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG/TWICE PER DAY/ORAL		Grand Mal Convulsion Joint Dislocation	Foreign Health Professional	Zyban Tablet - Zyban Aspirin Olanzapine	PS C C		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/04/99ISR Number: 3214713-0Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Angioneurotic Oedema Urticaria		Zyban Sustained Release Tablet	PS	Glaxo Wellcome	ORAL

Date:03/05/99ISR Number: 3214470-8Report Type:Expedited (15-DaCompany Report #A0082673  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Agitation	Health	Wellbutrin Tablet	PS		ORAL
Initial or Prolonged ORAL		Drug Level Above Therapeutic	Professional Other	Lithium Carbonate Tablet	SS		ORAL
ORAL		Feeling Drunk		Olanzapine Tablet	SS		ORAL
ORAL		Nausea		Benzatropine Tablet	SS		ORAL
ORAL		Tremor		Gabapentin Tablet	SS		ORAL
ORAL				Clonazepam Tablet	SS		ORAL
ORAL				Zolpidem Tartrate Tablet	SS		ORAL
ORAL				Sertraline Hydrochloride Tablet	SS		ORAL

Date:03/05/99ISR Number: 3214640-9Report Type:Expedited (15-DaCompany Report #A0074224  
 Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / PER		Asthenia	Health	Wellbutrin	PS		ORAL

Initial or Prolonged Blood Pressure Increased Professional  
DAY

Dizziness Atenolol C  
Nausea Prempro C

Date:03/08/99ISR Number: 3214861-5Report Type:Expedited (15-DaCompany Report #A0084548  
Age:24 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL	Depression  Dyspnoea  Hypersensitivity Insomnia Nausea Oedema Pharyngeal Oedema Pruritus Skin Discolouration Urticaria	Consumer	Zyban Tablet- Zyban	PS		ORAL

Date:03/08/99ISR Number: 3214862-7Report Type:Expedited (15-DaCompany Report #A0080441  
Age:27 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged Other	Confusional State Disorientation Dizziness Dizziness Postural Headache

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Stupor Syncope	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL			Health Professional Company Representative	Wellbutrin Tablets -Controlled Release	PS		ORAL

Date:03/08/99ISR Number: 3214863-9Report Type:Expedited (15-DaCompany Report #A0078251  
Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other 150 MG TWICE PER DAY ORAL		Bipolar I Disorder Condition Aggravated Convulsion Hallucination	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:03/08/99ISR Number: 3215088-3Report Type:Expedited (15-DaCompany Report #A0076758  
Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG SINGLE Hospitalization - DOSE ORAL Initial or Prolonged		Anaemia Cardiac Failure Cerebrovascular Accident Chest Pain Coronary Artery Disease Gastric Ulcer Gastrointestinal Haemorrhage Haematemesis Insomnia Myocardial Infarction Pneumonia Respiratory Failure	Health Professional	Zyban Paracetamol	PS C		ORAL

Sepsis  
Vomiting

Date:03/08/99ISR Number: 3215565-5Report Type:Expedited (15-DaCompany Report #A0084700  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Toxicity Overdose	Consumer	Wellbutrin (Formulation Unknown)	PS		ORAL
100 MG / UNK							
/ ORAL							

Date:03/11/99ISR Number: 3217771-2Report Type:Direct Company Report #  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Bupropion	PS		ORAL
150 MG BID PO							
Hallucination, Auditory Tinnitus							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/12/99ISR Number: 3219018-XReport Type:Expedited (15-DaCompany Report #A0078500  
Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 15 MG / TWICE Initial or Prolonged PER DAY /		Fall	Health	Zyban	PS		ORAL
ORAL		Rib Fracture	Professional				
		Syncope					
		Upper Limb Fracture					

Date:03/12/99ISR Number: 3431285-6Report Type:Periodic Company Report #A0080806  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL	10 DAY	Excitability	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/12/99ISR Number: 3431290-XReport Type:Periodic Company Report #A0080807  
Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Rash Erythematous					
		Rash Papular					

Date:03/12/99ISR Number: 3434609-9Report Type:Periodic Company Report #A0080692  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG/TWICE PER DAY/ ORAL		Burning Sensation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Dermatitis					

Date:03/12/99ISR Number: 3443038-3Report Type:Periodic Company Report #1998001360-1  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dry Mouth	Consumer	Nicorette Otc 2 Mg Smithkline Beecham Consumer Healthcare	PS	Smithkline Beecham Consumer Healthcare	
BUCCAL	BUCCAL			Nicorette Otc 4 Mg Smithkline Beecham Consumer Healthcare	SS	Smithkline Beecham Consumer Healthcare	
BUCCAL	BUCCAL	44 DAY		Wellbutrin (Bupropion) Glaxo Wellcome	SS		ORAL
ORAL							

Date:03/15/99ISR Number: 3220366-8Report Type:Direct Company Report #  
 Age:42 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG QD X		Feeling Abnormal	Health	Zyban	PS	Glaxo/Wellcome	ORAL
3D PO 150 MG		Nervous System Disorder	Professional				
BID PO	11 DAY	Nightmare					
		Panic Attack		Effexor Xr	C		
		Thinking Abnormal		Haloperidol	C		
		Vomiting		Clonazepam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Estradiol C  
Tums/Roloids C

Date:03/17/99ISR Number: 3222051-5Report Type:Expedited (15-DaCompany Report #A0085087  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Injury	Health	Wellbutrin Tablet	PS		ORAL
UNK/ UNK/		Overdose	Professional				
ORAL							

Date:03/17/99ISR Number: 3222058-8Report Type:Expedited (15-DaCompany Report #A0080126  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Diabetes Mellitus	Health	Wellbutrin			
Hospitalization -		Inadequate Control	Professional	Tablet-Controlled			
Initial or Prolonged		Grand Mal Convulsion	Company	Release	PS		ORAL
150 MG / SEE							
Other		Insomnia	Representative				
TEXT/ ORAL				Insulin	C		

Date:03/17/99ISR Number: 3222323-4Report Type:Expedited (15-DaCompany Report #A0085804  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Health	Wellbutrin			
			Professional	Tablet-Controlled			
150 MG/TWICE			Company	Release	PS		ORAL
			Representative				
PER DAY/ORAL				Sildenafil Citrate	C		

Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/TWICE			Representative				
PER DAY/ORAL	1	MON		Tamoxifen	C		

Date:03/19/99ISR Number: 3225172-6Report Type:Periodic Company Report #8-97286-019L

Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cardiovascular Disorder	Consumer	Redux	PS		ORAL
15 MG TWICE		Palpitations					
DAILY ORAL		Pulmonary Hypertension		Fastin (Phentermine)	SS		ORAL
30 MG ONCE							
DAILY ORAL				Ionamin (Phentermine)	SS		ORAL
15-30 MG							
DAILY ORAL	18	WK		Pondimin (Fenfluramine) Tablet	SS		ORAL
20 MG THREE							
TIMES DAILY							
ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

100-150 MG  
 TWICE DAILY  
 ORAL

Wellbutrin  
 (Bupropion) SS ORAL

Date:03/22/99ISR Number: 3224264-5Report Type:Expedited (15-DaCompany Report #A0086553  
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG/TWICE	Angina Pectoris	Foreign	Zyban Tablet	PS		ORAL
PER DAY/ORAL		Myocardial Infarction	Health				
			Professional				

Date:03/22/99ISR Number: 3224298-0Report Type:Expedited (15-DaCompany Report #A0086052  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG TWICE	Diabetes Mellitus	Foreign	Zyban Tablet (Zyban)	PS		ORAL
Initial or Prolonged	DAILY ORAL	Polyuria	Health				
Other		Vision Blurred	Professional				

Date:03/22/99ISR Number: 3224381-XReport Type:Expedited (15-DaCompany Report #A0077049  
 Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	250 MG/ORAL	Bronchiolitis	Foreign	Wellbutrin	PS		ORAL
Hospitalization -		Dyspnoea Exacerbated	Health	Lithium Salt	C		
Initial or Prolonged		Hypoxia	Professional	Clomipramine Hcl	C		
Other		Lung Infiltration		Clonazepam	C		
		Pyrexia		Lithium Carbonate	C		

Date:03/22/99ISR Number: 3224382-1Report Type:Expedited (15-DaCompany Report #A0086318  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis	Consumer	Zyban	PS		ORAL
150 MG/TWICE		Dysphagia					
PER DAY/ORAL		Eyelid Oedema		Zyban	SS		ORAL
150 MG/SINGLE		Face Oedema					
DOSE/ORAL		Pharyngeal Oedema		Lansoprazole	C		
				Lorcet	C		

Date:03/22/99ISR Number: 3224414-0Report Type:Expedited (15-DaCompany Report #A0086224  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Medication Error	Foreign	Zyban	PS		ORAL
150 MG/TWICE		Myocardial Infarction	Literature				
PER DAY/ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/22/99ISR Number: 3224451-6Report Type:Expedited (15-DaCompany Report #A0081012

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complex Partial Seizures Grand Mal Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL	2 WK		Representative				

Date:03/22/99ISR Number: 3225548-7Report Type:Direct

Company Report #

Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia		Zyban	PS		ORAL
ORAL DAILY		Malaise Pyrexia Serum Sickness Urticaria White Blood Cell Count Increased					

Date:03/23/99ISR Number: 3225607-9Report Type:Direct

Company Report #

Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Angioneurotic Oedema		Wellbutrin 150 Sr	PS		ORAL
ONE TAB AT		Arthralgia					
Intervention to							
8:30 AM AND		Dyspnoea					
Prevent Permanent							
12:00 PM							
Impairment/Damage		Eyelid Oedema Hypoaesthesia Insomnia Joint Swelling Pain Pharyngolaryngeal Pain Swelling					

Date:03/23/99ISR Number: 3420370-0Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #5686/20536

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Jittery Paranoia	Consumer Company	Nicotrol	PS	Pharmacia And Upjohn Co	
TRANSDERMAL	15 MG -		Representative				
1Q1DY; T.DERM				Wellbutrin Lithium	SS C		

Date:03/24/99ISR Number: 3226001-7Report Type:Direct  
Age:55 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Bupropion 100mg Sr Tab	PS		ORAL
Other							
100MG SR BID							
PO				Aspirin Temazepam Buspar Ranitidine Tamslosin	C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Multivitamin.Mineral C

Date:03/25/99ISR Number: 3226398-8Report Type:Expedited (15-DaCompany Report #A0085728  
 Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
TWICE PER DAY/ ORAL		Insomnia					
		Joint Swelling Pain					

Date:03/25/99ISR Number: 3226401-5Report Type:Expedited (15-DaCompany Report #A0086604  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL			Representative	Aspartame (Formulation Unknown)	SS		

Date:03/25/99ISR Number: 3226406-4Report Type:Expedited (15-DaCompany Report #A0086619  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL			Representative	Aspartame (Formulation Unknown)	SS		

Date:03/26/99ISR Number: 3227632-0Report Type:Expedited (15-DaCompany Report #A0085426  
Age:15 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Aspiration	Consumer	Wellbutrin			
Initial or Prolonged	Convulsion		Tablet-Controlled			
	Intentional Misuse		Release	PS		ORAL
3 G / SINGLE						
DOSE/ ORAL	Vomiting					

Date:03/26/99ISR Number: 3227814-8Report Type:Expedited (15-DaCompany Report #A0085560  
Age:16 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Convulsion	Consumer	Wellbutrin Tablet			
Initial or Prolonged	Intentional Misuse		-Controlled Release	PS		ORAL
3 G/SINGLE						
DOSE/ORAL	Medication Error					
	Nausea					
	Vomiting					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/26/99ISR Number: 3227845-8Report Type:Expedited (15-DaCompany Report #A0086438  
Age:70 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG ORAL	Amnesia	Consumer	Wellbutrin	PS		ORAL
Initial or Prolonged	Bipolar I Disorder Glaucoma Intraocular Pressure Increased Visual Acuity Reduced		Venlafaxine Hydrochloride Lithium Salt	C C		

Date:03/28/99ISR Number: 3332514-XReport Type:Periodic Company Report #A0087342  
Age:41 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose 150 MG / TWICE PER DAY / ORAL	Arthralgia Balance Disorder Burning Sensation Dermatitis Hypersensitivity Pain In Extremity Pruritus Swelling	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:03/29/99ISR Number: 3229434-8Report Type:Expedited (15-DaCompany Report #A0086620  
Age:33 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 150 MG TWICE PER DAY ORAL ORAL	Alcohol Interaction Completed Suicide Drug Toxicity Head Injury	Health Professional	Wellbutrin Tablet - Controlled Release Ethanol (Formulation Unknown)	PS SS		ORAL ORAL

ORAL

Dextropropoxyphene Hcl Tablet	SS
Buspirone Hydrochloride	C

Date:03/29/99ISR Number: 3229438-5Report Type:Expedited (15-DaCompany Report #A0077426  
 Age:22 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
100 MG TWICE							
PER DAY ORAL							
				Fluoxetine Hydrochloride (Formulation Unknown)	SS		ORAL
20 MG PER DAY							
ORAL				Metoprolol Tartrate Paracetamol	SS SS		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/99ISR Number: 3229443-9Report Type:Expedited (15-DaCompany Report #A0086553

Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Angina Pectoris	Foreign	Zyban Tablet	PS		ORAL
150 MG TWICE							
Other		Myocardial Infarction	Health				
PER DAY ORAL			Professional	Glucophage	C		
				Conjugated Estrogens	C		
				Atorvastatin Calcium	C		
				Glibenclamide	C		

Date:03/30/99ISR Number: 3230085-XReport Type:Expedited (15-DaCompany Report #9911866

Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Abdominal Pain Upper	Consumer	Zoloft Tablets	PS		ORAL
ORAL							
Intervention to		Drug Interaction		Wellbutrin	SS		
Prevent Permanent		Flatulence		Marijuana	SS		
Impairment/Damage				Beer	SS		

Date:03/31/99ISR Number: 3230925-4Report Type:Expedited (15-DaCompany Report #A0065816

Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aggression	Health	Zyban Tablet- Zyban	PS		ORAL
150 MG ORAL							
		Agitation	Professional	Medroxyprogesterone			
		Amnesia		Ace	C		
		Anger					
		Anxiety					
		Breast Pain					
		Condition Aggravated					
		Disorientation					
		Disturbance In Attention					
		Dizziness					
		Feeling Abnormal					
		Feeling Jittery					

Fluid Retention  
 Galactorrhoea  
 Insomnia  
 Migraine  
 Paranoia  
 Serotonin Syndrome  
 Suicidal Ideation  
 Tinnitus

Date:03/31/99ISR Number: 3230927-8Report Type:Expedited (15-DaCompany Report #A0082610

Age:55 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG PER Initial or Prolonged DAY ORAL	Anaphylactoid Reaction	Foreign	Zyban Tablet - Zyban	PS		ORAL
	Angioneurotic Oedema	Health				
	Pruritus	Professional	Thyroxine Sodium	C		
	Rash Erythematous		Hrt	C		
	Rash Morbilliform					
	Tongue Oedema					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/31/99ISR Number: 3230929-1Report Type:Expedited (15-DaCompany Report #A0086505

Age:23 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG THREE TIMES PER DAY ORAL	Confusional State Grand Mal Convulsion Hallucination Illusion Memory Impairment Sedation	Health Professional Company Representative	Wellbutrin Tablet- Controlled Release     Olanzapine (Formulation Unknown)	PS     SS		ORAL
10 MG PER DAY			Fluvoxamine Maleate Lithium Salt Propranolol Hydrochloride Lorazepam Thyroxine Sodium	C C C C C C		

Date:04/01/99ISR Number: 3231319-8Report Type:Expedited (15-DaCompany Report #A0087673

Age:30 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 150 MG/SINGLE DOSE	Death	Health Professional Company Representative	Zyban	PS		

Date:04/01/99ISR Number: 3231322-8Report Type:Expedited (15-DaCompany Report #A0087656

Age:27 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1	Anaphylactic Shock	Study	Zyban	PS		ORAL

TABLET/TWICE

Dizziness

Consumer

PER DAY/ORAL

Dysphagia

Dyspnoea

Pseudoephedrine Hcl C

Fall

Gingival Hypertrophy

Hypersensitivity

Hypoventilation

Influenza

Insomnia

Joint Swelling

Myalgia

Nasal Oedema

Oedema Peripheral

Pain In Extremity

Swelling

Tongue Oedema

Urticaria

Weight Decreased

Date:04/01/99ISR Number: 3231458-1Report Type:Expedited (15-DaCompany Report #9911957

Age: Gender:Female I/FU:I

Outcome

Hospitalization -

Initial or Prolonged

Required

Intervention to

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Prevent Permanent Impairment/Damage

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL			Bradycardia	Health	Zoloft Tablets	PS		ORAL
ORAL			Drug Interaction	Professional	Zyban	SS		ORAL
			Lethargy Syncope					

Date:04/02/99ISR Number: 3235621-5Report Type:Periodic Company Report #9809438  
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL			Visual Disturbance	Health	Cardura	PS		ORAL
ORAL				Professional	Wellbutrin	SS		ORAL
					Cyclosporine Zestril	C C		

Date:04/05/99ISR Number: 3233013-6Report Type:Direct Company Report #  
Age:41 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG PO QD Initial or Prolonged			Agitation		Wellbutrin Sr	PS		ORAL
			Convulsion Psychotic Disorder					

Date:04/05/99ISR Number: 3233084-7Report Type:Expedited (15-DaCompany Report #A0087951  
Age:24 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG/TWICE			Blood Glucose Decreased	Foreign	Zyban	PS		ORAL

PER DAY/ORAL Condition Aggravated Consumer

Convulsion  
Fall

Date:04/06/99ISR Number: 3234412-9Report Type:Direct Company Report #  
Age:19 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100 MG PO	Convulsion		Wellbutrin	PS		ORAL
Initial or Prolonged TID	Vomiting		Allegra	C		
			Azithromycin	C		
			Proventil Inhaler	C		
			Azmacort Inhaler	C		
			Prednix	C		

Date:04/07/99ISR Number: 3234550-0Report Type:Expedited (15-DaCompany Report #A0087443  
Age: Gender:Male I/FU:I

Outcome	PT
Congenital Anomaly	Atrial Septal Defect Cardiac Malposition Complications Of Maternal Exposure To Therapeutic Drugs Hypoplastic Left Heart



Required Metrorrhagia Zyban PS Glaxowellcome ORAL  
 300MG BID  
 Intervention to  
 ORAL  
 Prevent Permanent Depo Provera SS Upjohn  
 Impairment/Damage

Date:04/12/99ISR Number: 3237985-5Report Type:Expedited (15-DaCompany Report #A0088107

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Foreign	Wellbutrin			
Other		Fall	Health	Tablet-Controlled			
		Upper Limb Fracture	Professional	Release	PS		ORAL
ORAL WEEKS				Carbamazepine	C		

Date:04/12/99ISR Number: 3237987-9Report Type:Expedited (15-DaCompany Report #A0088588

Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Angina Pectoris	Foreign	Zyban Tablet-Zyban	PS		ORAL
Hospitalization -		Cardiac Disorder	Consumer				
150 MG/TWICE		Dyspnoea		Nicotine	C		
Initial or Prolonged		Hypertension					
PER DAY/ORAL		Swelling					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/12/99ISR Number: 3238231-9Report Type:Expedited (15-DaCompany Report #A0081071  
Age:44 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - Initial or Prolonged 150 MG/PER Other DAY/ORAL	Respiratory Distress Urticaria	Health Professional	Wellbutrin Tablet-Controlled Release  Aspirin	PS  C		ORAL

Date:04/12/99ISR Number: 3238232-0Report Type:Expedited (15-DaCompany Report #A0088466  
Age:79 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY ORAL	Dysphonia Feeling Jittery Tremor	Consumer	Zyban Tablet - Zyban  Ranitidine Hydrochloride Thyroxine Sodium Hydrochlorothiazide Digoxin Paracetamol Potassium Salt	PS  C C C C C C		ORAL

Date:04/13/99ISR Number: 3239039-0Report Type:Expedited (15-DaCompany Report #A0079587  
Age:32 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG /  TWICE PER DAY	Burning Sensation Dermatitis Dry Skin Eczema Pruritus Rash Erythematous	Health Professional	Zyban Tablet - Zyban	PS		ORAL

Rash Macular  
Rash Vesicular  
Skin Ulcer  
Urticaria

Date:04/13/99ISR Number: 3239041-9Report Type:Expedited (15-DaCompany Report #A0080356  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG / TWICE PER DAY		Angina Pectoris Blood Pressure Increased Cardiac Disorder Cardiac Murmur Coronary Artery Occlusion Dermatitis Oedema Peripheral Urticaria	Consumer	Zyban Tablets - Zyban  Dyazide	PS  C		ORAL

Date:04/13/99ISR Number: 3239043-2Report Type:Expedited (15-DaCompany Report #A0084557  
Age:49 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Anxiety Disorder Disorientation

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Psychotic Disorder Tremor				
Dose	Duration		Report Source	Product	Role	Manufacturer
			Health Professional	Zyban Tablet - Zyban	PS	
150 MG /			Company Representative			ORAL
TWICE PER DAY						

Date:04/14/99ISR Number: 3239702-1Report Type:Direct Company Report #  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia Hypoaesthesia Joint Stiffness Lacrimation Increased		Zyban	PS		

Date:04/14/99ISR Number: 3239860-9Report Type:Expedited (15-DaCompany Report #A0086275  
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hemiparesis Hypoaesthesia	Foreign	Zyban Tablet - Zyban	PS		ORAL
Disability			Health Professional				
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:04/14/99ISR Number: 3239896-8Report Type:Direct Company Report #  
Age:6 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS		ORAL
Other		Depressed Level Of					
50MG PO TID	6 WK						

Consciousness  
Gaze Palsy  
Movement Disorder  
Pharyngitis  
Pyrexia  
Speech Disorder  
Urinary Incontinence

Date:04/14/99ISR Number: 3239910-XReport Type:Direct  
Age:50 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema		Zyban	PS		
SMOKING		Dyspnoea					
CESSATION		Eyelid Oedema		Xanax	C		
		Face Oedema		Indenal	C		
		Pruritus					
		Urticaria					

Date:04/14/99ISR Number: 3239943-3Report Type:Direct  
Age:63 YR Gender:Female I/FU:I

Company Report #

Outcome  
Required  
Intervention to

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Freedom Of Information (FOI) Report

Prevent Permanent  
Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 BID	15 DAY	Renal Disorder	Health	Wellbutrin	PS		
1 QD	3 WK		Professional	Wellbutrin Sr 100mg	SS		
				Atenolol	C		
				Medroxyprogesterone	C		
				Hctz	C		
				Lisinopril	C		
				Cometidine	C		
				Zoloft	C		
				Premarin	C		

Date:04/15/99ISR Number: 3239857-9Report Type:Expedited (15-DaCompany Report #A0081012

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Breath Odour	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Complex Partial Seizures	Professional				
PER DAY /		Convulsion	Company				
ORAL		Drug Eruption	Representative	Allopurinol	C		
		Grand Mal Convulsion		Simvastatin	C		
		Lower Respiratory Tract		Aspirin	C		
		Infection		Frusemide	C		
		Rash Pruritic		Atenolol	C		
		Rhinitis		Omeprazole	C		
		Sinusitis		Nitroglycerin	C		
		Skin Exfoliation		Glucophage	C		
		Staphylococcal Infection		Isosorbide			
		Streptococcal Infection		Mononitrate	C		
		Urticaria		Potassium Chloride	C		
				Nsaid	C		

Date:04/16/99ISR Number: 3241403-0Report Type:Expedited (15-DaCompany Report #WAES 99040593

Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Back Pain	Literature	Tab Mevacor Unk	PS		
Initial or Prolonged		Grand Mal Convulsion	Other	Wellbutrin Unk	SS		
		Intentional Misuse					
		Medication Error					

Date:04/16/99ISR Number: 3241604-1Report Type:Expedited (15-DaCompany Report #A0087945

Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Convulsion					
Initial or Prolonged		Epistaxis		Claritin-D	C		
DAY / ORAL		Haematemesis		Nicotine	C		
Disability		Loss Of Consciousness					
		Nasal Congestion					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/19/99ISR Number: 3242407-4Report Type:Expedited (15-DaCompany Report #A0080313

Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Health	Zyban	PS		ORAL
150 MG TWICE		Anxiety	Professional				
PER DAY ORAL		Convulsion		Percocet	C		
		Dry Mouth		Diazepam	C		
		Haematemesis		Quinalbarbitone			
		Head Injury		Sodium	C		
		Insomnia					
		Photopsia					
		Pneumonia					
		Tremor					
		Vision Blurred					

Date:04/19/99ISR Number: 3242408-6Report Type:Expedited (15-DaCompany Report #A0086318

Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
Other		Dysphagia					
150 MG TWICE		Eyelid Oedema					
PER DAY; 150		Face Oedema					
MG SINGLE		Hypersensitivity		Lansoprazole	C		
DOSE ORAL		Pharyngeal Oedema		Lorcet	C		

Date:04/19/99ISR Number: 3242447-5Report Type:Expedited (15-DaCompany Report #A0087807

Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Difficulty In Walking	Health	Wellbutrin			
Disability		Diplopia	Professional	Tablet-Controlled			
Other							

150 MG /  
 TWICE PER DAY  
 / ORAL

Dyspnoea  
 Myasthenia Gravis  
 Sensation Of Heaviness

Release PS ORAL

Pyridostigmine  
 Bromide C  
 Prostigmine C  
 Beclomethasone  
 Dipropion C  
 Bromfed-Pd C  
 Sumatriptan  
 Succinate C  
 Tylenol No. 3 C

Date:04/19/99ISR Number: 3242554-7Report Type:Expedited (15-DaCompany Report #A0082447  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Health	Wellbutrin	PS		ORAL
150 MG /		Dermatitis	Professional				
TWICE PER DAY		Dry Throat					
/ ORAL		Oedema		Amoxicillin	C		
		Petechiae					
		Pharyngolaryngeal Pain					
		Pruritus					
		Throat Tightness					
		Urticaria					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/19/99ISR Number: 3242555-9Report Type:Expedited (15-DaCompany Report #A0085728  
 Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	TWICE PER DAY	Angioneurotic Oedema	Health	Wellbutrin	PS		ORAL
/ ORAL		Arthralgia	Professional				
		Depression					
		Dermographism					
		Dyspnoea					
		Emotional Distress					
		Insomnia					
		Joint Swelling					
		Myalgia					
		Pain					
		Pharyngeal Oedema					
		Pruritus					
		Serum Sickness					
		Urticaria					

Date:04/19/99ISR Number: 3242557-2Report Type:Expedited (15-DaCompany Report #A0086620  
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health	Wellbutrin			
		Head Injury	Professional	Tablet-Controlled Release	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
ORAL				Ethanol (Formulation Unknown)	SS		ORAL
ORAL				Dextropropoxyphene Hcl Tablet	SS		ORAL
ORAL				Buspirone Hydrochloride	C		

Date:04/19/99ISR Number: 3242666-8Report Type:Expedited (15-DaCompany Report #1154856A  
Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Health Professional	Unk Acetaminophen Product Wellbutrin Tablet-Controlled Release	PS		ORAL
	100 MG, BID, PO				SS		
				Fluoxetine Hydrochloride (Formulation Unknown)	SS		ORAL
	20 MG/DAY PO			Metoprolol Tartrate	SS		

Date:04/20/99ISR Number: 3320135-4Report Type:Periodic Company Report #1999SUS0049  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Effect Decreased Drug Interaction	Health Professional	Sustiva Wellbutrin	PS SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/21/99ISR Number: 3322188-6Report Type:Periodic  
 Age:40 YR Gender:Male I/FU:I

Company Report #99USA10029

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Health	Diovan Capsule 80 Mg			
		Dizziness	Professional	(Valsartan)	PS		ORAL
80 MG QD ORAL		Erectile Dysfunction		Wellbutrin Tablets			
				(Amfebutamone Hydrochloride)	SS		ORAL
ORAL							

Date:04/22/99ISR Number: 3244821-XReport Type:Expedited (15-DaCompany Report #A0089661  
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sudden Death	Health	Wellbutrin			
Death			Professional	Tablet-Controlled Release	PS		ORAL
150 MG /							
TWICE PER DAY							

/ ORAL

Olanzapine	C
Quetiapine Fumarate	C
Temazepam	C
Prempro	C
Dyazide	C
Ibuprofen	C

Date:04/22/99ISR Number: 3244824-5Report Type:Expedited (15-DaCompany Report #A0083661  
 Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alcohol Interaction	Foreign	Zyban Tablet - Zyban	PS		ORAL
Other		Blood Glucose Decreased	Health				
150 MG /							
TWICE PER							

DAY/ ORAL	Convulsion	Professional				
	Eye Rolling		Human Short-Act,			
	Hypoglycaemia		Insulin		C	
	Medication Error		Human Int/Long			
			Insulin		C	

Date:04/22/99ISR Number: 3244835-XReport Type:Expedited (15-DaCompany Report #A0089381  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain	Foreign	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/PER			Consumer				
ORAL/ORAL							

Date:04/26/99ISR Number: 3246876-5Report Type:Expedited (15-DaCompany Report #A0089898  
 Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Communication Disorder	Foreign	Zyban Tablet - Zyban	PS		ORAL
150MG/PER		Disturbance In Attention	Consumer				
DAY/ORAL		Feeling Abnormal					
		Feeling Drunk					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/26/99ISR Number: 3246877-7Report Type:Expedited (15-DaCompany Report #A0089735

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Zyban Tablet - Zyban	PS		
		Drug Interaction	Professional	Alprazolam (Formulation Unknown)	SS		

Date:04/26/99ISR Number: 3246879-0Report Type:Expedited (15-DaCompany Report #A0089717

Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris	Foreign	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE							
		Angiopathy	Consumer				
PER DAY/ORAL							

Date:04/26/99ISR Number: 3246898-4Report Type:Expedited (15-DaCompany Report #A0088445

Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Abdominal Pain	Foreign	Zyban Tablets	PS		ORAL
150 MG TWICE							
		Constipation	Health				
PER DAY ORAL	2 WK	Fluid Retention	Professional	Oral Contraceptive	C		
		Irritability					
		Oliguria					

Date:04/26/99ISR Number: 3246900-XReport Type:Expedited (15-DaCompany Report #A0089140

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Hallucination	Health	Wellbutrin Tablet-			
Initial or Prolonged		Intentional Misuse	Professional	Controlled Release	PS		ORAL
150 MG ORAL							

Other  
(SEE TEXT) Suicide Attempt

Date:04/26/99ISR Number: 3246901-1Report Type:Expedited (15-DaCompany Report #B0065751  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fatigue	Health	Wellbutrin	PS		ORAL
ORAL		Liver Function Test	Professional	Troglitazone Tablet	SS		ORAL
300 MG TWICE		Abnormal					
PER DAY ORAL		Weight Increased		Atorvastatin	C		
				Estrogen	C		
				Metformin	C		
				Lansoprazole	C		
				Clonazepam	C		
				Insulin	C		
				Thioridazine	C		
				Pyridoxine	C		

Date:04/26/99ISR Number: 3247143-6Report Type:Expedited (15-DaCompany Report #A0089368  
Age:42 YR Gender:Male I/FU:I

Outcome  
Death  
Hospitalization -  
Initial or Prolonged

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG		Alcohol Poisoning Cardiac Arrest Convulsion	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
		Loss Of Consciousness Overdose		Ethanol	SS		

Date:04/26/99ISR Number: 3250090-7Report Type:Periodic Company Report #A0088435  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other TWICE PER DAY		Agitation Convulsion Psychotic Disorder	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL							

Date:04/26/99ISR Number: 3250091-9Report Type:Periodic Company Report #A0088100  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG TWICE PER DAY ORAL	11 DAY	Grand Mal Convulsion Loss Of Consciousness	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:04/26/99ISR Number: 3250092-0Report Type:Periodic Company Report #A0087970  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Health	Wellbutrin			

150 MG ORAL  
Professional Company Release PS ORAL  
Representative

Date:04/26/99ISR Number: 3250093-2Report Type:Periodic Company Report #A0087647  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		

Date:04/26/99ISR Number: 3250094-4Report Type:Periodic Company Report #A0087481  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL

150 MG TWICE  
PER DAY ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/26/99ISR Number: 3250095-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0087281

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL			Representative				

Date:04/26/99ISR Number: 3250096-8Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0086240

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG ORAL			Representative	Tricyclin Antidepressant	C		

Date:04/26/99ISR Number: 3250097-XReport Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0085964

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Rash Generalised	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG TWICE				Alprazolam	C		
PER DAY ORAL							

Date:04/26/99ISR Number: 3250098-1Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0084704

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Cerebrovascular Accident Consumer Wellbutrin  
Dyspnoea Tablet-Controlled  
Headache Release PS ORAL  
150 MG TWICE  
PER DAY ORAL

Date:04/26/99ISR Number: 3250099-3Report Type:Periodic Company Report #A0084373  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysgeusia	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL				Thyroxine Sodium	C		

Date:04/26/99ISR Number: 3250100-7Report Type:Periodic Company Report #A0084031  
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG TWICE				Fluoxetine	C		
PER DAY ORAL				Clonazepam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Quetiapine Fumarate C

Date:04/26/99ISR Number: 3250101-9Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0083747

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL	3 WK	Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release  Olanzapine (Formulation Unknown)	PS  SS		ORAL

Date:04/26/99ISR Number: 3250102-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0082995

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Convulsion Overdose	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:04/26/99ISR Number: 3250103-2Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #A0082989

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Convulsion Mastication Disorder Tooth Injury	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release  Paroxetine Hydrochloride	PS  C		ORAL

Date:04/26/99ISR Number: 3250104-4Report Type:Periodic Company Report #A0082594  
Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG TWICE			Representative				
PER DAY ORAL							

Date:04/26/99ISR Number: 3250105-6Report Type:Periodic Company Report #A0082573  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Dyspnoea Hypersensitivity Pruritus	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG TWICE		Rash Generalised					
PER DAY ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/26/99ISR Number: 3250106-8Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #A0080712

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypertension Hypoaesthesia	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG PER DAY ORAL				Atorvastatin Calcium Conjugated Estrogens Amitriptyline	C C C		

Date:04/26/99ISR Number: 3250107-XReport Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0080556

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Stevens-Johnson Syndrome	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
ORAL							

Date:04/26/99ISR Number: 3250108-1Report Type:Periodic  
Age:10 YR Gender:Male I/FU:I

Company Report #A0080551

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG PER DAY ORAL				Methylphenidate Hcl Diphenhydramine Hcl	C C		

Date:04/26/99ISR Number: 3250109-3Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0080543

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Electroencephalogram Abnormal	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG TWICE							
PER DAY ORAL							
				Risperidone	C		
				Lithium Carbonate	C		
				Thyroxine Sodium	C		

Date:04/26/99ISR Number: 3250110-XReport Type:Periodic Company Report #A0080518  
Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG TWICE							
PER DAY ORAL							
				Antihistamine	C		
				Cetirizine			
				Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/26/99ISR Number: 3250111-1Report Type:Periodic  
Age:32 YR Gender:Male I/FU:I

Company Report #A0080224

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Overdose	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG THREE TIMES PER DAY			Representative				
ORAL				Amitriptyline Hcl	C		

Date:04/26/99ISR Number: 3250112-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0080213

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Paranoia	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG TWICE PER DAY ORAL				Nefazodne Hcl	C		
				Olanzapine	C		
				Pyridostigimine Bromide	C		

Date:04/26/99ISR Number: 3250113-5Report Type:Periodic  
Age:18 YR Gender:Female I/FU:I

Company Report #A0080125

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG TWICE PER DAY ORAL			Representative				
				Methytlphenidate Hcl	C		

Date:04/26/99ISR Number: 3250114-7Report Type:Periodic Company Report #A0079398  
 Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema	Health	Wellbutrin			
		Drug Toxicity	Professional	Tablet-Controlled			
		Eyelid Oedema	Company	Release	PS		ORAL
ORAL							
		Face Oedema	Representative				
		Oedema Peripheral					
		Pruritus					
		Rash Erythematous					
		Rash Generalised					
		Urticaria					

Date:04/29/99ISR Number: 3250150-0Report Type:Expedited (15-DaCompany Report #A0090004  
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardiac Arrest	Foreign	Zyban Tablet-	PS	Zyban	ORAL
150 MG ORAL							
		Chest Discomfort		Lorazepam	C		
		Dyspnoea		Cefurozime Axetil	C		
		Pulmonary Embolism		Combivent	C		
				Budesonide	C		
				Tetracycline	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Theophylline C

Date:04/29/99ISR Number: 3250159-7Report Type:Expedited (15-DaCompany Report #A0081071

Age:44 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 150 MG PER Initial or Prolonged DAY ORAL Other	Dermatitis Oedema Mouth Pharyngeal Oedema Pruritus Respiratory Distress Tongue Oedema Urticaria	Health Professional	Wellbutrin Tablets- Controlled Release  Aspirin	PS  C		ORAL

Date:04/29/99ISR Number: 3250162-7Report Type:Expedited (15-DaCompany Report #A0088100

Age:24 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Other 150 MG TWICE  PER DAY ORAL	Cyanosis Fatigue Grand Mal Convulsion Loss Of Consciousness Muscle Contractions Involuntary Myalgia	Health Professional	Wellbutrin Tablet- Controlled Release  Pethidine Hydrochloride Ibuprofen Darvocet-N	PS  C C C		ORAL

Date:04/29/99ISR Number: 3251132-5Report Type:Direct

Age: Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150MG PO Hospitalization - QD/BID	Arterial Occlusive Disease		Zyban	PS	Glaxo Wellcome	ORAL

Initial or Prolonged  
Other

Finger Amputation  
Gangrene  
Necrosis Ischaemic  
Pain  
Pain In Extremity  
Petechiae  
Pyrexia  
Skin Discolouration

Prednisone

C

Date:04/30/99ISR Number: 3251007-1Report Type:Expedited (15-DaCompany Report #A0089788

Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / PER Initial or Prolonged DAY / ORAL	Anxiety Headache Hypertension Loss Of Consciousness Tinnitus	Foreign Consumer	Zyban Tablet - Zyban	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/30/99ISR Number: 3251348-8Report Type:Direct  
 Age:73 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 100MG QD PO; Initial or Prolonged 100MG BID PO	Chest Pain  Rash Pruritic  Urticaria		Bupropion Sr 100mg	PS		ORAL
			Genkgo Biloba	C		
			Nizatidine	C		
			Estropipate	C		
			Vitamin E	C		
			Mult E	C		
			Multi=Vit	C		

Date:05/03/99ISR Number: 3252406-4Report Type:Expedited (15-DaCompany Report #A0090134  
 Age:65 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG/ TWICE  PER DAY /  ORAL	Depression  Dizziness  Feeling Abnormal  Listless Loss Of Consciousness	Foreign  Consumer	Zyban Tablet - Zban	PS		ORAL

Date:05/03/99ISR Number: 3252413-1Report Type:Expedited (15-DaCompany Report #A0089368  
 Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Hospitalization - Initial or Prolonged 200 TABLET / Other SINGLE DOSE/  ORAL	Cardiac Arrest Convulsion Drug Toxicity  Intentional Misuse  Loss Of Consciousness	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL

Suicide Attempt

Ethanol (Formulation  
Unknown)

SS

Date:05/03/99ISR Number: 3252782-2Report Type:Expedited (15-DaCompany Report #A0089575

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion Medication Error	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
600 MG / UNK							
/ ORAL				Mirtazapine Mytussin Ac	C C		

Date:05/03/99ISR Number: 3252784-6Report Type:Expedited (15-DaCompany Report #A0089899

Age:69 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Coordination Abnormal Demyelinating Polyneuropathy Disturbance In Attention Dysarthria Educational Problem Headache Hypoaesthesia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Memory Impairment Oedema Peripheral Pain Paraesthesia	Foreign Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:05/03/99ISR Number: 3252785-8Report Type:Expedited (15-DaCompany Report #A0090630  
Age: Gender:Unknown I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	UNK / UNK / ORAL	UNK / UNK / ORAL	Coma Convulsion Overdose	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release Ethanol Liquid	PS SS		ORAL ORAL

Date:05/03/99ISR Number: 3252786-XReport Type:Expedited (15-DaCompany Report #A0075655  
Age:49 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY / ORAL			Grand Mal Convulsion Status Epilepticus	Foreign Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/03/99ISR Number: 3253378-9Report Type:Expedited (15-DaCompany Report #A0090016  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Foreign	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Hypertension	Consumer				
TWICE PER DAY		Tremor		Acebutolol			
				Hydrochloride	C		
				Thyroxine Sodium	C		
				Desipramine	C		
				Clonazepam	C		
				Clonazepam	C		
				Blood Pressure			
				Medication	C		

Date:05/03/99ISR Number: 3253381-9Report Type:Expedited (15-DaCompany Report #A0090299

Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain	Foreign	Zyban Tablet - Zyban	PS		ORAL
150 MG		Chest Pain					
		Pericarditis					
		Vomiting					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/04/99ISR Number: 3252788-3Report Type:Direct  
Age:50 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis	Health	Bupropion 100 Mgm			
100MG BID		Urticaria	Professional	Bid	PS		
				Clotriazole	C		
				Furosemide	C		
				Hemorroidal Rectal			
				Supp	C		
				Lisinopril	C		
				Naproxen	C		
				Nifidepril	C		
				Nystaten	C		
				Kcl	C		
				Povodine - Iodine	C		
				Simvastatin	C		

Date:05/05/99ISR Number: 3254911-3Report Type:Direct  
Age:70 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Convulsion		Zyban	PS		ORAL
150MG BID							
Hospitalization -							
ORAL							
Initial or Prolonged				Lorazepam	C		
Required				Glipizide	C		
Intervention to				Diltiazem (Tiazac)	C		
Prevent Permanent				Trazodone Hcl	C		
Impairment/Damage				Pentoxifylline	C		

Date:05/06/99ISR Number: 3255610-4Report Type:Expedited (15-DaCompany Report #A0075934  
Age:42 YR Gender:Female I/FU:F

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dermatitis	Foreign	Zyban Tablet- Zyban	PS		ORAL
150 MG /							

Initial or Prolonged      Dizziness      Health  
TWICE PER      Lung Disorder      Professional  
DAY / ORAL      Pruritus

Date:05/10/99ISR Number: 3257946-XReport Type:Expedited (15-DaCompany Report #A0090921  
Age:51 YR    Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ PER		Bradycardia	Foreign	Zyban Tablet-Zyban	PS		ORAL
Initial or Prolonged DAY/ ORAL		Chest Pain	Consumer				
		Dyspnoea Hypotension Pneumonia		Multiple Medication	C		

Date:05/10/99ISR Number: 3257966-5Report Type:Expedited (15-DaCompany Report #A0090688  
Age:33 YR    Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150 MG		Amnesia Cerebrovascular Accident	Consumer	Wellbutrin Tablet - Controlled Release	PS		ORAL
TWICE PER DAY ORAL		Choking Hemiplegia Hypoaesthesia					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/11/99ISR Number: 3259696-2Report Type:Direct  
Age:45 YR Gender: I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Dyskinesia Dystonia		Wellbutrin 100mg Glaxo Wellcome	PS	Glaxo Wellcome	
100MG 2X 1ST DAY THEN 100MG 3X PER DAY				Zyban 150mg Wellcome	SS	Glaxo Wellcome	

Date:05/12/99ISR Number: 3260096-XReport Type:Expedited (15-DaCompany Report #A0080356  
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL Other		Angina Pectoris Angiopathy Blood Pressure Increased Cardiac Disorder Cardiac Murmur Dermatitis Oedema Peripheral Urticaria	Health Professional	Zyban Tablet-Zyban Dyazide	PS C		ORAL

Date:05/13/99ISR Number: 3261006-1Report Type:Direct  
Age:18 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 200MG BID ORAL		Grand Mal Convulsion		Wellbutrin	PS		ORAL

Date:05/13/99ISR Number: 3261242-4Report Type:Expedited (15-DaCompany Report #A0075934  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Dermatitis	Foreign	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dizziness	Health				
Hospitalization -		Pruritus	Professional				
TWICE PER DAY		Respiratory Disorder					
Initial or Prolonged		Urticaria					
/ORAL							

Date:05/13/99ISR Number: 3261243-6Report Type:Expedited (15-DaCompany Report #A0087951  
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Glucose Decreased	Foreign	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Convulsion	Health				
PER DAY/ ORAL		Fall	Professional	Humalog	C		
				Human Int/Long			
				Insulin	C		
				Short-Acting Insulin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/13/99ISR Number: 3261245-XReport Type:Expedited (15-DaCompany Report #A0090953  
 Age:16 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 15 GRAMS/ SINGLE DOSE/ ORAL	Overdose	Health Professional	Wellbutrin Tablet - Controlled Release	PS		ORAL

Date:05/13/99ISR Number: 3261328-4Report Type:Expedited (15-DaCompany Report #A0090907  
 Age:33 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ ORAL	Dermatitis Dyspnoea	Consumer	Zyban Tablet - Zyban	PS		ORAL
	Eating Disorder Face Oedema Tongue Oedema					

Date:05/13/99ISR Number: 3261329-6Report Type:Expedited (15-DaCompany Report #A0091460  
 Age:56 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG/ TWICE PER DAY/ ORAL	Dyspnoea Oedema	Foreign Health Professional	Zyban Tablet- Zyban Amitriptyline Hcl	PS C		ORAL

Date:05/13/99ISR Number: 3262492-3Report Type:Direct Company Report #  
 Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG PO BID		Chest Discomfort		Zyban	PS		ORAL
Initial or Prolonged		Chest Pain Pruritus Rash Generalised Urticaria					

Date:05/14/99ISR Number: 3262528-XReport Type:Direct  
 Age:43 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ONE TABLET  TWICE A DAY		Urticaria		Zyban 150mg	PS		

Date:05/14/99ISR Number: 3262553-9Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 1 TAB PO BID Intervention to Prevent Permanent Impairment/Damage		Crying  Panic Disorder Tremor		Wellbutrin Sr 150mg	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/14/99ISR Number: 3262569-2Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG BID PO		Dizziness		Wellbutrin	PS		ORAL
* STARTED BID		Ventricular Tachycardia					
DOSING				Betapace	C		

Date:05/14/99ISR Number: 3262717-4Report Type:Expedited (15-DaCompany Report #A0089203  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Arthralgia	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Chest Discomfort	Professional				
PER DAY/ ORAL		Disturbance In Attention					
		Memory Impairment					

Date:05/14/99ISR Number: 3262720-4Report Type:Expedited (15-DaCompany Report #A0091511  
 Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Confusional State	Foreign	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Dizziness	Consumer				
Initial or Prolonged		Mental Impairment					
PER DAY/ ORAL							

Date:05/14/99ISR Number: 3262724-1Report Type:Expedited (15-DaCompany Report #A0087656  
 Age:27 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other 1	Anaphylactic Shock	Study	Zyban Tablet - Zyban	PS	ORAL
TABLET/TWICE	Dizziness	Health			
PER DAY/ORAL	Dysphagia	Professional			
	Dyspnoea		Pseudoephedrine Hcl	C	
	Face Oedema				
	Gingivitis				
	Hypersensitivity				
	Hypoventilation				
	Influenza				
	Insomnia				
	Joint Swelling				
	Myalgia				
	Oedema Peripheral				
	Oral Intake Reduced				
	Pain In Extremity				
	Swelling				
	Tongue Oedema				
	Urticaria				
	Weight Decreased				

Date:05/14/99ISR Number: 3262742-3Report Type:Expedited (15-DaCompany Report #A0055443  
Age:49 YR Gender:Male I/FU:I

Outcome	PT	Report Source
Hospitalization -	Hepatic Enzyme Increased	Literature
Initial or Prolonged	Muscle Injury	Health
Other	Nasopharyngitis	Professional
	Rhabdomyolysis	Company

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Representative

Dose	Duration	Product	Role	Manufacturer	Route
150MG/TWICE		Wellbutrin Tablet-Controlled Release	PS		ORAL
PER DAY/ORAL		Glipizide	C		
		Metformin	C		
		Multivitamin	C		
		Insulin	C		
		Stool Softener	C		

Date:05/14/99ISR Number: 3262769-1Report Type:Expedited (15-DaCompany Report #A0086505  
Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Central Nervous System Stimulation Confusional State	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/THREE TIMES PER DA ORAL		Grand Mal Convulsion Illusion	Representative				
10 MG/PER DAY		Memory Impairment Sedation		Olanzapine (Formulation Unnown)	SS		
				Fluvoxamine Maleate Lithium Salt Propranolol Hydrochloride Lorazepam Thyroxine Sodium	C C C C C C		

Date:05/14/99ISR Number: 3262782-4Report Type:Expedited (15-DaCompany Report #A0092017  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Foreign	Zyban Tablet-Zyban	PS		ORAL
ORAL		Atrial Fibrillation Bronchospasm Tachycardia	Health Professional				

Date:05/14/99ISR Number: 3262784-8Report Type:Expedited (15-DaCompany Report #A0092018  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Foreign	Zyban Tablet - Zyban	PS		ORAL
ORAL		Atrial Fibrillation Bronchospasm Tachycardia	Health Professional				

Date:05/14/99ISR Number: 3262785-XReport Type:Expedited (15-DaCompany Report #A0091961  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/PER DAY/ORAL		Pruritus					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/14/99ISR Number: 3262787-3Report Type:Expedited (15-DaCompany Report #A0091658  
Age:46 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / PER	Anxiety	Foreign	Zyban Tablet - Zyban	PS		ORAL
Initial or Prolonged DAY / ORAL	Atrial Fibrillation	Health				
	Bronchospasm Tachycardia	Professional				

Date:05/14/99ISR Number: 3262789-7Report Type:Expedited (15-DaCompany Report #A0089238  
Age:21 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150 MG /	Arthralgia	Foreign	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY	Bronchospasm	Health				
/ ORAL	Conjunctival Disorder	Professional				
	Conjunctivitis Eyelid Oedema Face Oedema Myalgia Oedema Peripheral Serum Sickness Throat Tightness Urticaria					

Date:05/14/99ISR Number: 3264632-9Report Type:Direct Company Report #  
Age:67 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose 150MG BID	Hypoaesthesia		Bupropion	PS		
			Warfarin	C		
			Fosinopril	C		
			Simvoslatin	C		
			Caco3	C		

Date:05/17/99ISR Number: 3263494-3Report Type:Expedited (15-DaCompany Report #A0091793

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression	Foreign	Wellbutrin			
		Physical Assault	Health	Tablet-Controlled			
		Psychotic Disorder	Professional	Release	PS		
ORAL							

Date:05/17/99ISR Number: 3263495-5Report Type:Expedited (15-DaCompany Report #A0088811

Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Angina Pectoris	Foreign	Zyban Tablet -			
Other			Health	Zyban	PS		ORAL
150 MG/TWICE			Professional				
PER DAY/ORAL							
				Ramipril	C		
				Isosorbide	C		
				Metoprolol	C		
				Nicotinic Acid	C		
				Lorazepam	C		
				Nitroglycerin	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/17/99ISR Number: 3263802-3Report Type:Expedited (15-DaCompany Report #A0092189

Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE		Asthma	Foreign	Zyban	PS		ORAL
Initial or Prolonged PER DAY/ ORAL		Burning Sensation	Consumer				
		Eyelid Oedema Facial Palsy Hypersensitivity Pruritus					

Date:05/17/99ISR Number: 3263803-5Report Type:Expedited (15-DaCompany Report #A0092480

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Gait Disturbance	Foreign	Zyban	PS		ORAL
		Speech Disorder	Consumer				

Date:05/17/99ISR Number: 3263841-2Report Type:Expedited (15-DaCompany Report #A0092016

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Anxiety	Foreign	Zyban Tablet - Zyban	PS		ORAL
		Atrial Fibrillation Bronchospasm Tachycardia	Health Professional				

Date:05/18/99ISR Number: 3264987-5Report Type:Periodic Company Report #S99-USA-00106-01

Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 880 MG ONCE		Convulsion	Health	Celexa	PS		ORAL

Initial or Prolonged PO	Electrocardiogram Qt	Professional				
600 MG ONCE	Prolonged		Wellbutrin	SS		ORAL
PO	Intentional Misuse					
	Suicide Attempt		Ethanol	SS		

Date:05/19/99ISR Number: 3264878-XReport Type:Direct  
 Age:45 YR Gender:Male I/FU:I

Company Report #

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG 2 A DAY	Anxiety Apathy	Consumer	Zyban 150mg Glaxowellcome	PS	Glaxowellcome	
	Conversion Disorder Crying Depression Malaise					

Date:05/19/99ISR Number: 3264906-1Report Type:Direct  
 Age:43 YR Gender:Female I/FU:I

Company Report #

Outcome Dose Other	PT	Report Source	Product	Role	Manufacturer	Route
150MG BID	Dizziness Headache		Wellbutrin Sr 150mg Bid	PS		
	Influenza Like Illness Malaise Pyrexia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/19/99ISR Number: 3267142-8Report Type:Periodic  
 Age:36 YR Gender:Female I/FU:I

Company Report #8-99054-165A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 37.5 MG ONCE		Hypertension	Health	Effexor Xr Capsules	PS		ORAL
DAILY ORAL;		Tachycardia	Professional				
75 MG ONCE							
DAILY; 150 MG							
ONCE DAILY	1	WK		Wellbutrin (Bupropion)	SS		ORAL
450 MG DAILY							
DECREASED TO							
300 MG DAILY							
ORAL							

Date:05/19/99ISR Number: 3299510-2Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #S99-USA-00764-01

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 20 MG QD PO ;		Anorgasmia	Health	Celexa	PS		ORAL
10 MG QD PO		Nausea	Professional				
150 MG QD PO		Sedation		Wellbutrin	SS		ORAL

Date:05/20/99ISR Number: 3265559-9Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

100MG BID PO; Dizziness Wellbutrin PS ORAL  
 Ventricular Tachycardia  
 \* STARTED BID  
 DOSING Betapace C

Date:05/21/99ISR Number: 3267464-0Report Type:Expedited (15-DaCompany Report #A0085560  
 Age:15 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 3 GRAMS/ Initial or Prolonged SINGLE DOSE/ Disability ORAL Other		Convulsion Nausea Vomiting	Health Professional	Wellbutrin Tablet - Controlled Release	PS		ORAL

Date:05/21/99ISR Number: 3267728-0Report Type:Expedited (15-DaCompany Report #A0092017  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Anxiety Atrial Fibrillation Bronchospasm Tachycardia	Foreign Health Professional	Zyban Tablet-Zyban	PS		ORAL

Date:05/21/99ISR Number: 3267819-4Report Type:Expedited (15-DaCompany Report #A0091712  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Neuroleptic Malignant Syndrome	Health Professional	Wellbutrin (Formulation)			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL  
 Unknown) PS ORAL

Date:05/21/99ISR Number: 3267823-6Report Type:Expedited (15-DaCompany Report #A0092506  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous Metrorrhagia	Study Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL							

Date:05/24/99ISR Number: 3283646-6Report Type:Periodic Company Report #JAUSA-36367  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyskinesia	Health Professional	Propulsid (Cisapride), Janssen, Tablet 10 Mg	PS	Janssen	ORAL
10 MG 4 DAILY							
ORAL							
				Bupropion (Amfebutamone) Tablet 75 Mg	SS		ORAL
75 MG 3 DAILY							
ORAL							

Date:05/25/99ISR Number: 3269450-3Report Type:Expedited (15-DaCompany Report #A0085426  
 Age:16 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization - 3 GRAMS		Aspiration Convulsion	Health Professional	Wellbutrin Tablet- Controlled Release	PS		ORAL

Initial or Prolonged Intentional Misuse  
SINGLE DOSE  
Other Nausea  
ORAL Vomiting

Date:05/25/99ISR Number: 3269451-5Report Type:Expedited (15-DaCompany Report #A0092674  
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fatigue	Foreign	Zyban	PS		ORAL
150 MG TWICE		Hypersomnia	Consumer				
PER DAY ORAL		Insomnia					
		Sleep Disorder					
		Tachycardia					

Date:05/25/99ISR Number: 3269452-7Report Type:Expedited (15-DaCompany Report #A0092867  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blindness	Foreign	Zyban	PS		ORAL
ORAL			Consumer				



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/25/99ISR Number: 3269453-9Report Type:Expedited (15-DaCompany Report #A0089735

Age:27 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Zyban	PS		ORAL
150 MG TWICE							
PER DAY ORAL		Drug Interaction	Professional				
				Alprazolam (Formulation Unknown)	SS		ORAL
1 MG AS							
REQUIRED ORAL				Ranitidine Hydrochloride	C		

Date:05/25/99ISR Number: 3354072-6Report Type:Periodic Company Report #A0082509

Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Major Depression	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Nicotine	C		

Date:05/26/99ISR Number: 3270466-1Report Type:Expedited (15-DaCompany Report #A0092868

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Foreign	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY		Hypoaesthesia	Consumer				
/ ORAL		Paraesthesia					
		Sensation Of Heaviness					

Tremor

Date:05/26/99ISR Number: 3270469-7Report Type:Expedited (15-DaCompany Report #A0090923  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Alcohol Withdrawal	Foreign	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Syndrome	Health				
Hospitalization -		Coma	Professional				
TWICE PER DAY		Convulsion	Company				
Initial or Prolonged		Hypoxia	Representative				
/ ORAL		Liver Function Test					
		Abnormal					
		Pneumonia Aspiration					

Date:05/26/99ISR Number: 3270879-8Report Type:Direct Company Report #  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Coma	Health	Wellbutrin Sr 150 Mg			
Other		Grand Mal Convulsion	Professional	Bid	PS		ORAL
150 MG BID PO 3	MON	Urinary Incontinence		Ultram 50 Mg Bid	SS		ORAL
50 MG BID PO 3	MON			Seroquel	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/26/99ISR Number: 3272704-8Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #8-98271-019A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiovascular Disorder	Consumer	Redux	PS		ORAL
ORAL				Fastin (Phentermine)	SS		ORAL
ORAL				Pondimin (Fenfluramine) Tablet	SS		ORAL
ORAL				Pro-Fast (Phentermine)	SS		ORAL
ORAL				Welbutrin (Bupropion)	SS		ORAL
150 MG TWICE							
DAILY ORAL							

Date:05/27/99ISR Number: 3271216-5Report Type:Direct  
Age:43 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG PO BID 1 MON	Convulsion	Health	Zyban 150 Mg Po Bid	PS		ORAL
Initial or Prolonged		Dyskinesia	Professional	Amtriptyline	C		
		Saliva Altered		Effexer	C		
				Remeron	C		

Date:05/28/99ISR Number: 3272922-9Report Type:Expedited (15-DaCompany Report #A0091510  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG/TWICE	Angioneurotic Oedema	Foreign	Zyban Tablet- Zyban	PS		ORAL
Initial or Prolonged	PER DAY/ORAL	Dermatitis	Health				
Disability		Leukocytosis	Professional				

Obstructive Airways  
Disorder  
Respiratory Disorder  
Serum Sickness  
Swelling  
Urticaria

Date:05/28/99ISR Number: 3272927-8Report Type:Expedited (15-DaCompany Report #A0092789

Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL		Dermatitis	Foreign	Zyban Tablet - Zyban	PS		ORAL
		Dyspnoea	Health				
		Pruritus Speech Disorder Urticaria	Professional				

Date:05/28/99ISR Number: 3272929-1Report Type:Expedited (15-DaCompany Report #A0089661

Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Health	Wellbutrin Tablet-Controlled Release			
150 MG/TWICE PER DAY/ORAL			Professional		PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3273170-9Report Type:Expedited (15-DaCompany Report #A0093370

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blindness	Consumer	Zyban Tablet - Zyban	PS		

Date:05/28/99ISR Number: 3282842-1Report Type:Periodic Company Report #A0087404

Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain	Consumer	Zyban	PS		ORAL
		Face Oedema					
		Hypersensitivity					
		Urticaria					

Date:05/28/99ISR Number: 3282845-7Report Type:Periodic Company Report #A0088006

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Zyban	PS		ORAL
			Professional				
			Company Representative				

Date:05/28/99ISR Number: 3282847-0Report Type:Periodic Company Report #A0088853

Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Oedema	Consumer	Zyban	PS		ORAL
		Pruritus					

Urticaria

Triphasil

C

Date:05/28/99ISR Number: 3282849-4Report Type:Periodic Company Report #A0088991  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG/ UNK/	Anaphylactic Reaction	Health	Zyban	PS		ORAL
Hospitalization -	ORAL	Eyelid Oedema	Professional				
Initial or Prolonged		Face Oedema Pharyngeal Oedema					

Date:05/28/99ISR Number: 3282852-4Report Type:Periodic Company Report #A0089382  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG/ TWICE	Chest Pain	Consumer	Zyban	PS		ORAL
PER DAY/ ORAL		Ecchymosis	Company				
		Urticaria	Representative				

Date:05/28/99ISR Number: 3282854-8Report Type:Periodic Company Report #A0089444  
Age:13 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300 MG / SEE	Convulsion	Health	Zyban	PS		ORAL
TEXT/ ORAL		Overdose	Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3282857-3Report Type:Periodic Company Report #A0089660  
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG / PER	Convulsion	Consumer	Zyban	PS		ORAL
DAY/ ORAL							

Date:05/28/99ISR Number: 3282860-3Report Type:Periodic Company Report #A0090183  
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG/ TWICE	Eyelid Oedema	Health	Zyban	PS		ORAL
PER DAY/ ORAL							
			Company Representative	Tamoxifen	C		
				Alendronate Sodium	C		
				Lisinopril	C		

Date:05/28/99ISR Number: 3282861-5Report Type:Periodic Company Report #A0090576  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG/ UNK/	Dermatitis	Health	Zyban	PS		ORAL
Initial or Prolonged ORAL							
			Company Representative				

Date:05/28/99ISR Number: 3282863-9Report Type:Periodic Company Report #A0059506  
 Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Life-Threatening	Anorexia	Health	Zyban	PS	ORAL
150 MG /					
TWICE PER	Paranoia	Professional			
DAY/ ORAL	Suicide Attempt				

Date:05/28/99ISR Number: 3282864-0Report Type:Periodic Company Report #A0070204  
 Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fatigue	Consumer	Zyban	PS		ORAL
150 MG / PER		Headache					
DAY/ ORAL		Loss Of Consciousness					
		Nervousness					
		Pruritus					
		Pyrexia					
		Sedation					
		Tremor					

Date:05/28/99ISR Number: 3282865-2Report Type:Periodic Company Report #A0073851  
 Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Zyban	PS		ORAL
150 MG/ TWICE		Fall	Professional				
PER DAY/ ORAL		Hypoglycaemia		Glimepiride	C		
		Injury		Multivitamin	C		
		Tremor		Vitamin E	C		
				Ester-C	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Ampicillin C

Date:05/28/99ISR Number: 3282871-8Report Type:Periodic Company Report #A0076211  
 Age:30 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY / ORAL	Hypersensitivity Urticaria	Health Professional	Zyban	PS		ORAL

Date:05/28/99ISR Number: 3282873-1Report Type:Periodic Company Report #A0077336  
 Age:33 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY/ ORAL	Hypersensitivity Oedema Peripheral Rash Erythematous Urticaria Vomiting	Health Professional	Zyban Claritin - D Finasteride	PS C C		ORAL

Date:05/28/99ISR Number: 3282876-7Report Type:Periodic Company Report #A0077347  
 Age:63 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER Other DAY/ ORAL	Convulsion Syncope	Health Professional Company Representative	Zyban Oestradiol Salbutamol Sulphate	PS C C		ORAL

Theophylline C  
Allopurinol C  
Alendronate Sodium C

Date:05/28/99ISR Number: 3282880-9Report Type:Periodic Company Report #A0078443  
Age:23 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Capillary Leak Syndrome	Health	Zyban	PS		ORAL
150 MG/ UNK/		Hypersensitivity	Professional				
ORAL		Swelling	Company Representative	Omeprazole	C		

Date:05/28/99ISR Number: 3282885-8Report Type:Periodic Company Report #A0078477  
Age:32 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain Upper
Initial or Prolonged	Bone Pain
Other	Chills
	Diarrhoea
	Dizziness
	Dyspepsia
	Face Oedema
	Headache
	Hypersensitivity
	Pruritus

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pyrexia Rash Maculo-Papular Swelling	Report Source	Product	Role	Manufacturer	Route
150 MG /		Urticaria	Health	Zyban	PS		ORAL
TWICE PER		Vomiting	Professional				
DAY/ ORAL				Ibuprofen	C		

Date:05/28/99ISR Number: 3282888-3Report Type:Periodic Company Report #A0079702  
 Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Health	Zyban	PS		ORAL
Other		Syncope	Professional				
150 MG / WICE			Company	Cetirizine			
PER DAY/ ORAL			Representative	Hydrochloride	C		
				Fluoxymesterone	C		

Date:05/28/99ISR Number: 3282890-1Report Type:Periodic Company Report #A0080413  
 Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decreased Appetite	Health	Zyban	PS		ORAL
Hospitalization -		Disturbance In Attention	Professional				
150 MG /		Dysgeusia					
Initial or Prolonged		Hallucination		Naproxen Sodium	C		
TWICE PER		Headache					
Other		Insomnia					
DAY/ ORAL							

Date:05/28/99ISR Number: 3282892-5Report Type:Periodic  
Age:44 YR Gender:Female I/FU:F

Company Report #A0080693

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hyperglycaemia	Health	Zyban	PS		ORAL
150 MG /			Professional				
UNK/ORAL							

Choline	
Theophyllinate	C
Gemfibrozil	C
Atenolol	C
Human Int / Long	
Insulin	C
Human Short-Act.	
Insulin	C
Frusemide	C
Lansoprazole	C

Date:05/28/99ISR Number: 3282900-1Report Type:Periodic  
Age:20 YR Gender:Male I/FU:F

Company Report #A0080807

Outcome	PT
Hospitalization -	Burning Sensation
Initial or Prolonged	Dry Skin
	Dyspnoea
	Hypersensitivity
	Pruritus
	Rash Erythematous
	Rash Papular

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Freedom Of Information (FOI) Report

		Skin Irritation Urticaria	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Health	Zyban	PS		ORAL
150 MG /			Professional				
TWICE PER							
DAY/ ORAL							

Date:05/28/99ISR Number: 3282903-7Report Type:Periodic Company Report #A0081611  
 Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Health	Zyban	PS		ORAL
Hospitalization -		Throat Tightness	Professional				
150 MG /		Urticaria					
Initial or Prolonged				Weight Loss			
TWICE PER				Medication	C		
DAY/ ORAL							

Date:05/28/99ISR Number: 3282909-8Report Type:Periodic Company Report #A0081603  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharyngeal Oedema	Health	Zyban Tablet - Zyban	PS		ORAL
Other		Tongue Oedema	Professional				
150 MG /		Urticaria		Cyclobenzaprin Hcl			
TWICE PER DAY				Tablet	SS		ORAL
10 MG / THREE							
TIMES PER DA				Armour Thyroid	C		
				Naproxen	C		

Date:05/28/99ISR Number: 3282912-8Report Type:Periodic Company Report #A0082017  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Tremor					
DAY							

Date:05/28/99ISR Number: 3282914-1Report Type:Periodic Company Report #A0082065  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Face Oedema	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Mydriasis					
TWICE PER DAY		Nervousness					
		Oedema					
		Oedema Peripheral					
		Pruritus					
		Urticaria					

Date:05/28/99ISR Number: 3282917-7Report Type:Periodic Company Report #A0082357  
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Aggression	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Grand Mal Convulsion	Professional				
Hospitalization -		Loss Of Consciousness	Company	Simvastatin	C		
TWICE PER DAY		Mood Swings	Representative	Nicotine	C		
Initial or Prolonged							

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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3282920-7Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0082921

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban Tablet - Zyban	PS		ORAL
Other		Dermatitis	Professional				
150 MG /		Pruritus					
TWICE PER DAY		Swelling					

Date:05/28/99ISR Number: 3282921-9Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0083363

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban Tablet - Zyban	PS		ORAL
Hospitalization -		Convulsion	Professional				
150 MG			Company				
Initial or Prolonged			Representative				

Date:05/28/99ISR Number: 3282923-2Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #A0083631

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet - Zyban	PS		ORAL
Other		Pruritus					
150 MG /		Urticaria					
TWICE PER DAY				Birth Control	C		

Date:05/28/99ISR Number: 3282925-6Report Type:Periodic  
Age:59 YR Gender:Male I/FU:I

Company Report #A0083648

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet -			
Hospitalization -		Dermatitis		Zyban	PS		ORAL
Initial or Prolonged		Dyspnoea					
150MG ORAL							

Hypersensitivity

Date:05/28/99ISR Number: 3282927-XReport Type:Periodic Company Report #A0083649  
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
Initial or Prolonged		Dyspnoea Hypersensitivity					

Date:05/28/99ISR Number: 3282929-3Report Type:Periodic Company Report #A0084374  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG /		Convulsion	Health Professional	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY			Company Representative				

Date:05/28/99ISR Number: 3282930-XReport Type:Periodic Company Report #A0084412  
 Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG /		Convulsion	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vicodin C

Date:05/28/99ISR Number: 3282932-3Report Type:Periodic Company Report #A0084629  
 Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY		Myocardial Infarction	Health Professional	Zyban Tablet - Zyban	PS		ORAL
			Company Representative	Colestipol Pravastatin Sodium	C C		

Date:05/28/99ISR Number: 3282934-7Report Type:Periodic Company Report #A0084862  
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG / TWICE PER DAY		Convulsion	Study	Zyban Tablet - Zyban	PS		ORAL
		Hyperhidrosis	Consumer				
		Nasopharyngitis Pallor Tremor					

Date:05/28/99ISR Number: 3282936-0Report Type:Periodic Company Report #A0085003  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG / TWICE PER DAY		Anxiety	Health	Zyban Tablet - Zyban	PS		ORAL
		Convulsion	Professional				
		Insomnia Myalgia Nervousness Tremor					

Date:05/28/99ISR Number: 3282938-4Report Type:Periodic Company Report #A0085850  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Urticaria	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							

TWICE PER DAY

Date:05/28/99ISR Number: 3282941-4Report Type:Periodic Company Report #A0086000  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anaphylactic Reaction	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							

Initial or Prolonged Dermatitis

Date:05/28/99ISR Number: 3282943-8Report Type:Periodic Company Report #A0086021  
Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Dermatitis
Initial or Prolonged	Dyspnoea
	Face Oedema
	Oedema
	Tongue Oedema

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Freedom Of Information (FOI) Report

Urticaria

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG /		Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY			Claritin-D	C		
			Conjugated Estrogens	C		
			Medroxyprogesterone			
			Ace.	C		

Date:05/28/99ISR Number: 3282945-1Report Type:Periodic Company Report #A0086086  
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Arthralgia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dermatitis					
Initial or Prolonged		Pruritus					
TWICE PER DAY							

Date:05/28/99ISR Number: 3282947-5Report Type:Periodic Company Report #A0086828  
 Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / SEE		Influenza Like Illness					
TEXT		Pharyngeal Oedema		Nocotine	C		
		Urticaria					

Date:05/28/99ISR Number: 3282950-5Report Type:Periodic Company Report #A0087268  
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Myocardial Infarction Consumer Zyban Tablet - Zyban PS ORAL  
1500 MG /  
Initial or Prolonged Sinus Congestion  
TWICE PER

DAY

Simvastatin C  
Lisinopril C  
Metoprolol C  
Nitroglycerin C

Date:05/28/99ISR Number: 3332413-3Report Type:Periodic Company Report #A0086568  
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Accommodation Disorder	Consumer	Zyban Tablet - Zyban	PS		ORAL
PER DAY /		Dizziness					
ORAL		Hypertension					
		Muscle Contractions Involuntary Nausea Sensation Of Pressure		Ranitidine Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332414-5Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0086587

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ PER		Pruritus					
DAY / ORAL				Thyroid Medication	C		
				Prempro	C		
				Erythromycin	C		

Date:05/28/99ISR Number: 3332415-7Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0086696

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Balance Disorder	Consumer	Zyban Tablet- Zyban	PS		ORAL
150 MG/ TWICE		Insomnia					
PER DAY/ ORAL		Nausea		Excedrin	C		

Date:05/28/99ISR Number: 3332416-9Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #A0086698

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:05/28/99ISR Number: 3332417-0Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #A0086700

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperglycaemia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							

DAY / ORAL

Glibenclamide C  
Acarbose C  
Lisinopril C

Date:05/28/99ISR Number: 3332418-2Report Type:Periodic Company Report #A0086701  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Flatulence					
PER DAY/ ORAL		Insomnia		Prempro	C		
		Tachycardia		Pravastatin Sodium	C		
				Multivitamin	C		

Date:05/28/99ISR Number: 3332419-4Report Type:Periodic Company Report #A0086702  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Tachycardia					
TWICE PER		Tremor					
DAY/ ORAL				Cyclobenzaprine Hcl	C		
				Ibuprofen	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332420-0Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #A0086703

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							
DAY / ORAL							

Ibuprofen C

Date:05/28/99ISR Number: 3332421-2Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0086704

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Generalised	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							
DAY/ ORAL							

Multivitamin C

Date:05/28/99ISR Number: 3332422-4Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #A0086705

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dry Mouth					
TWICE PER		Dyspnoea					
DAY/ ORAL		Insomnia					
		Tachycardia					

Date:05/28/99ISR Number: 3332423-6Report Type:Periodic  
Age:72 YR Gender:Male I/FU:I

Company Report #A0086706

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG/ PER	Weight Decreased	Consumer	Zyban Tablet - Zyban	PS	ORAL
DAY/ ORAL			Blood Pressure Medication	C	
			Nifedipine	C	

Date:05/28/99ISR Number: 3332424-8Report Type:Periodic Company Report #A0086708  
 Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Epistaxis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Headache					
PER DAY/ ORAL		Irritability					

Date:05/28/99ISR Number: 3332425-XReport Type:Periodic Company Report #A0086709  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Diarrhoea	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE				Ibuprofen	C		
PER DAY/ ORAL				Baclofen	C		
				Diazepam	C		
				Alendronate Sodium	C		
				Multivitamin	C		
				Iron Salt	C		
				Calcium Salt	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Folic Acid C

Date:05/28/99ISR Number: 3332426-1Report Type:Periodic Company Report #A0086710  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY /							
ORAL							

Acyclovir C

Date:05/28/99ISR Number: 3332427-3Report Type:Periodic Company Report #A0086711  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Urticaria					
DAY / ORAL							

Date:05/28/99ISR Number: 3332428-5Report Type:Periodic Company Report #A0086712  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Headache					
VARIABLE DOSE		Hepatitis C					
/ ORAL		Muscle Twitching		Trazodone	C		
		Nausea		Clonazepam	C		
				Ranitidine			
				Hydrochloride	C		
				Naltrexone			

Date:05/28/99ISR Number: 3332429-7Report Type:Periodic  
Age:51 YR Gender:Male I/FU:I

Company Report #A0086713

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /	PER						
DAY / ORAL							

Date:05/28/99ISR Number: 3332430-3Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #A0086714

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3332431-5Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0086715

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet- Zyban	PS		ORAL
150 MG /		Amnesia					
TWICE PER DAY							
/ ORAL							
		Drug Ineffective					
		Smoker		Multivitamin	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332432-7Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0086716

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Dry Mouth	Professional				
DAY / ORAL		Eye Movement Disorder					

Date:05/28/99ISR Number: 3332433-9Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0087215

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Haematemesis	Professional				
TWICE PER DAY		Mood Swings					
/ ORAL							

Date:05/28/99ISR Number: 3332435-2Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #A0087216

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Mydriasis					
TWICE PER DAY							
/ ORAL							

Levofloxacin C  
Prednisone C  
Salbutamol Sulphate C

Date:05/28/99ISR Number: 3332436-4Report Type:Periodic  
Age:29 YR Gender:Male I/FU:I

Company Report #A0087246

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / AT NIGHT/		Dermatitis Drug Hypersensitivity Pruritus Urticaria	Health Professional	Zyban Tablet - Zyban	PS		

Date:05/28/99ISR Number: 3332438-8Report Type:Periodic Company Report #A0087256  
 Age:38 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL		Chest Pain Flatulence Gastrointestinal Disorder	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3332439-XReport Type:Periodic Company Report #A0087282  
 Age:46 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL		Asthenia Drug Hypersensitivity Hypoaesthesia Joint Stiffness Lacrimation Increased Neurological Symptom	Health Professional	Zyban Tablet - Zyban	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332440-6Report Type:Periodic  
 Age:50 YR Gender:Male I/FU:I

Company Report #A0087313

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Erectile Dysfunction	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER		Insomnia					
DAY/ ORAL				Amlodipine	C		

Date:05/28/99ISR Number: 3332442-XReport Type:Periodic  
 Age:39 YR Gender:Male I/FU:I

Company Report #A0087314

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Headache	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER							
DAY/ ORAL							

Date:05/28/99ISR Number: 3332443-1Report Type:Periodic  
 Age:40 YR Gender:Male I/FU:I

Company Report #A0087315

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Toxicologic Test Abnormal	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER							
DAY/ ORAL							

Date:05/28/99ISR Number: 3332444-3Report Type:Periodic  
 Age:47 YR Gender:Male I/FU:I

Company Report #A0087316

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Headache					
TWICE PER		Irritability					
DAY/ ORAL		Malaise					

Date:05/28/99ISR Number: 3332446-7Report Type:Periodic Company Report #A0087317  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ill-Defined Disorder	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Influenza Like Illness					

Date:05/28/99ISR Number: 3332447-9Report Type:Periodic Company Report #A0087318  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER							
DAY/ ORAL							

Date:05/28/99ISR Number: 3332448-0Report Type:Periodic Company Report #A0087319  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Irritability	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER							
DAY/ ORAL							



150 MG / Pruritus Consumer Zyban Tablet - Zyban PS ORAL  
TWICE PER DAY Urticaria  
/ ORAL

Date:05/28/99ISR Number: 3332453-4Report Type:Periodic Company Report #A0087323  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Prempro C  
Ziac C

Date:05/28/99ISR Number: 3332455-8Report Type:Periodic Company Report #A0087324  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Diarrhoea					
TWICE PER		Sedation					
DAY/ ORAL				Ascorbic Acid	C		





150 MG / ORAL  
Confusional State  
Consumer  
Zyban Tablet - Zyban PS  
ORAL  
Dizziness  
Headache  
Insomnia  
Nausea

Date:05/28/99ISR Number: 3332462-5Report Type:Periodic Company Report #A0087728  
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Constipation					
TWICE PER DAY		Hyperhidrosis					
/ ORAL		Insomnia					

Date:05/28/99ISR Number: 3332463-7Report Type:Periodic Company Report #A0087729  
Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL	7 DAY						
				Diltiazem			
				Hydrochloride	C		
				Digoxin	C		
				Gemfibrozil	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332465-0Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0087753

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Insomnia	Professional				
DAY / ORAL							

Date:05/28/99ISR Number: 3332466-2Report Type:Periodic  
 Age:36 YR Gender:Female I/FU:I

Company Report #A0087754

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Depression					
TWICE PER DAY		Disturbance In Attention					
/ ORAL		Drug Ineffective					
		Emotional Disorder					
		Fear					
		Nervousness					

Date:05/28/99ISR Number: 3332468-6Report Type:Periodic  
 Age:71 YR Gender:Female I/FU:I

Company Report #A0087755

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER							
DAY/ ORAL							
				Fluvastatin Sodium	C		
				Ketoprofen	C		

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Abnormal Dreams	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER		Back Pain					
DAY/ ORAL		Dry Skin					
		Fatigue		Pravastatin Sodium	C		
		Myalgia		Allopurinol	C		
		Neck Pain		Aspirin	C		
		Nightmare		Calcium Salt	C		
				Multivitamin	C		

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Constipation	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Diarrhoea					
/ ORAL				Amlodipine	C		
				Spiroinolactone	C		
				Hydrazaline			
				Hydrochloride	C		
				Nifedipine	C		
				Dilantin	C		
				Potassium Chloride	C		
				Aspirin	C		
				Finasteride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Frusemide C  
Prednisone C

Date:05/28/99ISR Number: 3332472-8Report Type:Periodic Company Report #A0087758  
Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL	5 DAY	Constipation Dry Mouth Dyspnoea	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3332473-XReport Type:Periodic Company Report #A0087759  
Age:57 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL

Nifedipine C  
Thyroxine Sodium C  
Atorvastatin Calcium C  
Hormones C  
Aspirin C

Date:05/28/99ISR Number: 3332474-1Report Type:Periodic Company Report #A0087760  
Age:42 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Dyspnoea	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3332475-3Report Type:Periodic Company Report #A0087761  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG / PER		Blood Pressure Increased	Consumer	Zyban Tablet - Zyban	PS		ORAL
DAY/ ORAL		Dissociation					
		Dizziness					
		Malaise					

Date:05/28/99ISR Number: 3332476-5Report Type:Periodic Company Report #A0087762  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER							
DAY/ ORAL							

Date:05/28/99ISR Number: 3332478-9Report Type:Periodic Company Report #A0087763  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332479-0Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0087764

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL		Anorexia Dermatitis Disturbance In Attention Dizziness Folliculitis Pruritus Rash Erythematous Weight Decreased	Health Professional	Zyban Tablet - Zyban  Nordette	PS  C		ORAL

Date:05/28/99ISR Number: 3332481-9Report Type:Periodic  
Age:73 YR Gender:Female I/FU:I

Company Report #A0087765

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Smoker	Consumer	Zyban Tablet - Zyban  Valsartan Sulphasalazine	PS  C C		ORAL

Date:05/28/99ISR Number: 3332482-0Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #A0087766

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Accommodation Disorder Headache	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3332483-2Report Type:Periodic Company Report #A0087329  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Vision Blurred					
DAY/ ORAL				Ortho Tri-Cyclen	C		

Date:05/28/99ISR Number: 3332484-4Report Type:Periodic Company Report #A0087767  
Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Constipation	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Drug Ineffective Smoker	Professional				

Date:05/28/99ISR Number: 3332486-8Report Type:Periodic Company Report #A0087768  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Burning Sensation	Consumer	Zyban	PS		ORAL
150 MG / ORAL		Dermatitis					





150 MG /  
TWICE PER DAY  
/ ORAL

Dizziness

Consumer

Zyban Tablet - Zyban PS

ORAL

Date:05/28/99ISR Number: 3332493-5Report Type:Periodic Company Report #A0087333  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pain					
TWICE PER DAY							
/ ORAL				Diazepam	C		

Date:05/28/99ISR Number: 3332495-9Report Type:Periodic Company Report #A0087334  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypertonia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY		Tremor					
/ ORAL				Diazepam	C		
				Clonidine	C		
				Dicyclomine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332496-0Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #A0087335

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dermatitis Contact Lichenification Pruritus Rash Erythematous Rash Papular	Health Professional	Zyban Tablet - Zyban   Triamcinolone Acetonide Oxymetazoline Hcl Clindamycin Phosphate	PS   C C C		ORAL

Date:05/28/99ISR Number: 3332497-2Report Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #A0087336

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Dermatitis Exfoliative Dry Mouth Insomnia Myalgia Oedema Peripheral Pallor	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3332500-XReport Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0087337

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Insomnia Irritability	Consumer	Zyban Tablet - Zyban  Estrogen	PS  C		ORAL

Date:05/28/99ISR Number: 3332501-1Report Type:Periodic  
 Age:67 YR Gender:Female I/FU:I

Company Report #A0087338

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Anxiety					
TWICE PER DAY		Crying					
/ ORAL		Depression		Alendronate Sodium	C		
		Disturbance In Attention		Atorvastatin Calcium	C		
		Insomnia					
		Nervousness					
		Phobia					
		Tremor					
		Weight Decreased					

Date:05/28/99ISR Number: 3332504-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0087339

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							
DAY / ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332505-9Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #A0087340

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Abdominal Distension	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Micturition Urgency					
/ ORAL		Nervousness					
		Tinnitus		Alprazolam	C		
				Inhaler	C		
				Omeprazole	C		

Date:05/28/99ISR Number: 3332512-6Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #A0087341

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Pruritus					
/ ORAL				Alprazolam	C		
				Aspirin	C		

Date:05/28/99ISR Number: 3332515-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0087343

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG / ORAL	4 DAY	Palpitations	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3332517-5Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #A0087344

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Insomnia					
/ ORAL		Pruritus					
		Urticaria		Clonazepam	C		

Date:05/28/99ISR Number: 3332518-7Report Type:Periodic Company Report #A0087345  
 Age:50 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Myalgia	Health	Zyban Tablet - Zyban	PS		ORAL
		Sleep Disorder	Professional	Multivitamin	C		

Date:05/28/99ISR Number: 3332519-9Report Type:Periodic Company Report #A0087346  
 Age:49 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Drug Dependence	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Headache					
		Myalgia					
		Neck Pain					
		Paraesthesia					
		Sinus Headache					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332520-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0087347

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Depression					

Date:05/28/99ISR Number: 3332522-9Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0087348

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/ORAL		Professional					

Date:05/28/99ISR Number: 3332524-2Report Type:Periodic  
 Age:45 YR Gender:Male I/FU:I

Company Report #A0087349

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Decreased Appetite					
TWICE PER DAY		Feeling Drunk					
/ ORAL		Insomnia					
		Judgement Impaired					

Date:05/28/99ISR Number: 3332526-6Report Type:Periodic  
 Age:37 YR Gender:Female I/FU:I

Company Report #A0087350

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							
DAY / ORAL							

Date:05/28/99ISR Number: 3332527-8Report Type:Periodic Company Report #A0087351  
 Age:56 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Disturbance In Attention Dizziness Dry Mouth Sleep Disorder Syncope Tremor	Health Professional	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3332529-1Report Type:Periodic Company Report #A0087352  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / UNK / ORAL		Anxiety Dry Mouth Dyspnoea Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332530-8Report Type:Periodic  
 Age:47 YR Gender:Male I/FU:I

Company Report #A0087353

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Condition Aggravated					
TWICE PER DAY		Dry Mouth					
/ ORAL		Dysgeusia		Diazepam	C		
		Dysphagia		Unknown	C		
		Oesophageal Spasm		Diltiazem			
				Hydrochloride	C		

Date:05/28/99ISR Number: 3332531-XReport Type:Periodic  
 Age:34 YR Gender:Female I/FU:I

Company Report #A0087411

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Excitability					
TWICE PER DAY		Insomnia					
/ ORAL							

Date:05/28/99ISR Number: 3332532-1Report Type:Periodic  
 Age:50 YR Gender:Female I/FU:I

Company Report #A0087412

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dissociation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3332533-3Report Type:Periodic Company Report #A0087461  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK / UNK /		Insomnia	Consumer	Zyban Tablet - Zyban	PS		
UNK		Night Sweats					
		Throat Tightness		Amoxicillin	C		

Date:05/28/99ISR Number: 3332535-7Report Type:Periodic Company Report #A0087466  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
DAY / ORAL		Depression					
		Suicidal Ideation		Ranitidine Hydrochloride	C		

Date:05/28/99ISR Number: 3332536-9Report Type:Periodic Company Report #A0087597  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / UNK		Balance Disorder	Health	Zyban Tablet - Zyban	PS		ORAL
/ ORAL		Dizziness	Professional				
		Gait Disturbance	Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332537-0Report Type:Periodic  
 Age:50 YR Gender:Female I/FU:I

Company Report #A0087666

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Change Of Bowel Habit	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Flatulence					
TWICE PER DAY							
/ ORAL				Pravastatin Sodium	C		

Date:05/28/99ISR Number: 3332540-0Report Type:Periodic  
 Age:55 YR Gender:Male I/FU:I

Company Report #A0087669

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flatulence	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Gastrointestinal Disorder					
TWICE PER DAY							
/ ORAL				Atenolol	C		
				Simvastatin	C		

Date:05/28/99ISR Number: 3332542-4Report Type:Periodic  
 Age:52 YR Gender:Female I/FU:I

Company Report #A0087719

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Hormones	C		

Date:05/28/99ISR Number: 3332544-8Report Type:Periodic Company Report #A0087720  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3332546-1Report Type:Periodic Company Report #A0087721  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Generalised	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / UNK							
/ ORAL							
Urticaria							
Ortho-Cept							
Multivitamin							
C							
C							

Date:05/28/99ISR Number: 3332548-5Report Type:Periodic Company Report #A0087722  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menstruation Irregular	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / UNK							
/ ORAL							
8 WK							
Pruritus							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332550-3Report Type:Periodic  
 Age:44 YR Gender:Female I/FU:I

Company Report #A0087723

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL				Thyroid	C		

Date:05/28/99ISR Number: 3332552-7Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #A0087724

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Swelling					
TWICE PER DAY		Urticaria					
/ ORAL							

Date:05/28/99ISR Number: 3332555-2Report Type:Periodic  
 Age:29 YR Gender:Female I/FU:I

Company Report #A0087725

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cold Sweat	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Difficulty In Walking					
TWICE PER DAY		Disturbance In Attention					
ORAL		Malaise					
		Speech Disorder					
		Tremor					

Date:05/28/99ISR Number: 3332556-4Report Type:Periodic Company Report #A0087726  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / UNK			Health				
/ ORAL			Professional				
			Other				

Date:05/28/99ISR Number: 3332560-6Report Type:Periodic Company Report #A0086410  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cough	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG		Fluid Retention					
TWICE PER DAY		Hypersensitivity					
ORAL		Lacrimation Increased		Zinc Salt	C		
		Nasopharyngitis		Vitamin B Complex	C		
		Pulmonary Congestion		Ascorbic Acid	C		
		Sneezing		Vitamin E	C		
				Thyroxine Sodium	C		
				Estrogen	C		

Date:05/28/99ISR Number: 3332561-8Report Type:Periodic Company Report #A0086411  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG							
TWICE PER DAY							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL

Oral Contraceptive C

Date:05/28/99ISR Number: 3332563-1Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0086412

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
PER DAY		Insomnia					
ORAL		Major Depression					
		Tremor					

Date:05/28/99ISR Number: 3332565-5Report Type:Periodic  
Age:21 YR Gender:Female I/FU:I

Company Report #A0086413

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG		Gingival Bleeding	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							

ORAL

Oral Contraceptive C

Date:05/28/99ISR Number: 3332567-9Report Type:Periodic  
Age:59 YR Gender:Female I/FU:I

Company Report #A0086414

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Insomnia					

ORAL

Estrogen C

Calcium Salt

C

Date:05/28/99ISR Number: 3332568-0Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0086415

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG							
TWICE PER DAY							
ORAL							

Thyroxine Sodium C

Date:05/28/99ISR Number: 3332570-9Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0086416

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG		Nausea					
TWICE PER DAY							
ORAL							

Date:05/28/99ISR Number: 3332583-7Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #A0086417

Outcome	PT
	Dermatitis
	Dysphagia
	Dysuria
	Ecchymosis



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Pharyngeal Oedema					
		Pruritus					
		Urinary Tract Infection	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG							
TWICE PER DAY							
ORAL							

Ibuprofen C

Date:05/28/99ISR Number: 3332585-0Report Type:Periodic Company Report #A0086418  
 Age:49 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
Dose		Insomnia					
150 MG		Personality Change					
TWICE PER DAY							
ORAL							

Date:05/28/99ISR Number: 3332586-2Report Type:Periodic Company Report #A0086419  
 Age:48 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Tinnitus	Consumer	Zyban Tablet - Zyban	PS		ORAL
Dose							
150 MG							
TWICE PER DAY							
ORAL							

Date:05/28/99ISR Number: 3332589-8Report Type:Periodic Company Report #A0086420  
 Age:70 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose							

150 MG Abdominal Distension Consumer Zyban Tablet - Zyban PS ORAL  
 TWICE PER DAY Anorexia  
 ORAL Gastric Dilatation  
 Oedema Peripheral  
 Weight Increased

Date:05/28/99ISR Number: 3332591-6Report Type:Periodic Company Report #A0086421  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG		Dermatitis	Consumer	Zyban Tablet - Zyban PS			ORAL
ORAL			Company Representative				

Date:05/28/99ISR Number: 3332593-XReport Type:Periodic Company Report #A0086422  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG		Pruritus	Health	Zyban Tablet - Zyban PS			ORAL
ORAL			Professional				

Date:05/28/99ISR Number: 3332597-7Report Type:Periodic Company Report #A0086423  
 Age: Gender: I/FU:I

Outcome	PT	Report Source
	Tremor	Health Professional

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Company Representative	Product	Role	Manufacturer	Route
150 MG			Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY						
ORAL						

Date:05/28/99ISR Number: 3332599-0Report Type:Periodic Company Report #A0086424  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG		Dermatitis					
Local Swelling							
ORAL							

Date:05/28/99ISR Number: 3332600-4Report Type:Periodic Company Report #A0086425  
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG		Pruritus					
TWICE PER DAY							
ORAL							

Date:05/28/99ISR Number: 3332603-XReport Type:Periodic Company Report #A0086426  
 Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG		Drug Ineffective					

TWICE PER DAY

ORAL

Date:05/28/99ISR Number: 3332606-5Report Type:Periodic Company Report #A0086427  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG		Diarrhoea	Professional				
ORAL		Headache		Nicotine Patch	SS		
TRANSDERMAL	TRANSDERMAL	Therapeutic Response		Nicotine Gum	SS		ORAL
2 MG	AS	Unexpected					
REQUIRED							
ORAL				Inhaler	C		
				Estropipate	C		

Date:05/28/99ISR Number: 3332607-7Report Type:Periodic Company Report #A0086428  
Age:75 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG	PER						
DAY	ORAL			Methyldopa	C		
				Diltiazem			
				Hydrochloride	C		
				M.V.I.	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332610-7Report Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #A0086541

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 TABLET PER DAY ORAL		Blood Pressure Decreased Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
				Multivitamin	C		

Date:05/28/99ISR Number: 3332614-4Report Type:Periodic  
Age:25 YR Gender:Female I/FU:I

Company Report #A0084833

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL		Urticaria	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3332616-8Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #A0084834

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Dermatitis Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3332617-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0084835

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ ORAL		Hyperphagia Weight Increased	Consumer	Zyban Tablet - Zyban Purified Water	PS C		ORAL

Date:05/28/99ISR Number: 3332620-XReport Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #A0084836

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/ ORAL			Professional Other	Prednisone (Formulation Unknown)	SS		
				Nicotine Patch	SS		
				Pseudoephedrine (Formulation Unknown)	SS		

Date:05/28/99ISR Number: 3332624-7Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0084837

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ PER DAY/ ORAL		Influenza					
				Theraflu	C		
				Diazepam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332625-9Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0084838

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menopause	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Oligomenorrhoea					
PER DAY/ ORAL				Birth Control	C		

Date:05/28/99ISR Number: 3332629-6Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #A0084841

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Condition Aggravated					
PER DAY/ ORAL		Crying		Conj.			
		Decreased Activity		Estrogen+Medroxyprog	C		
		Depression		Multivitamin	C		
		Insomnia					
		Lethargy					
		Nervous System Disorder					
		Tremor					

Date:05/28/99ISR Number: 3332631-4Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0084842

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Migraine	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Nasopharyngitis					
PER DAY/ ORAL		Nausea					
		Productive Cough					

Date:05/28/99ISR Number: 3332634-XReport Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0084843

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hair Texture Abnormal		Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Muscle Spasms					
PER DAY/ORAL		Tongue Ulceration		Lotrel	C		
				Omeprazole	C		
				Vitamin E	C		
				Calcium Salt	C		
				Garlic	C		
				Hypericum	C		
				Kava	C		
				Hyoscyamine Sulphate	C		
				Metronidazole	C		
				Estrogen	C		
				Allergy Medication	C		
				Trimethoprim	C		

Date:05/28/99ISR Number: 3332637-5Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0084844

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Cough					
PER DAY/ORAL		Dyspnoea		Naproxen Sodium	C		
		Fatigue		Cyanocobalamin	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332640-5Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0084845

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Fatigue					
PER DAY/ORAL		Increased Appetite					

Date:05/28/99ISR Number: 3332642-9Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0084846

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sinusitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL							

Date:05/28/99ISR Number: 3332645-4Report Type:Periodic  
Age:62 YR Gender:Male I/FU:I

Company Report #A0084847

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/PER		Pruritus					
DAY/ORAL		Tremor		Multivitamin Zinc Salt	C C		

Date:05/28/99ISR Number: 3332648-XReport Type:Periodic  
Age:54 YR Gender:Male I/FU:I

Company Report #A0084848

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE							

PER DAY/ORAL

Date:05/28/99ISR Number: 3332649-1Report Type:Periodic Company Report #A0084849  
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Drug Withdrawal Syndrome					
		Irritability					
		Nervousness					

PER DAY/ORAL

Date:05/28/99ISR Number: 3332654-5Report Type:Periodic Company Report #A0084850  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE							
				Cholestyramine	C		
				Diazepam	C		
				Trifluoperazine	C		

PER DAY/ORAL

Date:05/28/99ISR Number: 3332656-9Report Type:Periodic Company Report #A0084851  
Age:47 YR Gender:Male I/FU:I

Outcome	PT
	Anxiety
	Disturbance In Attention

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Insomnia Stress				
Dose	Duration		Report Source	Product	Role	Manufacturer
			Consumer	Zyban Tablet - Zyban	PS	
150 MG/TWICE						ORAL
PER DAY/ORAL						
				Etodolac	C	
				Allegra-D	C	
				Tramadol		
				Hydrochloride	C	

Date:05/28/99ISR Number: 3332657-0Report Type:Periodic Company Report #A0084852  
 Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Escherichia Infection					
PER DAY/ORAL							
				Prempro	C		
				Ranitidine			
				Hydrochloride	C		
				Thyroxine Sodium	C		

Date:05/28/99ISR Number: 3332660-0Report Type:Periodic Company Report #A0084853  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ ORAL		Pruritus					

Date:05/28/99ISR Number: 3332681-8Report Type:Periodic Company Report #A0084854  
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG/TWICE                      Dermatitis                      Health                      Zyban Tablet - Zyban   PS                      ORAL  
PER DAY/ ORAL                      Professional

Date:05/28/99ISR Number: 3332685-5Report Type:Periodic                      Company Report #A0084855  
Age:35 YR   Gender:Male                      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL

150 MG/TWICE  
PER DAY/ORAL

Date:05/28/99ISR Number: 3332686-7Report Type:Periodic                      Company Report #A0084856  
Age:28 YR   Gender:Male                      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL

150 MG/TWICE  
PER DAY/ORAL   3   WK

Date:05/28/99ISR Number: 3332688-0Report Type:Periodic                      Company Report #A0084857  
Age:55 YR   Gender:Male                      I/FU:I

Outcome                      PT  
                                  Insomnia  
                                  Pruritus

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Rash Pruritic Swelling Urticaria	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/TWICE							
PER DAY/ORAL							

Date:05/28/99ISR Number: 3332690-9Report Type:Periodic Company Report #A0084858  
 Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/TWICE							
PER DAY/ORAL							

Nicotine C

Date:05/28/99ISR Number: 3332691-0Report Type:Periodic Company Report #A0085002  
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/TWICE							
PER DAY/ORAL							

Ostradiol C

Date:05/28/99ISR Number: 3332694-6Report Type:Periodic Company Report #A0085004  
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/TWICE							
PER DAY/ORAL							
		Dry Mouth					
		Insomnia					

Nausea

Date:05/28/99ISR Number: 3332696-XReport Type:Periodic Company Report #A0085005  
Age:32 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Panic Disorder	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/TWICE							
PER DAY/ORAL							

Date:05/28/99ISR Number: 3332698-3Report Type:Periodic Company Report #A0085006  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/TWICE		Drug Effect Decreased					
PER DAY/ORAL		Insomnia		Verapamil	C		

Date:05/28/99ISR Number: 3332700-9Report Type:Periodic Company Report #A0085007  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pyrexia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/TWICE		Rash Generalised					
PER DAY/ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332703-4Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0085008

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/PER DAY/ORAL		Dermatitis Pruritus	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL

Date:05/28/99ISR Number: 3332706-XReport Type:Periodic  
Age:27 YR Gender:Female I/FU:I

Company Report #A0085009

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Dysgeusia Nausea	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL

Date:05/28/99ISR Number: 3332707-1Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0085010

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Alcoholic Hangover Confusional State Disorientation Dizziness	Consumer	Zyban Tablet - Zyban Thyroid Hormones	PS C C	Zyban	ORAL

Date:05/28/99ISR Number: 3332709-5Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #A0085011

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Dermatitis Fungal Infection	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL

Pruritus

Hrt

C

Date:05/28/99ISR Number: 3332711-3Report Type:Periodic Company Report #A0085012  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/PER		Feeling Abnormal					
DAY/ORAL		Nausea		Prempro	C		
		Tremor		Salmetrol Xinafoate	C		

Date:05/28/99ISR Number: 3332713-7Report Type:Periodic Company Report #A0085013  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/TWICE							
PER DAY/ORAL							

Date:05/28/99ISR Number: 3332714-9Report Type:Periodic Company Report #A0085014  
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Neck Pain	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/PER							
DAY/ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3332716-2Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #A0085015

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity	Health	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/PER		Overdose	Professional				
DAY/ORAL		Pruritus		Nicotine Patch	SS		
TRANSDERMAL	21 MG/PER	Rash Erythematous					
DAY/TRANSDERM		Rash Papular					
AL		Urticaria					

Date:05/28/99ISR Number: 3332721-6Report Type:Periodic  
Age:26 YR Gender:Male I/FU:I

Company Report #A0085016

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/TWICE		Pruritus	Professional				
PER DAY/ORAL		Urticaria					

Date:05/28/99ISR Number: 3332723-XReport Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0085017

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/TWICE							
PER DAY/ORAL							

Date:05/28/99ISR Number: 3332724-1Report Type:Periodic  
Age:30 YR Gender:Male I/FU:I

Company Report #A0085018



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333424-4Report Type:Periodic  
Age:68 YR Gender:Male I/FU:I

Company Report #A0082284

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Initial Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Insomnia		Omeprazole	C		
				Atorvastatin Calcium	C		
				Aspirin	C		

Date:05/28/99ISR Number: 3333425-6Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #A0082285

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Nausea					
/ ORAL				Omeprazole	C		

Date:05/28/99ISR Number: 3333426-8Report Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #A0082287

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Nausea	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Nightmare					
/ ORAL		Tachycardia					
		Tremor					

Date:05/28/99ISR Number: 3333427-XReport Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0082288

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Muscle Spasms Muscle Twitching					

Date:05/28/99ISR Number: 3333428-1Report Type:Periodic Company Report #A0082289  
 Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chorea	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Tremor					
TWICE PER DAY							
/ ORAL				Atorvastatin Calcium	C		
				Ramipril	C		
				Maxzide	C		

Date:05/28/99ISR Number: 3333429-3Report Type:Periodic Company Report #A0082290  
 Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pain					
TWICE PER DAY		Pruritus					
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333430-XReport Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0082291

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Erectile Dysfunction Priapism Sexual Dysfunction	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3333431-1Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0082292

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Insomnia	Consumer	Zyban Tablet- Zyban	PS		ORAL

Frusemide	C
Potassium Salt	C
Multivitamin	C

Date:05/28/99ISR Number: 3333432-3Report Type:Periodic  
 Age:27 YR Gender:Female I/FU:I

Company Report #A0082354

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Headache	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL

Birth Control	C
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Date:05/28/99ISR Number: 3333433-5Report Type:Periodic  
 Age:80 YR Gender:Male I/FU:I

Company Report #A0082368

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nightmare	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL							

Date:05/28/99ISR Number: 3333434-7Report Type:Periodic Company Report #A0082380  
 Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arrhythmia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							
DAY / ORAL							

Date:05/28/99ISR Number: 3333435-9Report Type:Periodic Company Report #A0082381  
 Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Skin Odour Abnormal	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Verapamil C  
 Frusemide C  
 Potassium Salt C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333436-0Report Type:Periodic  
 Age:46 YR Gender:Male I/FU:I

Company Report #A0082382

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Headache					
/ ORAL		Hypersensitivity					
		Pruritus					
		Rash Maculo-Papular					
		Swelling					

Date:05/28/99ISR Number: 3333437-2Report Type:Periodic  
 Age:49 YR Gender:Male I/FU:I

Company Report #A0082383

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Constipation	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3333438-4Report Type:Periodic  
 Age:40 YR Gender:Male I/FU:I

Company Report #A0082384

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Dyspnoea					
/ ORAL		Headache					
		Insomnia					
		Night Sweats					
		Pruritus					
		Urticaria					

Date:05/28/99ISR Number: 3333439-6Report Type:Periodic Company Report #A0082385  
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diplopia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ORAL				Pancreatic Enzyme	C		

Date:05/28/99ISR Number: 3333440-2Report Type:Periodic Company Report #A0082386  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Fatigue	Professional				
DAY / ORAL		Hypertension		Desogen	C		
		Nausea					
		Orthostatic Hypotension					
		Vision Blurred					

Date:05/28/99ISR Number: 3333441-4Report Type:Periodic Company Report #A0082387  
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Glossodynia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Tongue Disorder					
TWICE PER DAY							

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Freedom Of Information (FOI) Report

/ ORAL 10 DAY

Doxazosin Mesylate C

Date:05/28/99ISR Number: 3333442-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0088471

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ UNK /							
ORAL	1 WK						

Date:05/28/99ISR Number: 3333443-8Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #A0088496

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Rash Erythematous					
PER DAY /		Skin Disorder					
ORAL		Swelling		Conjugated Estrogens	C		
		Urticaria		Alprazolam	C		

Date:05/28/99ISR Number: 3333444-XReport Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #A0088516

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cold Sweat	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Nausea					
PER DAY /		Pain					
ORAL		Palpitations		Conjugated Estrogens	C		
		Tremor		Alprazolam	C		
		Visual Disturbance		Hydroxyzine	C		

Simvastatin C  
Aspirin C

Date:05/28/99ISR Number: 3333445-1Report Type:Periodic Company Report #A0088517  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Hyperaesthesia					
PER DAY /		Thinking Abnormal					
ORAL							

Date:05/28/99ISR Number: 3333446-3Report Type:Periodic Company Report #A0088520  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Flushing					
PER DAY /		Insomnia					
ORAL							

Pentetrazol C  
Estrogen C  
Nabumetone C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333447-5Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0088521

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Decreased Appetite					
PER DAY /		Dissociation					
ORAL		Dry Mouth		Conjugated Estrogens	C		
		Dysgraphia		Folic Acid	C		
		Insomnia		Extra Strength			
		Tremor		Tylenol Pm	C		
				Cyanocobalamin	C		

Date:05/28/99ISR Number: 3333448-7Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #A0088522

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Salivary Hypersecretion	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY /							
ORAL				Premphase	C		

Date:05/28/99ISR Number: 3333449-9Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #A0088523

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Sedation					
PER DAY /							
ORAL							

Date:05/28/99ISR Number: 3333450-5Report Type:Periodic Company Report #A0088524  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Headache					
PER DAY /							
ORAL				Amitriptyline	C		

Date:05/28/99ISR Number: 3333451-7Report Type:Periodic Company Report #A0088525  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Disturbance In Attention					
PER DAY /		Tremor					
ORAL				Librium	C		
				Conjugated Estrogens	C		
				Vitamin	C		

Date:05/28/99ISR Number: 3333452-9Report Type:Periodic Company Report #A0088544  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoaesthesia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY /							
ORAL							

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150 MG/ PER Abdominal Pain Consumer Zyban Tablet - Zyban PS ORAL  
 DAY / ORAL Abdominal Pain Upper  
 Anxiety  
 Disorientation  
 Nervousness  
 Tremor

Date:05/28/99ISR Number: 3333457-8Report Type:Periodic Company Report #A0088708  
 Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ TWICE		Drug Effect Decreased	Consumer	Zyban Tablet - Zyban	PS		ORAL
PER DAY /		Dry Mouth					
ORAL		Insomnia					
				Multivitamin	C		
				Mineral Supplements	C		
				Omega - 3 Marine			
				Triglyceri	C		

Date:05/28/99ISR Number: 3333458-XReport Type:Periodic Company Report #A0088709  
 Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ PER		Tremor	Consumer	Zyban Tablet - Zyban	PS		ORAL
DAY / ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333459-1Report Type:Periodic  
 Age:34 YR Gender:Female I/FU:I

Company Report #A0088710

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Urticaria					
PER DAY /							
ORAL							

Date:05/28/99ISR Number: 3333460-8Report Type:Periodic  
 Age:65 YR Gender:Female I/FU:I

Company Report #A0088712

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY /							
ORAL							

Prempro	C
Aspirin	C
Calcium Salt	C
Verapamil	C

Date:05/28/99ISR Number: 3333461-XReport Type:Periodic  
 Age:58 YR Gender:Female I/FU:I

Company Report #A0088860

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Insomnia					
PER DAY /							
ORAL							

Irbesartan	C
Conjugated Estrogens	C

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Tachycardia					
TWICE PER DAY							
/ ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Burning Sensation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Cardiovascular Disorder					
DAY / ORAL		Dermatitis		Lisinopril	C		
		Dry Mouth					
		Hypoaesthesia					
		Insomnia					
		Keratoconjunctivitis					
		Sicca					
		Palpitations					
		Paraesthesia					
		Peripheral Coldness					



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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333464-5Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0089910

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
ORAL				Nicotine Gum	SS		ORAL

Date:05/28/99ISR Number: 3333465-7Report Type:Periodic  
Age:83 YR Gender:Female I/FU:I

Company Report #A0089911

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pruritus					
TWICE PER DAY							
/ ORAL							
				Frusemide	C		
				Clopidogrel			
				Bisulphate	C		
				Multivitamin	C		

Date:05/28/99ISR Number: 3333466-9Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0089912

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cough	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Fluid Retention					
TWICE PER DAY							
/ ORAL							
				Mefenidramium			
				Metilsulfat	C		

Date:05/28/99ISR Number: 3333467-0Report Type:Periodic Company Report #A0089913  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pruritus					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3333468-2Report Type:Periodic Company Report #A0089914  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL							
				Verapamil			
				Hydrochloride	C		
				Conjugated Estrogens	C		
				Pravastatin Sodium	C		

Date:05/28/99ISR Number: 3333469-4Report Type:Periodic Company Report #A0089915  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL							
				Nifedipine	C		

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Oestradiol C  
 Mefenidramium C  
 Metilsulfat C  
 Calcium Salt C

Date:05/28/99ISR Number: 3333470-0Report Type:Periodic Company Report #A0089916  
 Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Hypersensitivity					
PER DAY/ ORAL		Pruritus		Estrogen	C		

Date:05/28/99ISR Number: 3333471-2Report Type:Periodic Company Report #A0089917  
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Distension	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Dry Mouth					
PER DAY/ORAL		Mental Impairment		Omeprazole	C		
		Nausea		Trazodone			
		Tremor		Hydrochloride	C		

Date:05/28/99ISR Number: 3333472-4Report Type:Periodic Company Report #A0089918  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Pruritus					
PER DAY/ORAL							

Date:05/28/99ISR Number: 3333473-6Report Type:Periodic Company Report #A0089919  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Asthenia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Blood Pressure Abnormal					
PER DAY/ORAL		Dysarthria		Atenolol	C		
		Tremor		Digoxin	C		

Date:05/28/99ISR Number: 3333474-8Report Type:Periodic Company Report #A0090065  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Face Oedema					
PER DAY/ORAL		Tongue Oedema		Oestradiol	C		

Date:05/28/99ISR Number: 3333475-XReport Type:Periodic Company Report #A0090176  
Age:53 YR Gender:Female I/FU:I

Outcome	PT
	Cough
	Dizziness
	Influenza Like Illness
	Insomnia

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Pain Personality Change Rhinorrhoea Sneezing	Consumer	Zyban Tablet - Zyban	PS		ORAL
				Imipramine Hydrochloride	C		
				Nadolol	C		
				Clonazepam	C		
				Naproxen Sodium	C		
				Estrogen	C		

Date:05/28/99ISR Number: 3333476-1Report Type:Periodic Company Report #A0090177  
Age:71 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dizziness Syncope	Consumer	Zyban Tablet - Zyban	PS		ORAL
				Combivent	C		
				Triamcinolone Acetonide	C		

Date:05/28/99ISR Number: 3333477-3Report Type:Periodic Company Report #A0090178  
Age:58 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
				Metoprolol	C		
				Lisinopril	C		

Pravastatin Sodium	C
Aspirin	C
Clopidogrel	
Bisulphate	C
Magnesium Salt	C
Hypericum	C

Date:05/28/99ISR Number: 3333478-5Report Type:Periodic Company Report #A0090179  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Excitability					
TWICE PER DAY		Hyperhidrosis					
/ ORAL				Aspirin	C		

Date:05/28/99ISR Number: 3333479-7Report Type:Periodic Company Report #A0090180  
 Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Micturition Urgency	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pollakiuria					
TWICE PER DAY		Polyuria					
/ ORAL		Urine Abnormality		Bcg Vaccine	C		

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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333480-3Report Type:Periodic  
Age:59 YR Gender:Male I/FU:I

Company Report #A0090181

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY / ORAL		Dizziness Drug Tolerance Decreased Malaise Nausea	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3333481-5Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0090182

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dermatitis Headache	Consumer	Zyban Tablet - Zyban	PS		ORAL
				Ibuprofen	C		

Date:05/28/99ISR Number: 3333482-7Report Type:Periodic  
Age:31 YR Gender:Male I/FU:I

Company Report #A0090719

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dizziness Irritability Nausea Vomiting	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL

Date:05/28/99ISR Number: 3333483-9Report Type:Periodic  
Age:64 YR Gender:Male I/FU:I

Company Report #A0090720

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/	TWICE	Nausea					
PER DAY /							
ORAL				Atorvastatin Calcium	C		

Date:05/28/99ISR Number: 3333484-0Report Type:Periodic Company Report #A0090721  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /		Feeling Abnormal					
TWICE PER DAY		Hyperaesthesia					
/ ORAL		Hypoaesthesia		Nelfinavir Mesylate	C		
				Combivir	C		

Date:05/28/99ISR Number: 3333485-2Report Type:Periodic Company Report #A0090722  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG / PER		Dyspnoea					
DAY / ORAL							



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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333486-4Report Type:Periodic  
 Age:42 YR Gender:Male I/FU:I

Company Report #A0090723

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Insomnia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
TWICE PER DAY							
/ ORAL							
				Atenolol	C		
				Atorvastatin Calcium	C		
				Vitamin E	C		
				Folic Acid	C		
				Aspirin	C		

Date:05/28/99ISR Number: 3333487-6Report Type:Periodic  
 Age:62 YR Gender:Male I/FU:I

Company Report #A0090724

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dysgeusia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
TWICE PER DAY							
/ ORAL							
		Major Depression		Atenolol	C		
				Amlodipine	C		
				Pravastatin Sodium	C		
				Aspirin	C		

Date:05/28/99ISR Number: 3333488-8Report Type:Periodic  
 Age:56 YR Gender:Male I/FU:I

Company Report #A0090725

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Agitation	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
TWICE PER							
/ ORAL							
		Dermatitis					

Nervousness

DAY/ ORAL

Date:05/28/99ISR Number: 3333489-XReport Type:Periodic Company Report #A0090726  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sleep Disorder	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/ TWICE							
PER DAY /							
ORAL							

Multivitamin C  
Cetirizine  
Hydrochloride C

Date:05/28/99ISR Number: 3333490-6Report Type:Periodic Company Report #A0090727  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/ TIWCE		Headache					
PER DAY/ ORAL		Insomnia		Cephalexin	C		
		Neck Pain					
		Paraesthesia					

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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333491-8Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0090769

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoaesthesia	Health	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/ ORAL			Professional Company Representative	Cimetidine	C		

Date:05/28/99ISR Number: 3333492-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0090823

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/ PER DAY / ORAL							

Date:05/28/99ISR Number: 3333493-1Report Type:Periodic  
 Age:52 YR Gender:Female I/FU:I

Company Report #A0090824

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG / TWICE PER DAY / ORAL		Headache Nausea Neck Pain		Nicotine Glibenclamide	C C		

Date:05/28/99ISR Number: 3333494-3Report Type:Periodic  
 Age:39 YR Gender:Female I/FU:I

Company Report #A0090825

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG / PER DAY / ORAL  
Dry Mouth  
Headache  
Consumer  
Zyban Tablet - Zyban PS  
ORAL  
Clonazepam C

Date:05/28/99ISR Number: 3333495-5Report Type:Periodic Company Report #A0090826  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flatulence	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /		Insomnia					
TWICE PER DAY		Mouth Ulceration					
/ ORAL							

Date:05/28/99ISR Number: 3333496-7Report Type:Periodic Company Report #A0090827  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/ TWICE		Muscle Twitching					
PER DAY /		Weight Decreased					
ORAL				Paracetamol	C		

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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333497-9Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0090828

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ ORAL	WK	Abnormal Dreams Arthralgia Insomnia Lymphadenopathy Myalgia	Health Professional	Zyban Tablet - Zyban	PS	Zyban	ORAL

Date:05/28/99ISR Number: 3333498-0Report Type:Periodic  
Age:27 YR Gender:Male I/FU:I

Company Report #A0090829

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY / ORAL		Dizziness Insomnia Pruritus Rash Erythematous Urticaria	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL

Date:05/28/99ISR Number: 3333499-2Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #A0090830

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Anxiety Dysgeusia Insomnia Nausea Paranoia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Nausea					
TWICE PER DAY							
/ ORAL							
				Glipizide	C		
				Atenolol	C		
				Diltiazem			
				Hydrochloride	C		
				Clopidogrel			
				Bisulphate	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							
DAY/ ORAL							
				Nsaid	C		
				Cetirizine			
				Hydrochloride	C		
				Claritin-D	C		

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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333502-XReport Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0082388

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Cough					
PER DAY/ ORAL		Emotional Disorder		Multivitamin	C		
		Salivary Hypersecretion		Naproxen Sodium	C		
				Ibuprofen	C		

Date:05/28/99ISR Number: 3333503-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0082389

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sleep Disorder	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ ORAL							

Date:05/28/99ISR Number: 3333504-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0082390

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ ORAL		Rash Erythematous					
2.5 WEEKS		Skin Exfoliation					

Date:05/28/99ISR Number: 3333505-5Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #A0082391

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Sedation					
PER DAY/ ORAL							

Frusemide C  
Guanfacine  
Hydrochloride C  
Imipramine  
Hydrochloride C

Date:05/28/99ISR Number: 3333506-7Report Type:Periodic Company Report #A0082392  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Rash Generalised	Professional				
PER DAY/ ORAL		Urticaria					

Date:05/28/99ISR Number: 3333507-9Report Type:Periodic Company Report #A0082393  
Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Generalised	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							



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Date:05/28/99ISR Number: 3333508-0Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0082466

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Zyban Tablet - Zyban	PS		ORAL
150 MG/		Dyspnoea					
SINGLE DOSE /		Heart Rate Increased					
ORAL		Vision Blurred					

Date:05/28/99ISR Number: 3333509-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0082469

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain	Consumer	Zyban Tablet -Zyban	PS	Zyban	ORAL
150 MG/ TWICE		Nervousness					
PER DAY/ ORAL				Nicotine	C		

Date:05/28/99ISR Number: 3333510-9Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0082470

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/ TWICE		Chest Pain					
PER DAY/ ORAL 6 DAY		Dry Mouth					
		Headache					
		Malaise					
		Nervousness					
		Sedation					

Date:05/28/99ISR Number: 3333511-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0082496

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Unevaluable Event	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ ORAL	1 WK						

Date:05/28/99ISR Number: 3333512-2Report Type:Periodic Company Report #A0082497  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Chest Pain	Professional				
PER DAY/ ORAL		Headache					
		Nausea					

Date:05/28/99ISR Number: 3333513-4Report Type:Periodic Company Report #A0082498  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Night Sweats	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ ORAL							

Date:05/28/99ISR Number: 3333514-6Report Type:Periodic Company Report #A0082499  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ ORAL	2 DAY	Pruritus					

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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333515-8Report Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #A0082500

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Face Oedema					
PER DAY/ ORAL				Clonidine	C		
				Spiroinolactone	C		
				Multivitamin	C		
				Aspirin	C		

Date:05/28/99ISR Number: 3333516-XReport Type:Periodic  
Age:75 YR Gender:Male I/FU:I

Company Report #A0082501

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Asthenia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/		Dizziness					
SINGLE DOSE/		Dry Mouth					
ORAL		Headache					
		Insomnia					
		Vision Blurred					

Date:05/28/99ISR Number: 3333517-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0082502

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis	Health	Zyban Tablet - Zyban	PS		ORAL
150MG/ORAL	3 WK		Professional				

Date:05/28/99ISR Number: 3333518-3Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #A0082503

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL							
				Doxepin Hydrochloride	C		
				Zestoretic	C		
				Nicotine	C		

Date:05/28/99ISR Number: 3333519-5Report Type:Periodic Company Report #A0082504  
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL							

Date:05/28/99ISR Number: 3333520-1Report Type:Periodic Company Report #A0082505  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fluid Retention	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Oedema					
PER DAY/ORAL		Weight Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333521-3Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0082506

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE			Professional				
PER DAY/ORAL							

Date:05/28/99ISR Number: 3333522-5Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0081652

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG / ORAL	5 DAY	Disturbance In Attention					
		Fatigue					
		Irritability					

Date:05/28/99ISR Number: 3333523-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0081653

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3333524-9Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #A0081660

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Health	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /							

Eructation Professional  
Nausea  
Loratadine C  
Oxaprozin C

Date:05/28/99ISR Number: 3333525-0Report Type:Periodic Company Report #A0081661  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /							

TWICE PER DAY  
/ ORAL

Date:05/28/99ISR Number: 3333526-2Report Type:Periodic Company Report #A0081865  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /							

TWICE PER DAY  
/ ORAL

ORAL				Wellbutrin Tablet - Controlled Release	SS		ORAL
				Trazodone Hydrochloride	C		
				Thyroid Medication	C		
				Antibiotics	C		



150 MG / PER	Diabetes Mellitus	Health	Zyban Tablet - Zyban	PS	Zyban	ORAL
DAY / ORAL	Hypoglycaemia	Professional				
			Trandolapril	C		
			Insulin	C		
			Salbutamol Sulphate	C		
			Beclomethasone			
			Dipropion	C		

Date:05/28/99ISR Number: 3333530-4Report Type:Periodic Company Report #A0082008  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pruritus	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Birth Control	C		

Date:05/28/99ISR Number: 3333531-6Report Type:Periodic Company Report #A0082009  
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Glossodynia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Clarithromycin	C		
				Combivent	C		
				Omeprazole	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333532-8Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #A0082010

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dysgeusia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL

Date:05/28/99ISR Number: 3333553-5Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0082011

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Dermatitis	Health Professional Company Representative	Zyban Tablet - Zyban	PS	Zyban	ORAL

Date:05/28/99ISR Number: 3333554-7Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0082012

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL	2 WK	Dermatitis Food Allergy Urticaria	Health Professional Company Representative	Zyban Tablet - Zyban	PS	Zyban	ORAL

Date:05/28/99ISR Number: 3333555-9Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #A0082013

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Balance Disorder	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL

Muscle Spasms

TWICE PER DAY

/ ORAL

Clonidine C  
Hydrochlorothiazide C

Date:05/28/99ISR Number: 3333556-0Report Type:Periodic

Company Report #A0082014

Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
		Nervousness					

150 MG /

TWICE PER DAY

/ ORAL

Conjugated Estrogens C  
Omeprazole C

Date:05/28/99ISR Number: 3333557-2Report Type:Periodic

Company Report #A0082015

Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
		Nervousness					

150 MG / PER

DAY / ORAL

19-Aug-2005 12:44 PM

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333559-6Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0082016

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG / PER		Keratoconjunctivitis					
DAY / ORAL		Sicca					
		Nervousness					
		Tachycardia					
		Tremor					

Date:05/28/99ISR Number: 3333560-2Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #A0082018

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypercholesterolaemia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /		Hypertriglyceridaemia					
TWICE PER DAY		Liver Function Test					
/ ORAL		Abnormal					

Date:05/28/99ISR Number: 3333562-6Report Type:Periodic  
Age:80 YR Gender:Male I/FU:I

Company Report #A0082019

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cough	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3333563-8Report Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #A0082020

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Hypoaesthesia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3333564-XReport Type:Periodic Company Report #A0070207  
 Age:50 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dyspepsia	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL							
Gastrooesophageal Reflux Disease							
Insomnia Pruritus							

Date:05/28/99ISR Number: 3333566-3Report Type:Periodic Company Report #A0077455  
 Age:27 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER		Arthralgia	Health	Zyban Tablet - Zyban	PS		ORAL
DAY / ORAL							
150 MG /							
TWICE PER DAY							
/ ORAL							
Bronchospasm							
Dermatitis Face Oedema							
Pyrexia							
Urticaria							
Paracetamol C							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333567-5Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0080328

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Syncope					
TWICE PER DAY							
/ ORAL				Amisulpride	C		
				Fluticasone			
				Propionate	C		
				Birth Control	C		

Date:05/28/99ISR Number: 3333568-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0080869

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / UNK		Drug Ineffective					
/ ORAL		Major Depression					
		Suicidal Ideation					
		Weight Increased					

Date:05/28/99ISR Number: 3333570-5Report Type:Periodic  
Age:65 YR Gender:Female I/FU:I

Company Report #A0081514

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Deafness	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Tinnitus	Professional				
TWICE PER DAY		Tympanic Membrane					
/ ORAL		Disorder		Minocycline	C		
				Ciprofloxacin Hcl	C		
				Antibiotics	C		

Date:05/28/99ISR Number: 3333571-7Report Type:Periodic Company Report #A0081515  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Metrorrhagia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Birth Control	C		

Date:05/28/99ISR Number: 3333573-0Report Type:Periodic Company Report #A0081517  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Depressed Mood					
TWICE PER DAY		Emotional Disorder					
/ ORAL		Feeling Abnormal		Birth Control	C		
		Major Depression					
		Tachycardia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333574-2Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0081518

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Accommodation Disorder Agitation Drug Ineffective Insomnia Irritability	Consumer	Zyban Tablet - Zyban    Prempro	PS    C		ORAL

Date:05/28/99ISR Number: 3333576-6Report Type:Periodic  
 Age:53 YR Gender:Female I/FU:I

Company Report #A0081519

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL	1 WK	Agitation Metrorrhagia Night Sweats Panic Disorder	Consumer	Zyban Tablet - Zyban    Prempro Diazepam	PS    C C		ORAL

Date:05/28/99ISR Number: 3333577-8Report Type:Periodic  
 Age:39 YR Gender:Male I/FU:I

Company Report #A0081520

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Abdominal Pain Dermatitis Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3333579-1Report Type:Periodic Company Report #A0081595  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / UNK		Depression					
/ ORAL	9 DAY	Insomnia					
		Nausea					

Date:05/28/99ISR Number: 3333581-XReport Type:Periodic Company Report #A0081635  
Age:75 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Tolazamide	C		
				Cimetidine	C		
				Potassium Salt	C		

Date:05/28/99ISR Number: 3333583-3Report Type:Periodic Company Report #A0081637  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Psychomotor Hyperactivity					
TWICE PER DAY							
/ ORAL							
				Multivitamin	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333584-5Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0081638

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Face Oedema	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Headache					
TWICE PER DAY		Oedema Peripheral					
/ ORAL		Pruritus		Hormones	C		
		Skin Test Positive		Claritin-D	C		
		Tongue Oedema					
		Urticaria					

Date:05/28/99ISR Number: 3333586-9Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #A0081639

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3333588-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0081640

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / UNK							
/ ORAL	6 WK						

Date:05/28/99ISR Number: 3333589-4Report Type:Periodic  
Age:68 YR Gender:Female I/FU:F

Company Report #A0081648

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150 MG /		Anxiety	Health	Zyban Tablet - Zyban	PS		ORAL
		TWICE PER DAY	Dizziness	Professional				
	/ ORAL		Nervousness					
			Tachycardia		Quinine	C		
					Blood Pressure Medication	C		
					Prednisone	C		
					Diclofenac Sodium	C		
					Hydrochlorothiazide	C		

Date:05/28/99ISR Number: 3333591-2Report Type:Periodic Company Report #A0081649  
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150 MG / UNK		Tremor	Consumer	Zyban Tablet - Zyban	PS		ORAL
	/ ORAL	3 WK						

Date:05/28/99ISR Number: 3333593-6Report Type:Periodic Company Report #A0081650  
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150 MG / UNK		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
	/ ORAL		Nightmare					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333594-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0081651

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Abdominal Pain Upper Discomfort Ill-Defined Disorder Insomnia Vomiting	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3333596-1Report Type:Periodic  
 Age:33 YR Gender:Female I/FU:I

Company Report #A0082101

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Anxiety Chest Pain Dyspnoea Tachycardia Tremor	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3333597-3Report Type:Periodic  
 Age:34 YR Gender:Female I/FU:I

Company Report #A0082103

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dizziness Dysphagia Face Oedema Tachycardia Throat Tightness Urticaria	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3333599-7Report Type:Periodic Company Report #A0082108  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rectal Haemorrhage	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Oxaprozin Centrum	C C		

Date:05/28/99ISR Number: 3333601-2Report Type:Periodic Company Report #A0082158  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Food Allergy Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Atorvastatin Calcium M.V. I Torasemide Metoprolol Succinate Irbesartan ...	C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333602-4Report Type:Periodic  
 Age:36 YR Gender:Female I/FU:I

Company Report #A0082264

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Apathy Depression	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Nausea					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3333603-6Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:I

Company Report #A0082265

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Excitability Insomnia	Health Professional	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Ibuprofen C

Date:05/28/99ISR Number: 3333604-8Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0082266

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							ORAL

Date:05/28/99ISR Number: 3333606-1Report Type:Periodic  
 Age:47 YR Gender:Female I/FU:I

Company Report #A0082268

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS	ORAL
150 MG /				
TWICE PER DAY				
/ ORAL				
		Atenolol	C	
		Lisinopril	C	
		Salbutamol Sulphate	C	

Date:05/28/99ISR Number: 3333607-3Report Type:Periodic Company Report #A0082269  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pruritus Rash Generalised					
TWICE PER DAY		Urticaria					
/ ORAL				Thyroxine Sodium	C		

Date:05/28/99ISR Number: 3333608-5Report Type:Periodic Company Report #A0082270  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Visual Disturbance					
TWICE PER DAY							
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333610-3Report Type:Periodic  
 Age:52 YR Gender:Female I/FU:I

Company Report #A0082271

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Eyelid Disorder Muscle Twitching	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL							

Nutritional Supplement C

Date:05/28/99ISR Number: 3333611-5Report Type:Periodic  
 Age:56 YR Gender:Female I/FU:I

Company Report #A0082273

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Agitation Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Cystitis					
/ ORAL		Dizziness					

Atenolol C  
 Quinapril  
 Hydrochloride C  
 Prempro C  
 Thyroxine Sodium C  
 Clopidogrel  
 Bisulphate C  
 Fluvastatin Sodium C  
 Aspirin C

Date:05/28/99ISR Number: 3333613-9Report Type:Periodic  
 Age:31 YR Gender:Female I/FU:I

Company Report #A0082274

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypersensitivity	Consumer	Zyban Tablet -			

150 MG / Pruritus Zyban PS ORAL  
Urticaria  
TWICE PER DAY  
/ ORAL

Date:05/28/99ISR Number: 3333615-2Report Type:Periodic Company Report #A0082275  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chills Sensation Of Heaviness	Health Professional	Zyban Tablet - Zyban	PS		ORAL

150 MG /  
TWICE PER DAY  
/ ORAL

Date:05/28/99ISR Number: 3333616-4Report Type:Periodic Company Report #A0082276  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cold Sweat Nightmare	Consumer	Zyban Tablet - Zyban	PS		ORAL

1.5 TABLET /  
THREE TIMES  
PE ORAL

Diclofenac C  
Codeine C  
Nicotine C



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333617-6Report Type:Periodic  
 Age:42 YR Gender:Female I/FU:I

Company Report #A0082277

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Fatigue	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Irritability					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3333619-XReport Type:Periodic  
 Age:41 YR Gender:Female I/FU:I

Company Report #A0082278

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3333620-6Report Type:Periodic  
 Age:42 YR Gender:Male I/FU:I

Company Report #A0082279

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Condition Aggravated					
TWICE PER DAY		Confusional State					
/ ORAL		Disorientation Migraine Smoker		Ibuprofen Paracetamol	C C		

Date:05/28/99ISR Number: 3333622-XReport Type:Periodic Company Report #A0082280  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3333623-1Report Type:Periodic Company Report #A0082281  
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pruritus					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3333689-9Report Type:Periodic Company Report #A0083898  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Urticaria					
TWICE PER DAY							
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333692-9Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #A0083919

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Disorientation					
TWICE PER DAY		Dyspnoea					
/ ORAL		Euphoric Mood					
		Hypoaesthesia					
		Panic Disorder					

Date:05/28/99ISR Number: 3333693-0Report Type:Periodic  
Age:57 YR Gender:Male I/FU:I

Company Report #A0083922

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Pruritus	Professional				
DAY / ORAL							

Date:05/28/99ISR Number: 3333694-2Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0083923

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Back Pain					
DAY / ORAL		Headache		Percocet	C		
				Ranitidine			
				Hydrochloride	C		

Date:05/28/99ISR Number: 3333697-8Report Type:Periodic  
Age:26 YR Gender:Male I/FU:I

Company Report #A0083925

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3333699-1Report Type:Periodic Company Report #A0083927  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased	Consumer	Zyban Tablet -Zyban	PS		ORAL
150 MG /		Cough					
TWICE PER DAY		Drug Dependence					
/ ORAL	4 WK	Dry Mouth		Metoprolol Tartate	C		
		Headache		Clorazepate			
		Insomnia		Dipotassium	C		
		Nervousness		Indapamide	C		

Date:05/28/99ISR Number: 3333700-5Report Type:Periodic Company Report #A0083977  
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Palpitations					
TWICE PER DAY							
/ ORAL				Glipizide	C		
				Loratadine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333701-7Report Type:Periodic  
Age:71 YR Gender:Male I/FU:I

Company Report #A0083980

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /			Professional				
TWICE PER DAY							
/ ORAL				Omeprazole	C		
				Multivitamin	C		
				Aspirin	C		

Date:05/28/99ISR Number: 3333703-0Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0084028

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Health	Zyban Tablet - Zyban	PS		ORAL
3 TABLETS /		Dyspnoea	Professional				
UNK / ORAL		Urticaria					

Date:05/28/99ISR Number: 3333704-2Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0084043

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Formication	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							
DAY / ORAL							

Date:05/28/99ISR Number: 3333705-4Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #A0084044

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG / PER	Irritability	Consumer	Zyban Tablet - Zyban	PS	ORAL	
DAY / ORAL						
Date:05/28/99ISR Number: 3333707-8Report Type:Periodic Company Report #A0084418						
Age:40 YR Gender:Female I/FU:I						
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
150 MG /	Chest Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY	Dyspnoea					
/ ORAL	Feeling Drunk					
	Flushing					

Date:05/28/99ISR Number: 3333710-8Report Type:Periodic Company Report #A0084632						
Age:62 YR Gender:Female I/FU:I						
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
UNK / UNK /	White Blood Cell Count	Health	Zyban Tablet - Zyban	PS		ORAL
ORAL	Decreased	Professional				
	9 WK	Company Representative	Fluvastatin Sodium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333712-1Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0084702

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dermatitis Dyspnoea Face Oedema Hypersensitivity Tremor Urticaria	Consumer	Zyban Tablet - Zyban   Ibuprofen	PS   C		ORAL

Date:05/28/99ISR Number: 3333715-7Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #A0084800

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Headache	Consumer	Zyban Tablet - Zyban  Cimetidine	PS  C		ORAL

Date:05/28/99ISR Number: 3333716-9Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0084823

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Bronchospasm Dermatitis Dyspnoea Pruritus Urticaria	Consumer	Zyban Tablet - Zyban   Valaciclovir	PS   C		ORAL

Date:05/28/99ISR Number: 3333718-2Report Type:Periodic Company Report #A0084825  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Breast Tenderness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Dry Mouth					
DAY / ORAL							

Date:05/28/99ISR Number: 3333745-5Report Type:Periodic Company Report #A0084828  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Oedema Peripheral					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3333746-7Report Type:Periodic Company Report #A0084829  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Hypersensitivity					
TWICE PER DAY							
/ ORAL				Birth Control	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333748-0Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #A0084831

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / UNK / ORAL		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Erythema	Health				
UNK / UNK / UNK		Pain	Professional	Nicotine Patch	SS		
		Pruritus	Other				
		Swelling					

Date:05/28/99ISR Number: 3333753-4Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0085267

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Crying	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3333755-8Report Type:Periodic  
Age:57 YR Gender:Male I/FU:I

Company Report #A0085328

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3333759-5Report Type:Periodic  
Age:26 YR Gender:Male I/FU:I

Company Report #A0085329

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Dyspnoea					

Insomnia  
Pharyngeal Oedema  
Pruritus

Date:05/28/99ISR Number: 3333762-5Report Type:Periodic Company Report #A0085330  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Halitosis					
TWICE PER DAY							
/ ORAL				Atorvastatin Calcium	C		

Date:05/28/99ISR Number: 3333765-0Report Type:Periodic Company Report #A0085331  
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperaesthesia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Naproxen	C		
				Glipizide	C		
				Hydrocodone +			
				Paracetamol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333767-4Report Type:Periodic  
 Age:65 YR Gender:Male I/FU:I

Company Report #A0085332

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Nausea					
/ ORAL							

Date:05/28/99ISR Number: 3333769-8Report Type:Periodic  
 Age:56 YR Gender:Male I/FU:I

Company Report #A0085333

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Swelling					
/ ORAL		Urticaria					

Date:05/28/99ISR Number: 3333771-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0085335

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Bipolar I Disorder	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Psychotic Disorder					

Date:05/28/99ISR Number: 3333775-3Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:I

Company Report #A0085336

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Face Oedema	Health	Zyban Tablet - Zyban	PS		ORAL

Herpes Simplex Professional  
TWICE PER DAY  
Pruritus  
/ ORAL  
Urticaria

Date:05/28/99ISR Number: 3333777-7Report Type:Periodic Company Report #A0085337  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Influenza Like Illness					
TWICE PER DAY		Insomnia					
/ ORAL		Lethargy		Cold Medication	C		
		Pruritus		M.V.I	C		

Date:05/28/99ISR Number: 3333779-0Report Type:Periodic Company Report #A0085338  
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dizziness					
TWICE PER DAY		Syncope					
/ ORAL				Insulin	C		
				Thyroxine Sodium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333781-9Report Type:Periodic  
 Age:39 YR Gender:Female I/FU:I

Company Report #A0085339

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Paraesthesia					
TWICE PER DAY							
/ ORAL							
				Coffee (Formulation Unknown)	SS		ORAL
ORAL				Bupirone Hydrochloride	C		

Date:05/28/99ISR Number: 3333783-2Report Type:Periodic  
 Age:55 YR Gender:Female I/FU:I

Company Report #A0085340

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Migraine	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Estrogen	C		
				Thyroxine Sodium	C		
				Esgic	C		

Date:05/28/99ISR Number: 3333785-6Report Type:Periodic  
 Age:55 YR Gender:Female I/FU:I

Company Report #A0085341

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL							
				Fluoxetine Hydrochloride	C		

Date:05/28/99ISR Number: 3333788-1Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0085342

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							
DAY / ORAL							

Glucophage	C
Simvastatin	C
Famotidine	C
Conjugated Estrogens	C
Fluticasone	
Propionate	C
Salbutamol Sulphate	C
Ipratropium Bromide	C

Date:05/28/99ISR Number: 3333790-XReport Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #A0085343

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia	Professional				
TWICE PER DAY							
/ ORAL							

Alendronate Sodium	C
Conjugated Estrogens	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333794-7Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0085344

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /			Professional				
ORAL;							
DURATION: ??							
WEEKS							

Date:05/28/99ISR Number: 3333797-2Report Type:Periodic  
 Age:35 YR Gender:Female I/FU:I

Company Report #A0085345

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthritis	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Difficulty In Walking	Professional				
TWICE PER DAY		Hypersensitivity					
/ ORAL		Inflammation					
		Oedema Peripheral					
		Pain					
		Pruritus					
		Rash Erythematous					
		Urticaria					

Date:05/28/99ISR Number: 3333800-XReport Type:Periodic  
 Age:43 YR Gender:Male I/FU:I

Company Report #A0085373

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Angioneurotic Oedema	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Discomfort	Professional				
DAY / ORAL		Feeling Hot	Company	Amlodipine	C		
		Oedema Peripheral	Representative	Hydrochlorothiazide	C		

Pain  
Paraesthesia  
Rash Erythematous

Metoprolol Tartrate C

Date:05/28/99ISR Number: 3333801-1Report Type:Periodic Company Report #A0085554  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Unevaluable Event	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL							

Date:05/28/99ISR Number: 3333803-5Report Type:Periodic Company Report #A0085657  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/PER DAY/ORAL							

Date:05/28/99ISR Number: 3333806-0Report Type:Periodic Company Report #A0085658  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/PER DAY/ORAL							
				Estrogen	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333808-4Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #A0085659

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/PER DAY/ORAL		Abdominal Pain Chest Pain	Consumer	Zyban Tablet-Zyban	PS		ORAL

Date:05/28/99ISR Number: 3333811-4Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #A0085660

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/PER DAY/ORAL		Headache Nausea	Consumer	Zyban Tablet-Zyban	PS		ORAL

Prempro C

Date:05/28/99ISR Number: 3333813-8Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0085661

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Gingival Bleeding	Consumer	Zyban Tablet-Zyban	PS		ORAL

Nicotine C

Date:05/28/99ISR Number: 3333815-1Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0085662

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Gingival Bleeding	Consumer	Zyban Tablet-Zyban	PS		ORAL

Date:05/28/99ISR Number: 3333817-5Report Type:Periodic Company Report #A0085663  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL							

Date:05/28/99ISR Number: 3333818-7Report Type:Periodic Company Report #A0085664  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE		Emotional Disorder					
PER DAY/ORAL							

Date:05/28/99ISR Number: 3333824-2Report Type:Periodic Company Report #A0085665  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Health	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE		Lip Dry	Professional				
PER DAY/ORAL		Urticaria		Librax	C		
				Herbal Medication	C		
				Famciclovir	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333826-6Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #A0085666

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE		Hypersensitivity	Professional				
PER DAY/ORAL		Rash Generalised		Vitamin	C		

Date:05/28/99ISR Number: 3333827-8Report Type:Periodic  
Age:71 YR Gender:Female I/FU:I

Company Report #A0085667

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Health	Zyban Tablet-Zyban	PS		ORAL
150 MG/PER		Muscle Twitching	Professional				
DAY/ORAL				Prempro	C		
				Atorvastatin Calcium	C		
				Verapamil			
				Hydrochloride	C		

Date:05/28/99ISR Number: 3333832-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0085668

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ORAL				Guaiphenesin (Formulation Unknown)	SS		
				Guaifenesin Dm (Formulation Unknown)	SS		

Date:05/28/99ISR Number: 3333834-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0085669

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ORAL		Dizziness Malaise Nausea Vomiting					

Date:05/28/99ISR Number: 3333837-0Report Type:Periodic Company Report #A0085670  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ORAL							

Date:05/28/99ISR Number: 3333839-4Report Type:Periodic Company Report #A0085671  
Age:51 YR Gender:Female I/FU:I

Outcome	PT
	Arthralgia Dermatitis Dysgeusia Face Oedema Myalgia Oral Candidiasis

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Swelling

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Consumer	Zyban Tablet-Zyban	PS		ORAL
			Thyroxine Sodium	C		

Date:05/28/99ISR Number: 3333842-4Report Type:Periodic Company Report #A0085672  
Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ORAL		Feeling Abnormal					
		Hypoaesthesia Paranoia		Birth Control	C		

Date:05/28/99ISR Number: 3333843-6Report Type:Periodic Company Report #A0085674  
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/PER DAY/ORAL		Disorientation					
		Dizziness					
		Dry Mouth Headache Paraesthesia Tremor Vomiting					

Date:05/28/99ISR Number: 3333846-1Report Type:Periodic Company Report #A0085675  
Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG/TWICE	Abdominal Pain	Consumer	Zyban Tablet-Zyban	PS	ORAL
PER DAY/ORAL	Headache				
			Ascorbic Acid	C	

Date:05/28/99ISR Number: 3333848-5Report Type:Periodic Company Report #A0085676  
 Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL							

- Lorazepam C
- Mirtazapine C
- Montelukast Sodium C
- Glibenclamide C
- Simvastatin C
- Verapamil C
- Hydrochloride C
- Prednisone C
- Salbutamol Sulphate C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333851-5Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #A0085677

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/TWICE		Parosmia					
PER DAY/ORAL				Antibiotics	C		

Date:05/28/99ISR Number: 3333962-4Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0087976

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TWICE		Constipation					
PER DAY/ ORAL		Crying		Alendronate Sodium	C		
		Headache		Alprazolam	C		
		Therapeutic Response		Prednisone	C		
		Unexpected		Unknown	C		

Date:05/28/99ISR Number: 3333978-8Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #A0087977

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia	Consumer	Zyban Tablet- Zyban	PS		ORAL
150 MG/ TWICE		Fatigue					
PER DAY/ ORAL		Insomnia					
		Nausea					
		Tachycardia					

Date:05/28/99ISR Number: 3333980-6Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0087978

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TWICE		Insomnia					
PER DAY/ ORAL		Micturition Urgency		Loratadine	C		
		Pollakiuria		Fluticasone			
				Propionate	C		
				Ipratropium Bromide	C		

Date:05/28/99ISR Number: 3333983-1Report Type:Periodic Company Report #A0087979  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:05/28/99ISR Number: 3333985-5Report Type:Periodic Company Report #A0087980  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TWICE		Dry Mouth					
PER DAY/ ORAL		Insomnia		Conj. Estrogen +			
		Nausea		Medroxyprog	C		
		Tremor		Paracetamol	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3333987-9Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #A0087981

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vomiting	Health	Zyban Tablet-Zyban	PS		ORAL
150 MG/ PER			Professional				
DAY/ ORAL				Temazepam	C		

Date:05/28/99ISR Number: 3333991-0Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0087982

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoaesthesia	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TWICE		Hypoaesthesia Oral					
PER DAY/ ORAL							

Date:05/28/99ISR Number: 3334043-6Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #A0087983

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Health	Zyban Tablet- Zyban	PS		ORAL
150 MG/ TWICE		Dysarthria	Professional				
PER DAY/ ORAL		Hypotension		Atenolol	C		
		Sedation		Digoxin	C		
		Tremor					

Date:05/28/99ISR Number: 3334046-1Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0087984

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TWICE							

PER DAY/ ORAL

Date:05/28/99ISR Number: 3334048-5Report Type:Periodic Company Report #A0087985  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TWICE		Glossodynia					
PER DAY/ ORAL		Mouth Ulceration		Conjugated Estrogens	C		
				Ibuprofen	C		
				Nicotine Polacrilex	C		

Date:05/28/99ISR Number: 3334051-5Report Type:Periodic Company Report #A0087986  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TWICE		Pruritus					
PER DAY/ ORAL							

Date:05/28/99ISR Number: 3334053-9Report Type:Periodic Company Report #A0087987  
Age:40 YR Gender:Female I/FU:I

Outcome	PT
	Dermatitis
	Face Oedema
	Pain

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Pruritus Rash Erythematous Urticaria	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ PER DAY/ ORAL				Atenolol Tyleno No. 3 Opium	C C C		

Date:05/28/99ISR Number: 3334070-9Report Type:Periodic Company Report #A0087988  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Affective Disorder	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ ORAL							

Date:05/28/99ISR Number: 3334073-4Report Type:Periodic Company Report #A0087989  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL							

Date:05/28/99ISR Number: 3334076-XReport Type:Periodic Company Report #A0087990  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL		Dizziness		Thyroxine Sodium Ranitidine	C		

Hydrochloride C  
Prempro C  
Esgic-Plus C

Date:05/28/99ISR Number: 3334077-1Report Type:Periodic Company Report #A0087991  
Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Accommodation Disorder	Consumer	Zyban Tablet-Zyban	PS		ORAL
PER DAY/ ORAL		Crying					
150 MG/ TWICE		Headache		Wellbutrin			
PER DAY/ ORAL		Malaise		Tablet-Controlled			
		Nausea		Release	SS		ORAL
		Nervousness					
		Sedation		Demulen	C		

Date:05/28/99ISR Number: 3334080-1Report Type:Periodic Company Report #A0087992  
Age:41 YR Gender:Male I/FU:I

Outcome PT  
Abnormal Dreams  
Aggression  
Depression  
Diarrhoea  
Dysgeusia  
Fatigue

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Headache Libido Decreased Parosmia	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ PER DAY/ ORAL							

Date:05/28/99ISR Number: 3334084-9Report Type:Periodic Company Report #A0087993  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphemia	Consumer	Zyban Tablet- Zyban	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL		Dyspnoea					
		Hypoglycaemia		Tylenol No. 3	C		
		Muscle Twitching		Loratadine	C		
		Vision Blurred		Temazepam	C		
				Famotidine	C		
				Diphenoxylate	C		
				Ranitidine			
				Hydrochloride	C		

Date:05/28/99ISR Number: 3334087-4Report Type:Periodic Company Report #A0087994  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet-Zyban	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL		Nervousness					
		Stomach Discomfort		Salbutamol Sulphate	C		

Date:05/28/99ISR Number: 3334088-6Report Type:Periodic Company Report #A0087995  
Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Furuncle	Consumer	Zyban Tablet- Zyban	PS		ORAL
150 MG/ TWICE		Pain					
PER DAY/ ORAL	7 DAY	Swelling					
		Urticaria					

Date:05/28/99ISR Number: 3334133-8Report Type:Periodic Company Report #A0090968  
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pruritus					
TWICE PER DAY		Urticaria					
/ ORAL				Nicotine	C		
				Isoniazid	C		

Date:05/28/99ISR Number: 3334137-5Report Type:Periodic Company Report #A0090974  
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cold Sweat	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Dermatitis					
		Dizziness					
		Headache					
		Vomiting					



150 MG / PER	Amnesia	Consumer	Zyban Tablet - Zyban	PS	ORAL
DAY / ORAL	Disturbance In Attention				
	Tremor				
Date:05/28/99ISR Number: 3334147-8Report Type:Periodic Company Report #A0090980					
Age:39 YR	Gender:Male	I/FU:I			
Outcome	PT	Report Source	Product	Role	Manufacturer
Dose	Duration				Route
150 MG /	Dry Mouth	Consumer	Zyban Tablet - Zyban	PS	ORAL
TWICE PER DAY	Nausea				
/ ORAL	Tachycardia				
Date:05/28/99ISR Number: 3334152-1Report Type:Periodic Company Report #A0091163					
Age:37 YR	Gender:Female	I/FU:I			
Outcome	PT	Report Source	Product	Role	Manufacturer
Dose	Duration				Route
150 MG /	Rash Generalised	Consumer	Zyban Tablet - Zyban	PS	ORAL
TWICE PER DAY	Urticaria				
/ ORAL					
Date:05/28/99ISR Number: 3334153-3Report Type:Periodic Company Report #A0091167					
Age:68 YR	Gender:Female	I/FU:I			
Outcome	PT				
	Dermatitis Bullous				
	Oedema Peripheral				
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Pruritus Rash Erythematous Rash Macular	Health	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Skin Nodule	Professional				
/ ORAL		Urticaria					

Prempro	C
Amlodipine	C

Date:05/28/99ISR Number: 3334156-9Report Type:Periodic Company Report #A0091168  
 Age:43 YR Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Constipation	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Decreased Appetite					
/ ORAL		Dry Mouth					
		Insomnia		Naproxen Sodium	C		
		Therapeutic Response					
		Unexpected					

Date:05/28/99ISR Number: 3334157-0Report Type:Periodic Company Report #A0091169  
 Age:44 YR Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Constipation	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Flatulence					
/ ORAL;							

DURATION :

WEEKS

Date:05/28/99ISR Number: 3334158-2Report Type:Periodic Company Report #A0091170  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Lower	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Back Pain					
TWICE PER DAY		Pollakiuria					
/ ORAL				Antibiotics	C		

Date:05/28/99ISR Number: 3334160-0Report Type:Periodic Company Report #A0091171  
Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Birth Control	C		

Date:05/28/99ISR Number: 3334163-6Report Type:Periodic Company Report #A0091172  
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pain In Extremity					
TWICE PER DAY							
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3334164-8Report Type:Periodic  
 Age:51 YR Gender:Female I/FU:I

Company Report #A0091173

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Diarrhoea					
DAY / ORAL		Headache					
		Pain					
		Pain In Extremity					

Date:05/28/99ISR Number: 3334307-6Report Type:Periodic  
 Age:48 YR Gender:Female I/FU:I

Company Report #A0091176

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Eye Irritation					
TWICE PER DAY		Insomnia					
/ ORAL		Urticaria		Prempro	C		

Date:05/28/99ISR Number: 3334308-8Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:I

Company Report #A0091178

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Balance Disorder					
DAY / ORAL		Dry Mouth		Alprazolam	C		
		Nausea		Zovia	C		
		Tremor					
		Visual Disturbance					

Date:05/28/99ISR Number: 3334309-XReport Type:Periodic Company Report #A0091180  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL							

Date:05/28/99ISR Number: 3334310-6Report Type:Periodic Company Report #A0091181  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Malaise					
DAY / ORAL				Birth Control	C		

Date:05/28/99ISR Number: 3334311-8Report Type:Periodic Company Report #A0091182  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Arthritis	Professional				
		Myalgia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3334790-6Report Type:Periodic  
 Age:68 YR Gender:Male I/FU:I

Company Report #A0090409

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dermatitis					
		Urticaria					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3334791-8Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0090577

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Dermatitis	Professional Company Representative				

Date:05/28/99ISR Number: 3334792-XReport Type:Periodic  
 Age:66 YR Gender:Male I/FU:I

Company Report #A0090683

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Amnesia					
		Coordination Abnormal					
TWICE PER DAY		Disorientation					
/ ORAL		Dizziness Headache Speech Disorder					

Date:05/28/99ISR Number: 3334794-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0090701

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Chest Pain Headache	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3334795-5Report Type:Periodic Company Report #A0090702  
 Age:49 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Dysgeusia	Consumer	Zyban Tablet - Zyban Iron Supplements	PS C		ORAL

Date:05/28/99ISR Number: 3334797-9Report Type:Periodic Company Report #A0090703  
 Age:42 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Nausea	Consumer	Zyban Tablet - Zyban Atenolol Propafenone Hydrochloride	PS C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3334798-0Report Type:Periodic  
 Age:52 YR Gender:Female I/FU:I

Company Report #A0090704

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Arthralgia Dermatitis Hypersensitivity Oedema Peripheral Pain Urticaria	Consumer	Zyban Tablet - Zyban   Quinapril Hydrochloride Amlodipine Loratadine Metoprolol Succinate	PS   C C C C		ORAL

Date:05/28/99ISR Number: 3334799-2Report Type:Periodic  
 Age:45 YR Gender:Female I/FU:I

Company Report #A0090705

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Apathy Decreased Appetite Dizziness Hyperhidrosis Nervousness Throat Tightness	Consumer	Zyban Tablet - Zyban   Prednisone Simvastatin Salbutamol Sulphate	PS   C C C		ORAL

Date:05/28/99ISR Number: 3334801-8Report Type:Periodic  
 Age:55 YR Gender:Female I/FU:I

Company Report #A0090706

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dermatitis Dry Mouth Oedema Peripheral	Consumer	Zyban Tablet - Zyban	PS		ORAL

Pruritus

Diazepam C  
Conjugated Estrogens C  
Simvastatin C  
Amitriptyline Hcl C

Date:05/28/99ISR Number: 3334803-1Report Type:Periodic Company Report #A0090707  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pruritus					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3334804-3Report Type:Periodic Company Report #A0090708  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Omeprazole C  
Diltiazem  
Hydrochloride C  
Enalapril Maleate C  
Amlodipine C  
Metoprolol C  
Glipizide C



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3334806-7Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0090710

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Multivitamin	C		
				Calcium Salt	C		
				Ascorbic Acid	C		
				Vitamin E	C		

Date:05/28/99ISR Number: 3334807-9Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0090711

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dry Mouth					
TWICE PER DAY		Dysgeusia					
/ ORAL		Headache					
		Night Sweats					
		Paraesthesia					
		Weight Decreased					

Date:05/28/99ISR Number: 3334809-2Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0090712

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Atorvastatin Calcium C  
 Isosorbide  
 Mononitrate C  
 Amlodipine C  
 Metoprolol C

Date:05/28/99ISR Number: 3334810-9Report Type:Periodic Company Report #A0090713  
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Zafirlukast C  
 Estrogen C

Date:05/28/99ISR Number: 3334812-2Report Type:Periodic Company Report #A0090714  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pruritus					
TWICE PER DAY							
/ ORAL							

Ortho-Novum C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3334814-6Report Type:Periodic  
Age:28 YR Gender:Male I/FU:I

Company Report #A0090715

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dizziness Hallucination Psychomotor Hyperactivity Syncope	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3334815-8Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0090716

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Insomnia Tremor	Consumer	Zyban Tablet - Zyban	PS		ORAL
				Terbinafine	C		

Date:05/28/99ISR Number: 3334816-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0090717

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Constipation Drug Dependence Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3334817-1Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #A0090718

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL

Insomnia

TWICE PER

DAY/ ORAL

Date:05/28/99ISR Number: 3334818-3Report Type:Periodic Company Report #A0077079  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Mouth Ulceration	Professional				

TWICE PER DAY

/ ORAL

Date:05/28/99ISR Number: 3334819-5Report Type:Periodic Company Report #A0077145  
Age:14 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							

TWICE PER DAY

/ ORAL

Date:05/28/99ISR Number: 3334820-1Report Type:Periodic Company Report #A0077952  
Age:38 YR Gender:Female I/FU:F

Outcome	PT
	Dysphagia
	Hyperhidrosis
	Insomnia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tachycardia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG /		Health	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Professional				
/ ORAL			Hydroxyzine	C		

Date:05/28/99ISR Number: 3334821-3Report Type:Periodic Company Report #A0077953  
 Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Crying	Professional				
TWICE PER DAY		Depression					
/ ORAL		Headache					

Date:05/28/99ISR Number: 3334822-5Report Type:Periodic Company Report #A0078431  
 Age:69 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Apathy	Professional				
TWICE PER DAY		Asthenia					
/ ORAL		Hypoaesthesia		Ascorbic Acid	C		
		Lethargy					
		Nausea					
		Vertigo					

Date:05/28/99ISR Number: 3334824-9Report Type:Periodic Company Report #A0078434  
 Age:52 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Insomnia	Health	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Paranoia	Professional				
/ ORAL		Psychotic Disorder					
		Sleep Disorder		Lovastatin	C		

Date:05/28/99ISR Number: 3334826-2Report Type:Periodic Company Report #A0078822  
Age:39 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Disorientation	Health	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY			Professional				
/ ORAL							

Date:05/28/99ISR Number: 3334828-6Report Type:Periodic Company Report #A0078836  
Age:56 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Abdominal Distension	Health	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Feeling Abnormal	Professional				
/ ORAL		Headache					
		Hypoaesthesia		Salmeterol Xinafoate	C		
				Salbutamol Sulphate	C		
				Famotidine	C		
				Beclomethasone			
				Dipropion	C		

Freedom Of Information (FOI) Report

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Date:05/28/99ISR Number: 3334830-4Report Type:Periodic Company Report #A0079576  
 Age:42 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Drug Interaction	Health	Zyban Tablet - Zyban	PS		ORAL
		Hyperhidrosis	Professional				
		Influenza Like Illness					
20 MG / ORAL		Nausea Serotonin Syndrome		Fluoxetine Hydrochloride Tablet	SS		ORAL
		Tachycardia Tremor Vomiting		Conjugated Estrogens	C		

Date:05/28/99ISR Number: 3334831-6Report Type:Periodic Company Report #A0079617  
 Age:30 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Chest Discomfort	Health	Zyban Tablet - Zyban	PS		ORAL
		Dizziness	Professional				
		Dyspnoea Feeling Abnormal Hyperventilation Nausea Pruritus Syncope Tachycardia Tremor Urticaria		Birth Control	C		

Date:05/28/99ISR Number: 3334833-XReport Type:Periodic Company Report #A0080055  
 Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Hypoglycaemia	Professional				
TWICE PER DAY							
/ ORAL							
				Human Insulin (Formulation Unknown)	SS		
SUBCUTANEOUS	AS DIRECTED /						
SUBCUTANEOUS							
				Captopril	C		
				Frusemide	C		
				Digoxin	C		
				Isotretinoin	C		
				Warfarin Sodium	C		
				Amlodipine	C		
				Isosorbide	C		
				Int./Long-Acting Insulin	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3334834-1Report Type:Periodic  
Age:58 YR Gender:Female I/FU:F

Company Report #A0080060

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Dermatitis	Professional				
		Pruritus					

Date:05/28/99ISR Number: 3334835-3Report Type:Periodic  
Age:27 YR Gender:Male I/FU:F

Company Report #A0080096

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL	2 WK	Urticaria	Professional				

Date:05/28/99ISR Number: 3334837-7Report Type:Periodic  
Age:44 YR Gender:Female I/FU:F

Company Report #A0080372

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Tremor					
DAY / ORAL							

Date:05/28/99ISR Number: 3334839-0Report Type:Periodic  
Age:39 YR Gender:Male I/FU:F

Company Report #A0080700

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Disturbance In Attention	Professional				
DAY / ORAL		Dizziness					
		Headache					
		Vomiting					

Date:05/28/99ISR Number: 3334840-7Report Type:Periodic Company Report #A0080878  
Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgraphia	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Tremor	Professional				
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3334843-2Report Type:Periodic Company Report #A0081148  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Nightmare	Professional Company Representative				

Date:05/28/99ISR Number: 3334845-6Report Type:Periodic Company Report #A0081198  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL			Professional Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3334848-1Report Type:Periodic  
 Age:33 YR Gender:Female I/FU:F

Company Report #A0081199

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL			Professional Company Representative				

Date:05/28/99ISR Number: 3334851-1Report Type:Periodic  
 Age:41 YR Gender:Female I/FU:F

Company Report #A0081332

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pain	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pruritus	Professional				
TWICE PER DAY		Rash Erythematous					
/ ORAL		Urticaria		Nicotine (Formulation Unknown)	SS		
TRANSDERMAL	TRANSDERMAL						

Date:05/28/99ISR Number: 3334853-5Report Type:Periodic  
 Age:38 YR Gender:Male I/FU:I

Company Report #A0090184

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gingival Bleeding	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3334856-0Report Type:Periodic  
 Age:59 YR Gender:Male I/FU:I

Company Report #A0090185

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Irritability	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Nervousness					
/ ORAL				Lovastatin	C		
				Atenolol	C		

Date:05/28/99ISR Number: 3334857-2Report Type:Periodic Company Report #A0090186  
 Age:42 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Pruritus	Health	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Swelling	Professional				
/ ORAL		Urticaria					

Date:05/28/99ISR Number: 3334860-2Report Type:Periodic Company Report #A0090188  
 Age:42 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Abdominal Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3334861-4Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #A0090189

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Hypoaesthesia					
TWICE PER DAY							
/ ORAL				Desogen	C		

Date:05/28/99ISR Number: 3334862-6Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0090190

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Dizziness					

Date:05/28/99ISR Number: 3334863-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0090191

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL							

Date:05/28/99ISR Number: 3334865-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0090192

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3334867-5Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0090193

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Psychomotor Hyperactivity					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3334869-9Report Type:Periodic  
Age:77 YR Gender:Female I/FU:I

Company Report #A0090194

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Famotidine	C
Theophylline	C
Conjugated Estrogens	C
Potassium Salt	C
Ramipril	C
Diltiazem	
Hydrochloride	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3334871-7Report Type:Periodic  
 Age:64 YR Gender:Female I/FU:I

Company Report #A0090195

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Musculoskeletal Stiffness					
/ ORAL		Tremor					
				Sinemet	C		
				Conjugated Estrogens	C		
				Medroxyprogesterone			
				Ace.	C		

Date:05/28/99ISR Number: 3334874-2Report Type:Periodic  
 Age:55 YR Gender:Female I/FU:I

Company Report #A0090196

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Alopecia					
/ ORAL							
				Conjugated Estrogens	C		
				Diazepam	C		
				Entex	C		
				Fexofenadine			
				Hydrochlorid	C		
				Acyclovir	C		

Date:05/28/99ISR Number: 3334878-XReport Type:Periodic  
 Age:33 YR Gender:Male I/FU:I

Company Report #A0090197

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Dry Mouth					

/ ORAL  
Epistaxis  
Nervousness

Date:05/28/99ISR Number: 3334882-1Report Type:Periodic Company Report #A0090245  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Chest Pain					
TWICE PER DAY		Nervousness					
/ ORAL		Pulmonary Congestion		Terazosin Hydrochloride	C		

Date:05/28/99ISR Number: 3334884-5Report Type:Periodic Company Report #A0090329  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Asthenia					
TWICE PER DAY		Dry Mouth					
/ ORAL		Insomnia		Birth Control	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3334886-9Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0090404

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3334887-0Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0090405

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dry Skin					
TWICE PER DAY							
/ ORAL							

Thyroxine Sodium C

Date:05/28/99ISR Number: 3334891-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0090406

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dry Mouth					
TWICE PER DAY		Insomnia					
/ ORAL	5 DAY	Tremor					

Date:05/28/99ISR Number: 3334892-4Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0090407

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dizziness					
TWICE PER DAY		Insomnia					
/ ORAL				Aspirin	C		
				Tramadol			
				Hydrochloride	C		
				Calcium Salt	C		
				Nitroglycerin	C		
				Salsalate	C		

Date:05/28/99ISR Number: 3334895-XReport Type:Periodic Company Report #A0090408  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Abnormal	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Hyperhidrosis	Professional	Insulin	C		

Date:05/28/99ISR Number: 3335283-2Report Type:Periodic Company Report #A0089570  
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Insomnia					
DAY / ORAL		Tachycardia					
		Trismus					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3335284-4Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0089571

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL							

Date:05/28/99ISR Number: 3335285-6Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0089572

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL							

Date:05/28/99ISR Number: 3335286-8Report Type:Periodic  
 Age:55 YR Gender:Female I/FU:I

Company Report #A0089604

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Sleep Disorder					
TWICE PER DAY							
/ ORAL							

Atenolol	C
Nabumentone	C
Inhaler	C

Date:05/28/99ISR Number: 3335287-XReport Type:Periodic  
 Age:50 YR Gender:Female I/FU:I

Company Report #A0089605

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Rash Erythematous					
DAY / ORAL							

Conjugated Estrogens	C
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Atenolol

C

Date:05/28/99ISR Number: 3335288-1Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #A0089606

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Pruritus		Estrogen	C		
		Urticaria		Armour Thyroid	C		
				Actifed	C		
				Multivitimin	C		

Date:05/28/99ISR Number: 3335289-3Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0089607

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Flushing					
TWICE PER DAY		Insomnia					
/ ORAL							

Date:05/28/99ISR Number: 3335290-XReport Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #A0089608

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Urticaria					
TWICE PER DAY							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

/ ORAL 4 WK

Hormones C

Date:05/28/99ISR Number: 3335291-1Report Type:Periodic Company Report #A0089609  
 Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Amnesia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Mental Impairment					
TWICE PER DAY							

/ ORAL

Date:05/28/99ISR Number: 3335292-3Report Type:Periodic Company Report #A0089610  
 Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nausea	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Sedation					
DAY / ORAL							

Date:05/28/99ISR Number: 3335293-5Report Type:Periodic Company Report #A0089611  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pruritus					
TWICE PER DAY							

/ ORAL

Diltiazem  
 Hydrochloride C  
 Estrogen C

Date:05/28/99ISR Number: 3335294-7Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #A0089612

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Pruritus					
/ ORAL	1	MON					

Blood Pressure Medication C

Date:05/28/99ISR Number: 3335295-9Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0089613

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Drug Dependence	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL							

Amitriptyline Hcl C  
Tylenol No. 3 C  
Human Insulin C  
Naproxen C  
Ticlopidine C  
Oestradiol C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3335296-0Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #A0089614

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3335297-2Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #A0089615

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gingival Bleeding	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Oedema Peripheral					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3335298-4Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #A0089616

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accommodation Disorder	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Agitation					
DAY / ORAL		Feeling Abnormal					
		Nausea					
		Paraesthesia					
		Sensation Of Heaviness					

Date:05/28/99ISR Number: 3335299-6Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #A0089617

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Oestradiol	C		

Date:05/28/99ISR Number: 3335300-XReport Type:Periodic Company Report #A0089618  
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dyspepsia					
TWICE PER DAY		Flatulence					
/ ORAL	2 WK	Gastrointestinal Disorder					

Date:05/28/99ISR Number: 3335301-1Report Type:Periodic Company Report #A0089619  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							
DAY / ORAL				Ascorbic Acid	C		
				Ginseng	C		
				Hypericum	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3335302-3Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #A0089620

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dyspnoea					
TWICE PER DAY		Heart Rate Increased					
/ ORAL		Paraesthesia		Thyroxine Sodium	C		
				Nordette	C		
				Cerivastatin	C		

Date:05/28/99ISR Number: 3335303-5Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #A0089621

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dry Skin					
TWICE PER DAY		Insomnia					
/ ORAL		Irritability					
		Nausea					
		Palpitations					
		Tremor					

Date:05/28/99ISR Number: 3335304-7Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #A0089622

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL				Amlodipine	C		

Hydrochlorothiazide C  
Potassium Chloride C

Date:05/28/99ISR Number: 3335305-9Report Type:Periodic Company Report #A0089623  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Bone Pain					
TWICE PER DAY							
/ ORAL				Conjugated Estrogens	C		

Date:05/28/99ISR Number: 3335306-0Report Type:Periodic Company Report #A0089624  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Urticaria					
TWICE PER DAY							
/ ORAL				Digoxin	C		
				Atenolol	C		
				Prempro	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3335307-2Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0089635

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dry Mouth Erectile Dysfunction	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3335308-4Report Type:Periodic  
 Age:58 YR Gender:Male I/FU:I

Company Report #A0089636

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL

Aspirin C

Date:05/28/99ISR Number: 3335309-6Report Type:Periodic  
 Age:50 YR Gender:Male I/FU:I

Company Report #A0089637

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Agitation Anorexia Cold Sweat Headache Muscle Twitching Nightmare	Consumer	Zyban Tablet - Zyban Diavan Omeprazole Oxycodone Hydrochloride	PS C C C		ORAL

Date:05/28/99ISR Number: 3335310-2Report Type:Periodic Company Report #A0089638  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3335311-4Report Type:Periodic Company Report #A0089639  
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL	6	DAY					

Thyroid  
Blood Pressure  
Medication C  
C

Date:05/28/99ISR Number: 3335313-8Report Type:Periodic Company Report #A0089640  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3335315-1Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #A0089641

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Fatigue	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3335317-5Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #A0089642

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL	10 DAY	Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3335322-9Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #A0089643

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dry Mouth	Health	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL							
		Ejaculation Disorder	Professional				
		Erectile Dysfunction					
		Insomnia		Minoxidil	C		
		Urticaria		Clarithromycin	C		
				Entex	C		

Date:05/28/99ISR Number: 3335323-0Report Type:Periodic  
Age:61 YR Gender:Male I/FU:I

Company Report #A0089901

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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150 MG /  
TWICE PER DAY  
/ ORAL  
Dermatitis  
Urticaria  
Consumer  
Zyban Tablet - Zyban PS  
ORAL

Date:05/28/99ISR Number: 3335325-4Report Type:Periodic Company Report #A0089902  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Nausea	Health	Nicotine Patch	SS		
INTRADERMAL	INTRADERMAL	Nervousness	Professional				
		Rash Erythematous	Other				

Date:05/28/99ISR Number: 3335326-6Report Type:Periodic Company Report #A0089903  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dry Mouth					
TWICE PER DAY		Insomnia					
/ ORAL		Lip Disorder		Ciprofloxacin Hcl	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3335327-8Report Type:Periodic  
 Age:76 YR Gender:Male I/FU:I

Company Report #A0089904

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dry Mouth					
TWICE PER DAY		Insomnia					
/ ORAL				Terbutaline Sulphate	C		
				Theophylline	C		
				Salbutamol Sulphate	C		
				Ipratropium Bromide	C		
				Zinc Oxide	C		

Date:05/28/99ISR Number: 3335331-XReport Type:Periodic  
 Age:56 YR Gender:Male I/FU:I

Company Report #A0089905

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Diazepam	C		
				Percocet	C		

Date:05/28/99ISR Number: 3335332-1Report Type:Periodic  
 Age:50 YR Gender:Male I/FU:I

Company Report #A0089906

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Major Depression					
DAY / ORAL							

Date:05/28/99ISR Number: 3335333-3Report Type:Periodic Company Report #A0089907  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Nicotine	C		
				Clorazepate			
				Dipotassium	C		
				Metolazone	C		

Date:05/28/99ISR Number: 3335335-7Report Type:Periodic Company Report #A0091739  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Birth Control	C		

Date:05/28/99ISR Number: 3335336-9Report Type:Periodic Company Report #A0091746  
Age:52 YR Gender:Male I/FU:I

Outcome	PT
	Chest Pain
	Increased Appetite



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Muscle Spasms Nausea Sedation	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3335338-2Report Type:Periodic Company Report #A0092020  
 Age: Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
Dose							
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3335340-0Report Type:Periodic Company Report #A0092114  
 Age:45 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Urticaria	Consumer	Zyban Tablet - Zyban	PS		ORAL
Dose							
150 MG/ TWICE							
PER DAY /							
ORAL							

Date:05/28/99ISR Number: 3335342-4Report Type:Periodic Company Report #A0092115  
 Age:50 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose							

150 MG /  
 TWICE PER DAY  
 / ORAL

Chills  
 Dyspnoea  
 Tremor

Consumer

Zyban Tablet - Zyban PS

ORAL

Date:05/28/99ISR Number: 3335343-6Report Type:Periodic Company Report #A0092116  
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Flushing					
TWICE PER DAY		Hyperhidrosis					
/ ORAL	25 DAY	Nausea		Cetirizine Hydrochloride	C		
				Fluticasone Propionate	C		

Date:05/28/99ISR Number: 3335344-8Report Type:Periodic Company Report #A0092144  
 Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /				Betaxolol Hydrochloride	C		
TWICE PER DAY				Ipratropium Bromide	C		
/ ORAL				Digoxin	C		
				Diltiazem	C		
				Warfarin Sodium	C		

Freedom Of Information (FOI) Report

Lorazepam C  
Salbutamol Sulphate C

Date:05/28/99ISR Number: 3335346-1Report Type:Periodic Company Report #A0065235  
Age:66 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Back Disorder Chest Wall Pain Condition Aggravated	Health Professional	Zyban Tablet - Zyban	PS		ORAL
		Depression Insomnia Pruritus Psychomotor Hyperactivity Weight Increased		Atorvastatin Calcium Ascorbic Acid Chromium Salt Aspirin Back Problems Tenderness: Chest Region	C C C C C C C		

Date:05/28/99ISR Number: 3335348-5Report Type:Periodic Company Report #A0066257  
Age:71 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / SINGLE DOSE / ORAL		Conjunctival Hyperaemia Discomfort Eye Disorder	Health Professional	Zyban Tablet - Zyban	PS		ORAL
		Eye Irritation Paraesthesia Rash Erythematous Retinal Vascular Disorder Transient Ischaemic Attack Visual Acuity Reduced Visual Disturbance		Amlodipine	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Constipation	Professional				
TWICE PER DAY		Depression					
/ ORAL		Dizziness		Amlodipine			
		Hypercholesterolaemia		(Formulation			
		Insomnia		Unknown)	SS		
5 MG		Nausea		Aspirin	C		
		Orthostatic Hypotension		Famotidine	C		
		Palpitations		Estropipate	C		
		Syncope		Metoclopramide Hcl	C		
		Weight Decreased		Thyroxine Sodium	C		
				Amitriptyline	C		
				Timolol Maleate	C		
				Brimonidine Tartrate	C		
				Rimexolone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3335351-5Report Type:Periodic  
 Age:38 YR Gender:Male I/FU:F

Company Report #A0073811

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Depression Hallucination Insomnia Nausea	Health Professional	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3335353-9Report Type:Periodic  
 Age:41 YR Gender:Female I/FU:F

Company Report #A0073830

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL		Bronchospasm Dermatitis Pruritus Rales Rash Erythematous Throat Tightness Urticaria	Health Professional	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3335354-0Report Type:Periodic  
 Age:38 YR Gender:Male I/FU:F

Company Report #A0074641

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Burning Sensation Chest Pain Dermatitis Dyspepsia Gastritis Hypersensitivity Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL

Pruritus  
Rash Erythematous  
Skin Irritation  
Swelling

Date:05/28/99ISR Number: 3335359-XReport Type:Periodic Company Report #A0075297  
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Depression	Professional				
TWICE PER DAY		Insomnia					
/ ORAL		Nausea		Hrt	C		
				Multivitamin	C		

Date:05/28/99ISR Number: 3335370-9Report Type:Periodic Company Report #A0076043  
Age:40 YR Gender:Male I/FU:F

Outcome

PT

Dermatitis

Dizziness

Face Oedema

Headache

Hunger

Hypersensitivity

Nausea

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pain Pruritus Pyrexia Sedation	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL			Consumer	Zyban Tablet - Zyban	PS		ORAL
50 MG / AS REQUIRED / ORAL				Imitrex Tablet	SS		ORAL

Date:05/28/99ISR Number: 3335372-2Report Type:Periodic Company Report #A0076416  
Age:66 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dizziness Nausea Vomiting	Health Professional	Zyban Tablet - Zyban	PS		ORAL
				Salbutamol Sulphate	C		

Date:05/28/99ISR Number: 3335374-6Report Type:Periodic Company Report #A0076427  
Age:52 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Hypersensitivity	Health Professional	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3335376-XReport Type:Periodic  
Age:54 YR Gender:Female I/FU:F

Company Report #A0076894

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Hypersensitivity	Professional				
TWICE PER DAY		Urticaria					
/ ORAL							

Date:05/28/99ISR Number: 3335380-1Report Type:Periodic  
Age:67 YR Gender:Female I/FU:F

Company Report #A0076971

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Agitation	Professional				
TWICE PER DAY		Anxiety					
/ ORAL		Dyspnoea Exertional		Cojugated Estrogens	C		
		Insomnia		Cimetidine	C		
		Malaise		Oxybutynin			
		Nervousness		Hydrochloride	C		
		Stress		Multivitamin	C		
		Tremor		Beclomethasone			
				Dipropion.	C		
				...	C		
				...	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3335381-3Report Type:Periodic  
Age:43 YR Gender:Female I/FU:F

Company Report #A0076978

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Diarrhoea	Professional				
TWICE PER DAY		Dizziness					
/ ORAL		Nausea		Ranitidine			
		Tremor		Hydrochloride	C		
				Conjugated Estrogens	C		

Date:05/28/99ISR Number: 3352572-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0091183

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Face Oedema	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/ ORAL			Professional Company Representative				

Date:05/28/99ISR Number: 3352574-XReport Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #A0091184

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ PER		Hyperventilation					
DAY/ ORAL		Nervousness					
		Palpitations					
		Tachycardia					

Date:05/28/99ISR Number: 3352575-1Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #A0091186

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hearing Impaired	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Insomnia					
PER DAY/ ORAL							

Date:05/28/99ISR Number: 3352576-3Report Type:Periodic Company Report #A0091187  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL	17 DAY			Nicotine	C		

Date:05/28/99ISR Number: 3352578-7Report Type:Periodic Company Report #A0091188  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Elevated Mood	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Insomnia					
PER DAY/ ORAL		Tremor		Cetirizine Hydrochloride	C		
				Zafirlukast	C		
				Salbutamol Sulphate	C		
				Fluticasone Propionate	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nicotine C

Date:05/28/99ISR Number: 3352580-5Report Type:Periodic Company Report #A0091189  
 Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:05/28/99ISR Number: 3352581-7Report Type:Periodic Company Report #A0091190  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nightmare	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Psychotic Disorder					
PER DAY/ ORAL							

Date:05/28/99ISR Number: 3352582-9Report Type:Periodic Company Report #A0091191  
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Headache					
PER DAY/ ORAL		Neck Pain		Ascorbic Acid	C		

Date:05/28/99ISR Number: 3352584-2Report Type:Periodic Company Report #A0091192  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG/ TWICE	Feeling Abnormal	Consumer	Zyban Tablet - Zyban	PS	ORAL
PER DAY/ ORAL	Nausea				
	Night Sweats		Simvastatin	C	
	Tremor				

Date:05/28/99ISR Number: 3352586-6Report Type:Periodic Company Report #A0091194  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Influenza Like Illness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ PER		Insomnia					
DAY/ ORAL		Malaise					

Date:05/28/99ISR Number: 3352588-XReport Type:Periodic Company Report #A0091229  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL	2 WK	Face Oedema					
		Oedema					
		Pruritus					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3352589-1Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0091529

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/	ORAL 3 MON	Weight Increased		Nicotine Polacrilex	C		

Date:05/28/99ISR Number: 3352590-8Report Type:Periodic  
 Age:31 YR Gender:Male I/FU:I

Company Report #A0091530

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/	ORAL						

Date:05/28/99ISR Number: 3352592-1Report Type:Periodic  
 Age:57 YR Gender:Female I/FU:I

Company Report #A0091531

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/	TWICE	Major Depression					
PER DAY/	ORAL	Tobacco Abuse					

Date:05/28/99ISR Number: 3352594-5Report Type:Periodic  
 Age:54 YR Gender:Female I/FU:I

Company Report #A0091532

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/	TWICE	Anxiety	Professional				
PER DAY /		Brief Psychotic Disorder	Company				
ORAL		Without Marked Stressors	Representative				
		Psychotic Disorder					

Date:05/28/99ISR Number: 3352595-7Report Type:Periodic Company Report #A0091734  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Depressed Mood	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
		Drug Dependence					
PER DAY/ ORAL							
				Glucophage	C		
				Simvastatin	C		
				Famotidine	C		
				Conjugated Estrogens	C		

Date:05/28/99ISR Number: 3352597-0Report Type:Periodic Company Report #A0091735  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Chest Discomfort	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
		Nausea					
PER DAY/ ORAL							
				Methimazole	C		

Date:05/28/99ISR Number: 3352599-4Report Type:Periodic Company Report #A0091736  
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
		Urticaria					
PER DAY/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3352601-XReport Type:Periodic  
Age:72 YR Gender:Female I/FU:I

Company Report #A0091737

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flushing	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Diltiazem			
				Hydrochloride	C		
				Conjugated Estrogens	C		

Date:05/28/99ISR Number: 3352603-3Report Type:Periodic  
Age:77 YR Gender:Female I/FU:I

Company Report #A0091738

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							
				Verapamil	C		
				Amlodipine	C		
				Warfarin Sodium	C		
				Clonidine			
				Hydrochloride	C		

Date:05/28/99ISR Number: 3352604-5Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0089123

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL							
			Professional				

Date:05/28/99ISR Number: 3352605-7Report Type:Periodic  
Age:28 YR Gender:Male I/FU:I

Company Report #A0089125

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3352606-9Report Type:Periodic Company Report #A0089126  
Age:48 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3352609-4Report Type:Periodic Company Report #A0089127  
Age:50 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Urticaria	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL							



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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3352610-0Report Type:Periodic  
 Age:48 YR Gender:Male I/FU:I

Company Report #A0089128

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ PER							
DAY / ORAL							

Date:05/28/99ISR Number: 3352611-2Report Type:Periodic  
 Age:47 YR Gender:Male I/FU:I

Company Report #A0089179

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Deafness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY /							
ORAL							

Lansoprazole C

Date:05/28/99ISR Number: 3352612-4Report Type:Periodic  
 Age:67 YR Gender:Female I/FU:I

Company Report #A0089184

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/ TWICE							
PER DAY /		Gait Disturbance					
ORAL							

Diazepam C  
 Aspirin C  
 Lisinopril C  
 Imipramine  
 Hydrochloride C  
 Trifluoperazine Hcl C

Date:05/28/99ISR Number: 3352613-6Report Type:Periodic Company Report #A0089190  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL	2 YR	Drug Dependence Major Depression					

Date:05/28/99ISR Number: 3352616-1Report Type:Periodic Company Report #A0089191  
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Diarrhoea					
DAY / ORAL							

Date:05/28/99ISR Number: 3352618-5Report Type:Periodic Company Report #A0089192  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							
				Atenolol	C		
				Amlodipine	C		
				Multivitamin	C		
				Estrogen	C		
				Sulphasalazine	C		

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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3352620-3Report Type:Periodic  
 Age:35 YR Gender:Female I/FU:I

Company Report #A0089193

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Dysgeusia					
PER DAY /							
ORAL							

Thyroid Medication C

Date:05/28/99ISR Number: 3352622-7Report Type:Periodic  
 Age:37 YR Gender:Female I/FU:I

Company Report #A0089194

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flatulence	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY /							
ORAL							

Birth Control C

Date:05/28/99ISR Number: 3352630-6Report Type:Periodic  
 Age:33 YR Gender:Female I/FU:I

Company Report #A0089196

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Dysgeusia					
PER DAY /							
ORAL							

Paroxetine  
 Hydrochloride C  
 Doxycycline C

Date:05/28/99ISR Number: 3352631-8Report Type:Periodic Company Report #A0089197  
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ PER		Dizziness					
DAY/ ORAL				Venlafaxine			
				Hydrochloride	C		
				Conjugated Estrogens	C		

Date:05/28/99ISR Number: 3352632-XReport Type:Periodic Company Report #A0089204  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Drug Ineffective					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3352637-9Report Type:Periodic Company Report #A0089205  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Spasms	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Myalgia					
TWICE PER DAY							
/ ORAL				Claritin- D	C		

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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3352640-9Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0089255

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Malaise	Health	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG / PER			Professional				
DAY / ORAL							

Date:05/28/99ISR Number: 3352641-0Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0089256

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Cephalexin	C
Histussin	C

Date:05/28/99ISR Number: 3352642-2Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0089257

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG / PER		Excitability					
DAY / ORAL				Diazepam	C		

Date:05/28/99ISR Number: 3352643-4Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0089258

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG/ TWICE	Oedema Peripheral	Consumer	Zyban Tablet -Zyban	PS	ORAL
PER DAY /	Rash Erythematous				
ORAL			Paracetamol	C	

Date:05/28/99ISR Number: 3352653-7Report Type:Periodic Company Report #A0088861  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Hypersensitivity	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/ PER		Eyelid Oedema	Professional				
DAY/ ORAL		Pruritus					
		Rash Erythematous					
		Rash Papular					
		Rubella					
		Scar					

Date:05/28/99ISR Number: 3352655-0Report Type:Periodic Company Report #A0088864  
 Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ PER		Diarrhoea					
DAY/ ORAL		Influenza Like Illness					
		Pruritus					
		Swelling					
		Urticaria					



150 MG/ TWICE	Confusional State	Consumer	Zyban Tablet - Zyban	PS	ORAL
	Dry Mouth				
PER DAY/ ORAL					
Date:05/28/99ISR Number: 3352664-1Report Type:Periodic Company Report #A0088998					
Age:70 YR Gender:Male I/FU:I					
Outcome	PT	Report Source	Product	Role	Manufacturer
Dose	Duration				Route
150 MG/ TWICE	Dermatitis	Health	Zyban Tablet - Zyban	PS	ORAL
PER DAY /	Pruritus	Professional			
ORAL	Urticaria				
			Triamcinolone Acetonide	C	

Date:05/28/99ISR Number: 3352666-5Report Type:Periodic Company Report #A0088999					
Age:48 YR Gender:Male I/FU:I					
Outcome	PT	Report Source	Product	Role	Manufacturer
Dose	Duration				Route
150 MG/ PER	Agitation	Consumer	Zyban Tablet - Zyban	PS	ORAL
DAY/ ORAL	Asthenia				
	Insomnia		Arthritis Medication	C	
	Tremor		Amoxicillin Trihydrate	C	



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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3352669-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0089027

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/	ORAL	Haemorrhage	Health	Zyban Tablet - Zyban	PS		ORAL
		Menstruation Irregular Metrorrhagia	Professional	Medroxyprogesterone Ace.	C		

Date:05/28/99ISR Number: 3352671-9Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0089088

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/	PER	Dysgeusia	Consumer	Zyban Tablet - Zyban	PS		ORAL
	DAY / ORAL	Insomnia					
		Nausea Neck Pain		Loestrin	C		

Date:05/28/99ISR Number: 3352674-4Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #A0089089

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/	TWICE	Delusion	Consumer	Zyban Tablet - Zyban	PS		ORAL
	PER DAY/ ORAL	Insomnia					
		Paranoia		Alprazolam Ibuprofen	C C		

Date:05/28/99ISR Number: 3352677-XReport Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #A0089090

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/	TWICE	Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL

PER DAY / Insomnia  
ORAL Irritability  
Birth Control C

Date:05/28/99ISR Number: 3352712-9Report Type:Periodic Company Report #A0089091  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Migraine					
DAY/ ORAL		Sedation		Cetirizine			
		Tremor		Hydrochloride	C		

Date:05/28/99ISR Number: 3352715-4Report Type:Periodic Company Report #A0089092  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Urticaria	Health	Nicotine Patch	SS		
INTRADERMAL	INTRADERMAL		Professional				
			Other				



Drug Ineffective Consumer Zyban Tablet - Zyban PS ORAL  
150 MG/ TWICE  
PER DAY/ ORAL  
Date:05/28/99ISR Number: 3352726-9Report Type:Periodic Company Report #A0089121  
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Hypotension					
PER DAY /							
ORAL				Isosorbide	C		
				Frusemide	C		
				Fluvastatin Sodium	C		

Date:05/28/99ISR Number: 3352727-0Report Type:Periodic Company Report #A0089122  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Stomatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY /							
ORAL				Conjugated Estrogens	C		
				Glucosamine			
				Chondroitin C	C		

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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3353692-2Report Type:Periodic  
Age:65 YR Gender:Female I/FU:I

Company Report #A0085853

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Nausea					
TWICE PER DAY							
/ ORAL				Ramipril	C		
				Diltiazem			
				Hydrochloride	C		
				Atenolol	C		

Date:05/28/99ISR Number: 3353693-4Report Type:Periodic  
Age:67 YR Gender:Female I/FU:I

Company Report #A0085854

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Agitation					
DAY / ORAL		Anxiety		Conjugated Estrogens	C		
		Hallucination		Ibuprofen	C		
		Insomnia		Beclomethasone			
		Nausea		Dipropion	C		
		Nervousness		Ipratropium Bromide	C		
		Panic Reaction					

Date:05/28/99ISR Number: 3353694-6Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #A0085855

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Nausea					
DAY / ORAL							

Date:05/28/99ISR Number: 3353695-8Report Type:Periodic Company Report #A0085856  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pruritus					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3353696-XReport Type:Periodic Company Report #A0086067  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Catatonia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Decreased Appetite					
TWICE PER DAY		Feeling Of Relaxation					
/ ORAL		Paraesthesia		Conjugated Estrogens	C		
		Paralysis		Multivitamin	C		
				Hypericum	C		

Date:05/28/99ISR Number: 3353697-1Report Type:Periodic Company Report #A0086105  
Age:68 YR Gender:Male I/FU:I

Outcome	PT
	Dysgeusia
	Dysphonia

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Freedom Of Information (FOI) Report

		Headache Insomnia				
Dose	Duration		Report Source	Product	Role	Manufacturer
			Consumer	Zyban Tablet - Zyban	PS	
150 MG/ TWICE PER DAY/ ORAL				Bromfenex	C	
				Vitamin E	C	
				Amlodipine	C	
				Atrovastatin Calcium	C	

Date:05/28/99ISR Number: 3353698-3Report Type:Periodic Company Report #A0086106  
Age:39 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer
Outcome	Duration		Consumer	Zyban Tablet - Zyban	PS	
Dose		Dermatitis				ORAL
150 MG / TWICE PER DAY / ORAL		Oedema Peripheral		Paracetamol	C	

Date:05/28/99ISR Number: 3353699-5Report Type:Periodic Company Report #A0086107  
Age:55 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer
Outcome	Duration		Consumer	Zyban Tablet - Zyban	PS	
Dose		Disturbance In Attention				ORAL
150 MG / TWICE PER DAY / ORAL		Feeling Abnormal		Fexofenadine Hydrochoride	C	

Date:05/28/99ISR Number: 3353700-9Report Type:Periodic Company Report #A0086108  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dermatitis					
TWICE PER DAY		Insomnia					
/ ORAL		Pruritus					
		Systemic Lupus					
		Erythematosis					

Date:05/28/99ISR Number: 3353701-0Report Type:Periodic Company Report #A0086109  
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Neck Pain	Professional				
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3353702-2Report Type:Periodic Company Report #A0086110  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL							



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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3353703-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0086111

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Drug Ineffective	Consumer Health Professional Other	Zyban Tablet - Zyban   Nicotine (Formulation Unknown)	PS   SS		ORAL

Date:05/28/99ISR Number: 3353704-6Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #A0086112

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Nausea	Consumer	Zyban Tablet - Zyban  Co-Trimoxazole	PS  C		ORAL

Date:05/28/99ISR Number: 3353705-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0086113

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL	WK	Dermatitis Drug Effect Decreased Pain Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3353706-XReport Type:Periodic  
 Age:35 YR Gender:Female I/FU:I

Company Report #A0086114

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dissociation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							
DAY / ORAL							

Date:05/28/99ISR Number: 3353707-1Report Type:Periodic Company Report #A0086115  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							
DAY / ORAL							

Date:05/28/99ISR Number: 3353708-3Report Type:Periodic Company Report #A0086116  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Influenza Like Illness					
TWICE PER DAY		Insomnia					
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3353710-1Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0086117

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Constipation	Professional				
TWICE PER DAY							
/ ORAL				Oestradiol	C		
				Multivitamin	C		

Date:05/28/99ISR Number: 3353712-5Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0086118

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Fatigue					
TWICE PER DAY		Headache					
/ ORAL		Urticaria					

Date:05/28/99ISR Number: 3353713-7Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0086198

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Angioneurotic Oedema	Health	Zyban Tablet - Zyban	PS		ORAL
ORAL	2 WK	Urticaria	Professional Company Representative				

Date:05/28/99ISR Number: 3353913-6Report Type:Periodic  
Age:27 YR Gender:Female I/FU:I

Company Report #A0083642

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		DERMATITIS	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Dry Mouth					
DAY / ORAL		Insomnia		Sleeping Medication	C		

Date:05/28/99ISR Number: 3353914-8Report Type:Periodic Company Report #A0083643  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		AGGRESSION	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Mood Altered					
TWICE PER DAY		Paraesthesia					
/ ORAL							

Date:05/28/99ISR Number: 3353916-1Report Type:Periodic Company Report #A0083644  
 Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		DERMATITIS	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Pruritus					
PER DAY /							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3353918-5Report Type:Periodic  
 Age:44 YR Gender:Male I/FU:I

Company Report #A0083645

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Early Morning Awakening					
TWICE PER DAY		Feeling Abnormal					
/ ORAL		Insomnia					
		Sleep Disorder					

Date:05/28/99ISR Number: 3353920-3Report Type:Periodic  
 Age:49 YR Gender:Male I/FU:I

Company Report #A0083646

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Anxiety					
TWICE PER DAY		Disturbance In Attention					
/ ORAL		Insomnia					

Date:05/28/99ISR Number: 3353922-7Report Type:Periodic  
 Age:79 YR Gender:Male I/FU:I

Company Report #A0083647

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis		Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL		Hypersensitivity					

Date:05/28/99ISR Number: 3353923-9Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #A0083858

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG/ TWICE	Aggression	Consumer	Zyban Tablet - Zyban	PS	ORAL
PER DAY /	Agitation				
ORAL	Anxiety				
	Disturbance In Attention				
	Irritability				

Date:05/28/99ISR Number: 3353924-0Report Type:Periodic Company Report #A0083864  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Chest Discomfort	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Headache					
/ ORAL		Nausea					

Date:05/28/99ISR Number: 3353926-4Report Type:Periodic Company Report #A0083867  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG / PER		Nausea	Health	Zyban Tablet - Zyban	PS		ORAL
DAY / ORAL		Vision Blurred	Professional				
				Desogen	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3353951-3Report Type:Periodic  
 Age:61 YR Gender:Female I/FU:I

Company Report #A0083874

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Rash Erythematous	Professional				
TWICE PER DAY		Urticaria					
/ ORAL				Digoxin	C		
				Propranolol			
				Hydrochloride	C		
				Omeprazole	C		
				Hydrochlorothiazide	C		
				Losartan Potassium	C		
				Conjugated Estrogens	C		

Date:05/28/99ISR Number: 3353984-7Report Type:Periodic  
 Age:55 YR Gender:Female I/FU:I

Company Report #A0083870

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Insomnia					
DAY / ORAL				Conjugated Estrogens	C		
				Carisoprodol	C		

Date:05/28/99ISR Number: 3353986-0Report Type:Periodic  
 Age:66 YR Gender:Female I/FU:I

Company Report #A0083875

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoaesthesia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3353989-6Report Type:Periodic Company Report #A0083879  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Skin Odour Abnormal					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3353991-4Report Type:Periodic Company Report #A0083885  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER							
DAY/ ORAL							

Date:05/28/99ISR Number: 3353992-6Report Type:Periodic Company Report #A0083888  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Influenza Like Illness					
TWICE PER DAY		Insomnia					
/ ORAL				Tetracycline	C		
				Oestradiol	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3353995-1Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0083893

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dyspnoea					
TWICE PER							
DAY/ ORAL	3 WK			Cefprozil (Formulation Unknown)	SS		

Date:05/28/99ISR Number: 3353997-5Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0083894

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Burning Sensation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Stomatitis					
TWICE PER DAY							
/ ORAL				Naproxen	C		
				Conjugated Estrogens	C		

Date:05/28/99ISR Number: 3353998-7Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #A0083895

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache Nausea	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Atenolol	C		
				Enalapril Maleate	C		

Date:05/28/99ISR Number: 3354002-7Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #A0083896

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diplopia	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Discomfort	Professional				
SINGLE DOSE /		Dyspnoea					
ORAL		Nervousness		Salbutamol Sulphate	C		
		Pharyngeal Oedema		Alprazolam	C		
		Vision Blurred		Zafirlukast	C		
				Progesterone	C		
				Conjugated Estrogens	C		

Date:05/28/99ISR Number: 3354004-0Report Type:Periodic  
Age:59 YR Gender:Female I/FU:I

Company Report #A0083897

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Asthenia	Professional				
TWICE PER DAY		Tremor	Company				
/ ORAL			Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354010-6Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #A0082955

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /		Dry Mouth					
TWICE PER DAY		Headache					
/ ORAL		Nervousness					
				Birth Control	C		

Date:05/28/99ISR Number: 3354012-XReport Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #A0082956

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG/ TWICE		Urticaria					
PER DAY /							
ORAL							

Date:05/28/99ISR Number: 3354014-3Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0082958

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /		Agitation					
TWICE PER DAY		Disturbance In Attention					
/ ORAL							

Date:05/28/99ISR Number: 3354016-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0082960

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG / ORAL							

Date:05/28/99ISR Number: 3354022-2Report Type:Periodic Company Report #A0082962  
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Ciprofloxacin Hcl	C		

Date:05/28/99ISR Number: 3354023-4Report Type:Periodic Company Report #A0082964  
 Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
		Pruritus	Professional				
		Urticaria					

Date:05/28/99ISR Number: 3354026-XReport Type:Periodic Company Report #A0083090  
 Age:67 YR Gender:Male I/FU:I

Outcome	PT
	Amnesia
	Confusional State
	Encephalopathy
	Mental Disorder



150 MG/ ORAL      Drug Ineffective      Consumer      Zyban Tablet - Zyban    PS    Zyban    ORAL  
Health      Nicotine      C  
Professional  
Other

Date:05/28/99ISR Number: 3354038-6Report Type:Periodic      Company Report #A0083632  
Age:70 YR    Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Blood Pressure Increased	Consumer	Zyban Tablet- Zyban	PS	Zyban	ORAL
TWICE PER DAY		Chest Discomfort					
/ ORAL		Diarrhoea					
		Dizziness		Gabapentin	C		
		Headache					
		Heart Rate Increased					
		Muscle Spasms					

Date:05/28/99ISR Number: 3354041-6Report Type:Periodic      Company Report #A0083633  
Age:33 YR    Gender:Female      I/FU:I

Outcome      PT  
Confusional State  
Drug Effect Decreased  
Hangover

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Headache Hyperphagia Pain	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG / PER DAY / ORAL							

Date:05/28/99ISR Number: 3354043-XReport Type:Periodic Company Report #A0083634  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG / TWICE PER DAY / ORAL		Insomnia Pruritus Swelling					

Date:05/28/99ISR Number: 3354047-7Report Type:Periodic Company Report #A0083635  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Generalised	Health	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG / ORAL			Professional Company Representative				

Date:05/28/99ISR Number: 3354051-9Report Type:Periodic Company Report #A0083636  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Health	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
TWICE PER DAY / ORAL							
Date:05/28/99ISR Number: 3354052-0Report Type:Periodic Company Report #A0083637 Age:31 YR Gender:Female I/FU:I							
150 MG /		Dry Skin	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
TWICE PER DAY / ORAL							
Birth Control Ibuprofen							
C C							
Date:05/28/99ISR Number: 3354055-6Report Type:Periodic Company Report #A0083638 Age:36 YR Gender:Female I/FU:I							
150 MG/ TWICE		Dermatitis	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
PER DAY / ORAL							
Pruritus							



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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354057-XReport Type:Periodic  
 Age:30 YR Gender:Female I/FU:I

Company Report #A0083639

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dizziness Dyspnoea Palpitations Tremor	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL

Date:05/28/99ISR Number: 3354060-XReport Type:Periodic  
 Age:58 YR Gender:Female I/FU:I

Company Report #A0083640

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TIWCE PER DAY / ORAL		Drug Effect Decreased Excitability Nervousness	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
				Thyroxine Sodium Hydroxyzine Hydrochloride Clonazepam	C C C		

Date:05/28/99ISR Number: 3354062-3Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:I

Company Report #A0083641

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL		Dermatitis Pruritus	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
				Thyroxine Sodium	C		

Date:05/28/99ISR Number: 3354065-9Report Type:Periodic Company Report #A0082507  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Pruritus					
PER DAY/ ORAL		Urticaria					

Date:05/28/99ISR Number: 3354068-4Report Type:Periodic Company Report #A0082508  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Dizziness					
PER DAY/ ORAL		Dry Mouth					
		Tachycardia					

Date:05/28/99ISR Number: 3354074-XReport Type:Periodic Company Report #A0082510  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Arthralgia	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Drug Hypersensitivity	Professional				
PER DAY/ ORAL		Joint Swelling		Hyzaar	C		
		Myalgia					
		Pruritus					
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354077-5Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0082553

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Colour Blindness	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG/	ORAL		Professional				

Date:05/28/99ISR Number: 3354079-9Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0082761

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Health	Zyban Tablet - Zyban	PS		ORAL
ORAL		Flushing Malaise Pyrexia	Professional	Buspirone Hydrochloride	C		

Date:05/28/99ISR Number: 3354082-9Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0082762

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Health	Zyban Tablet- Zyban	PS		ORAL
ORAL		Flushing Malaise Pyrexia	Professional				

Date:05/28/99ISR Number: 3354085-4Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0082763

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Health	Zyban Tablet - Zyban	PS		ORAL
ORAL		Flushing Malaise Pyrexia	Professional				

Date:05/28/99ISR Number: 3354088-XReport Type:Periodic Company Report #A0082903  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
ORAL	1 WK						

Date:05/28/99ISR Number: 3354091-XReport Type:Periodic Company Report #A0082918  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Glossodynia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE		Tongue Ulceration					
PER DAY/ ORAL							

Date:05/28/99ISR Number: 3354094-5Report Type:Periodic Company Report #A0082923  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL				Naproxen Sodium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354096-9Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0082926

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Urticaria	Health Professional Company Representative	Zyban Tablet - Zyban   Lotrel	PS   C		ORAL

Date:05/28/99ISR Number: 3354098-2Report Type:Periodic  
Age:84 YR Gender:Female I/FU:I

Company Report #A0082929

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dizziness	Health Professional	Zyban Tablet - Zyban	PS		ORAL
				Multivitamin	C		
				Metoprolol Succinate	C		
				Aspirin	C		
				Atorvastatin Calcium	C		
				Lisinopril	C		
				Thyroxine Sodium	C		
				Nabumetone	C		
				Isosorbide Mononitrate	C		

Date:05/28/99ISR Number: 3354100-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0082938

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Malaise	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3354103-3Report Type:Periodic Company Report #A0082940  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Generalised	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3354105-7Report Type:Periodic Company Report #A0082942  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Pain	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3354108-2Report Type:Periodic Company Report #A0082946  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG/ ORAL		Stomach Discomfort					

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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354110-0Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #A0082949

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperbilirubinaemia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3354112-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0082951

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eyelid Oedema	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Migraine					
TWICE PER DAY							
/ ORAL							

Salbutamol Sulphate C

Date:05/28/99ISR Number: 3354115-XReport Type:Periodic  
Age:74 YR Gender:Female I/FU:I

Company Report #A0082953

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Propranolol  
Hydrochloride C  
Cimetidine C  
Maxzide C  
Thyroxine Sodium C  
Griseofulvin C  
Triazolam C

Date:05/28/99ISR Number: 3354357-3Report Type:Periodic Company Report #A0087769  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flatulence	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG ORAL							

Date:05/28/99ISR Number: 3354362-7Report Type:Periodic Company Report #A0087770  
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Fluticasone  
Propionate C

Date:05/28/99ISR Number: 3354364-0Report Type:Periodic Company Report #A0087771  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dysgeusia					
TWICE PER DAY		Nausea					
/ ORAL							

Fluvastatin Sodium C



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Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354366-4Report Type:Periodic  
Age:62 YR Gender:Male I/FU:I

Company Report #A0087772

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3354368-8Report Type:Periodic  
Age:66 YR Gender:Male I/FU:I

Company Report #A0087773

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Dristan	C		
				Multivitamin	C		

Date:05/28/99ISR Number: 3354372-XReport Type:Periodic  
Age:67 YR Gender:Male I/FU:I

Company Report #A0087774

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY		Nervousness					
/ ORAL		Paraesthesia					
				Fluticasone			
				Propionate	C		
				Salmeterol Xinafoate	C		
				Ipratropium Bromide	C		

Date:05/28/99ISR Number: 3354374-3Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0087775

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Hypersensitivity	Professional				
TWICE PER DAY		Pruritus					
/ ORAL		Rash Erythematous		Cyclobenzaprine	C		
		Skin Nodule		Gentamicin	C		
		Swelling		Ibuprofen	C		
				Nizatidine	C		
				Misoprostol	C		
				Terbinafine	C		
				Methotrexate	C		

Date:05/28/99ISR Number: 3354376-7Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #A0087777

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alcoholic Hangover	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Feeling Abnormal					
TWICE PER DAY							
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354377-9Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0087778

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Flushing	Professional				
TWICE PER DAY		Insomnia					
/ ORAL							

Date:05/28/99ISR Number: 3354379-2Report Type:Periodic  
Age:68 YR Gender:Male I/FU:I

Company Report #A0087779

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgraphia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Headache					
TWICE PER DAY		Insomnia					
/ ORAL		Tremor		Simvastatin	C		
				Aspirin	C		

Date:05/28/99ISR Number: 3354381-0Report Type:Periodic  
Age:63 YR Gender:Male I/FU:I

Company Report #A0087780

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Headache					
DAY / ORAL		Nausea		Glucophage	C		
				Glipizide	C		
				Clonazepam	C		

Date:05/28/99ISR Number: 3354385-8Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #A0087781

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Drug Dependence Face Oedema	Consumer	Zyban Tablet - Zyban	PS		ORAL
				Diavan	C		

Date:05/28/99ISR Number: 3354387-1Report Type:Periodic Company Report #A0087782  
 Age:65 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Disturbance In Attention Insomnia Tremor	Health Professional	Zyban Tablet - Zyban	PS		ORAL
				Ibuprofen	C		

Date:05/28/99ISR Number: 3354390-1Report Type:Periodic Company Report #A0087898  
 Age:35 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / SEE TEXT / ORAL		Dyspnoea Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354392-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0087902

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Pressure Increased Nervousness Tremor	Consumer	Zyban Tablet - Zyban	PS		

Date:05/28/99ISR Number: 3354396-2Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #A0087971

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL	2 DAY	Vision Blurred	Health Professional	Zyban Tablet - Zyban	PS		ORAL
				Conjugated Estrogens	C		

Date:05/28/99ISR Number: 3354398-6Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0087972

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
				Paroxetine Hydrochloride	C		

Date:05/28/99ISR Number: 3354401-3Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0087973

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Amenorrhoea	Consumer	Zyban Tablet - Zyban	PS		ORAL

Oligomenorrhoea

TWICE PER DAY

/ ORAL

Date:05/28/99ISR Number: 3354404-9Report Type:Periodic Company Report #A0087974  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vomiting	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /							

TWICE PER DAY

/ ORAL

Date:05/28/99ISR Number: 3354406-2Report Type:Periodic Company Report #A0087975  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Swelling	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /			Professional				

TWICE PER DAY

/ ORAL

Quinapril	
Hydrochloride	C
Salbutamol Sulphate	C

Hydrocodone	C
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354512-2Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0088232

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Disturbance In Attention					
TWICE PER DAY		Nervousness					
/ ORAL				Alprazolam	C		

Date:05/28/99ISR Number: 3354514-6Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #A0088233

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Dermatitis					
DAY / ORAL		Dyspepsia					
		Pruritus					

Date:05/28/99ISR Number: 3354520-1Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0088234

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL				Fluoxetine			
				Hydrochloride	C		
				Naproxen	C		
				Omeprazole	C		
				Loratadine	C		

Date:05/28/99ISR Number: 3354521-3Report Type:Periodic Company Report #A0088235  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							
DAY / ORAL							
				Thyroxine Sodium	C		
				Birth Control	C		

Date:05/28/99ISR Number: 3354523-7Report Type:Periodic Company Report #A0088236  
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Atorvastatin Calcium	C		
				Zestoretic	C		
				Multivitamin	C		

Date:05/28/99ISR Number: 3354526-2Report Type:Periodic Company Report #A0088237  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Drug Ineffective					
TWICE PER DAY							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

/ ORAL

Date:05/28/99ISR Number: 3354530-4Report Type:Periodic Company Report #A0088238  
 Age:37 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Crying	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Dissociation					
/ ORAL		Hypokinesia					
		Vomiting					

Date:05/28/99ISR Number: 3354531-6Report Type:Periodic Company Report #A0088239  
 Age:46 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL				Nasal (Unspecified)	C		

Date:05/28/99ISR Number: 3354534-1Report Type:Periodic Company Report #A0088240  
 Age:34 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Blood Pressure Increased	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Headache					
/ ORAL		Heart Rate Increased					
		Nausea		Sertraline Hydrochloride	C		

Date:05/28/99ISR Number: 3354548-1Report Type:Periodic Company Report #A0088241  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Drug Dependence	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL	1	WK					

Date:05/28/99ISR Number: 3354550-XReport Type:Periodic Company Report #A0088242  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Blood Testosterone Decreased	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							
/ ORAL		Erectile Dysfunction					

Date:05/28/99ISR Number: 3354552-3Report Type:Periodic Company Report #A0088243  
 Age:32 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER		Aggression Disturbance In Attention Emotional Disorder Sedation	Consumer	Zyban Tablet - Zyban	PS		ORAL
DAY / ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354556-0Report Type:Periodic  
 Age:58 YR Gender:Male I/FU:I

Company Report #A0088244

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3354557-2Report Type:Periodic  
 Age:42 YR Gender:Female I/FU:I

Company Report #A0088274

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dizziness					
TWICE PER DAY		Drug Effect Decreased					
/ ORAL		Headache		Multivitamin	C		
		Insomnia					
		Middle Insomnia					
		Nausea					
		Tremor					

Date:05/28/99ISR Number: 3354559-6Report Type:Periodic  
 Age:37 YR Gender:Male I/FU:I

Company Report #A0088416

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Pruritus					
TWICE PER DAY		Urticaria					
/ ORAL							

Date:05/28/99ISR Number: 3354561-4Report Type:Periodic Company Report #A0088419  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Oestradiol	C		

Date:05/28/99ISR Number: 3354563-8Report Type:Periodic Company Report #A0088420  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Nausea					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3354566-3Report Type:Periodic Company Report #A0088421  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Migraine	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							
DAY / ORAL				Birth Control	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354568-7Report Type:Periodic  
Age:79 YR Gender:Female I/FU:I

Company Report #A0088422

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tremor	Consumer	Zyban Tablet - Zyban	PS		ORAL
	150 MG / ORAL			Prednisone	C		
				Thyroxine Sodium	C		
				Lorazepam	C		

Date:05/28/99ISR Number: 3354575-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0088423

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash Erythematous	Consumer	Zyban Tablet - Zyban	PS		ORAL
	150 MG /						
	TWICE PER DAY						
	/ ORAL						

Date:05/28/99ISR Number: 3354580-8Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #A0087996

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Flatulence	Consumer	Zyban Tablet - Zyban	PS		ORAL
	150 MG/ TWICE						
	PER DAY	Insomnia					
				Atorvastatin Calcium	C		
				Digoxin	C		
				Orciprenaline			
				Sulphate	C		

Date:05/28/99ISR Number: 3354582-1Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0087997

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Cough	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Libido Decreased					
TWICE PER DAY							

Date:05/28/99ISR Number: 3354584-5Report Type:Periodic Company Report #A0087998  
 Age:51 YR Gender:Male I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Dizziness	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Vomiting					
TWICE PER DAY							

Diltiazem  
 Hydrochloride C  
 Atenolol C  
 Doxazosin Mesylate C  
 Omeprazole C  
 Nitroglycerin C

Date:05/28/99ISR Number: 3354613-9Report Type:Periodic Company Report #A0087999  
 Age:60 YR Gender:Male I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Vomiting	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /			Professional				
TWICE PER DAY							

Terazosin  
 Hydrochloride C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354615-2Report Type:Periodic  
Age:63 YR Gender:Male I/FU:I

Company Report #A0088000

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Anxiety		Enalapril Maleate	C		
				Lovstatin	C		

Date:05/28/99ISR Number: 3354616-4Report Type:Periodic  
Age:34 YR Gender:Male I/FU:I

Company Report #A0088001

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG		Overdose	Consumer	Zyban Tablet - Zyban	PS		ORAL
		Rectal Haemorrhage					

Date:05/28/99ISR Number: 3354619-XReport Type:Periodic  
Age:30 YR Gender:Male I/FU:I

Company Report #A0088002

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY							

Date:05/28/99ISR Number: 3354621-8Report Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #A0088003

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Drug Ineffective	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY				Verapamil			

Hydrochloride C  
Hydrochlorothiazide C  
Omeprazole C

Date:05/28/99ISR Number: 3354623-1Report Type:Periodic Company Report #A0088004  
Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Paraesthesia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Rash Erythematous					
TWICE PER DAY							

Date:05/28/99ISR Number: 3354626-7Report Type:Periodic Company Report #A0088027  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Exfoliative	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Hyperaesthesia	Professional				
TWICE PER DAY		Hyperkeratosis		Nicotine	C		
		Myalgia		Vitamin	C		
		Rash Maculo-Papular		Calcium Salt	C		
		Skin Discolouration		Aspirin	C		
		Therapeutic Response		Nicotine Polacrilex	C		
		Unexpected					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354630-9Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #A0088044

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
TWICE PER DAY		Tongue Oedema	Consumer	Zyban Tablet - Zyban	PS		ORAL
				Minocycline Hydrochloride	C		

Date:05/28/99ISR Number: 3354632-2Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #A0088118

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY		Disorientation Dizziness Headache Insomnia	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3354633-4Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #A0088119

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY		Dermatitis	Health Professional	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3354635-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0088120

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG		Hypersensitivity	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3354636-XReport Type:Periodic Company Report #A0088121  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG	1 WK						

Date:05/28/99ISR Number: 3354640-1Report Type:Periodic Company Report #A0088122  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Urticaria					
TWICE PER DAY				Birth Control	C		

Date:05/28/99ISR Number: 3354664-4Report Type:Periodic Company Report #A0088123  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Psoriasis	Professional				
TWICE PER DAY				Baclofen	C		
				Oxycodone			
				Hydrochloride	C		
				Clonazepam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354669-3Report Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #A0088124

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY		Pruritus Urticaria	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3354672-3Report Type:Periodic  
Age:23 YR Gender:Male I/FU:I

Company Report #A0088144

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3354675-9Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0088169

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY		Face Oedema Liver Function Test Abnormal Urinary Tract Infection	Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:05/28/99ISR Number: 3354682-6Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0089259

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY		Agitation Irritability	Consumer	Zyban Tablet - Zyban	PS		ORAL

Tremor

/ ORAL

Aspirin

C

Date:05/28/99ISR Number: 3354686-3Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0089260

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Headache					
TWICE PER DAY							

/ ORAL

Prednisone

C

Salbutamol Sulphate

C

Theophylline

C

Diazepam

C

Date:05/28/99ISR Number: 3354689-9Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #A0089261

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Appetite Disorder	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER		Confusional State					
DAY / ORAL		Insomnia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354692-9Report Type:Periodic  
Age:54 YR Gender:Male I/FU:I

Company Report #A0089262

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dry Mouth	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Insomnia					
/ ORAL		Oliguria					
		Skin Odour Abnormal		Metoprolol	C		

Date:05/28/99ISR Number: 3354695-4Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0089263

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Condition Aggravated	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Dry Mouth					
/ ORAL		Fatigue					
		Hyperglycaemia		Human Int/Long Insulin	C		
				Vicodin	C		

Date:05/28/99ISR Number: 3354697-8Report Type:Periodic  
Age:53 YR Gender:Male I/FU:I

Company Report #A0089264

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Pruritus	Consumer	Zyban Tablet - Zyban	PS		ORAL
				Warfarin Sodium	C		

Date:05/28/99ISR Number: 3354700-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0089276

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Breast Discharge	Health	Zyban Tablet - Zyban	PS		ORAL
ORAL			Professional Company Representative				

Date:05/28/99ISR Number: 3354701-7Report Type:Periodic Company Report #A0089369  
 Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Sexual Dysfunction					
TWICE PER DAY							
/ ORAL							

Date:05/28/99ISR Number: 3354702-9Report Type:Periodic Company Report #A0089375  
 Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthma	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Atrovastatin Calcium	C		
				Metoprolol	C		
				Ipratropium Bromide	C		



150 MG /  
 TWICE PER DAY  
 / ORAL

Anxiety  
 Drug Withdrawal Syndrome

Consumer

Zyban Tablet - Zyban PS

ORAL

Cardiovascular  
 Medication C

Date:05/28/99ISR Number: 3354712-1Report Type:Periodic Company Report #A0089490  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							
DAY / ORAL							

Salbutamol Sulphate C  
 Steroid C

Date:05/28/99ISR Number: 3354713-3Report Type:Periodic Company Report #A0089491  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / ORAL							
		Pruritus		Multivitamins	C		
		Urticaria					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/99ISR Number: 3354714-5Report Type:Periodic  
 Age:32 YR Gender:Female I/FU:I

Company Report #A0089492

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Muscle Spasms	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Psychomotor Hyperactivity					
/ ORAL							

Date:05/28/99ISR Number: 3354716-9Report Type:Periodic  
 Age:52 YR Gender:Female I/FU:I

Company Report #A0089493

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dermatitis	Health	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Pruritus	Professional				
/ ORAL		Urticaria					

Date:05/28/99ISR Number: 3354717-0Report Type:Periodic  
 Age:41 YR Gender:Female I/FU:I

Company Report #A0089494

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Choking	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Insomnia					
/ ORAL				Thyroid Medication	C		

Date:05/28/99ISR Number: 3354718-2Report Type:Periodic  
 Age:50 YR Gender:Female I/FU:I

Company Report #A0089495

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL				Fluoxetine Hydrochloride	C		

Date:05/28/99ISR Number: 3354720-0Report Type:Periodic Company Report #A0089569  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban Tablet - Zyban	PS		ORAL
150 MG / PER							
DAY / ORAL				Salbutamol Sulphate Steroid	C C		

Date:06/01/99ISR Number: 3274163-8Report Type:Expedited (15-DaCompany Report #A0075934  
 Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Foreign	Zyban Tablet-Zyban	PS		ORAL
Life-Threatening		Dizziness	Health				
150 MG/TWICE		Dyspnoea	Professional				
Hospitalization -		Pruritus					
PER DAY/ORAL		Respiratory Disorder					
Initial or Prolonged		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/01/99ISR Number: 3274165-1Report Type:Expedited (15-DaCompany Report #A0092791  
Age:43 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG /TWICE	Dyspnoea	Foreign	Zyban Tablet - Zyban	PS		ORAL
Initial or Prolonged PER DAY/ ORAL	Laryngitis	Consumer				

Date:06/01/99ISR Number: 3275117-8Report Type:Expedited (15-DaCompany Report #A0075655  
Age:49 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG /	Amnesia	Foreign	Zyban Tablet - Zyban	PS		ORAL
Initial or Prolonged TWICE PER DAY	Dyspnoea	Consumer				
/ ORAL	Emotional Disorder					
	Grand Mal Convulsion Status Epilepticus					

Date:06/01/99ISR Number: 3275405-5Report Type:Expedited (15-DaCompany Report #A0093420  
Age:42 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG /	Chest Pain	Consumer	Zyban Tablet- Zyban	PS		ORAL
Initial or Prolonged TWICE PER DAY	Dermatitis					
	Dyspnoea					
	Hypoaesthesia					
	Pain In Jaw					
	Pharyngeal Oedema					
	Proctalgia					
	Pruritus					
	Urethral Pain					
	Urticaria					
	White Blood Cell Count Increased					

Date:06/01/99ISR Number: 3275421-3Report Type:Expedited (15-DaCompany Report #A0093419

Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain	Foreign	Zyban Tablets -			
		Insomnia	Consumer	Zyban	PS		ORAL
150 MG /							
TWICE PER DAY							

Date:06/03/99ISR Number: 3275549-8Report Type:Expedited (15-DaCompany Report #A0093694

Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Grand Mal Convulsion	Consumer	Wellbutrin Tablet-			
Initial or Prolonged		Spinal Fracture		Controlled Release	PS		ORAL
150 MG/PER							
Disability							
DAY/ ORAL	1	MON					

Date:06/03/99ISR Number: 3275550-4Report Type:Expedited (15-DaCompany Report #A0090630

Age:30 YR Gender:Female I/FU:F

Outcome  
Life-Threatening  
Hospitalization -  
Initial or Prolonged

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Disability

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Alcohol Poisoning Coma	Health Professional	Wellbutrin Tablet- Controlled Release	PS		ORAL
ORAL		Disorientation	Company Representative	Ethanol Liquid	SS		ORAL
		Dizziness Intentional Misuse Metabolic Encephalopathy Nystagmus Status Epilepticus					

Date:06/03/99ISR Number: 3355153-3Report Type:Periodic Company Report #1998UW48296  
Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Paraesthesia	Health Professional	Tenormin Zyban	PS SS		ORAL
150 MG BID PO				Quinapril Hydrochloride Metronidazole Dipyridamole Allopurinol	C C C C		

Date:06/05/99ISR Number: 3277700-2Report Type:Direct Company Report #  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral		Welbutrin 150mg Sr Glaxo Wellcome	PS	Glaxo Wellcome	ORAL
Required Intervention to 150MG 2 X DAY Prevent Permanent ORAL Impairment/Damage				Ritalin	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Heart Valve Replacement International Normalised Ratio Decreased	Health Professional Other	Wellbutrin (Formulation Unknown)	PS		ORAL
100 MG/PER							
DAY/ORAL				Mirtazapine	SS		ORAL
60 MG/PER							
DAY/ORAL				Warfarin Sodium	C		
				Atenolol	C		
				Beclomethasone			
				Dipropion	C		
				Clonazepam	C		
				Docusate Sodium	C		
				Terazosin			
				Hydrochloride	C		
				Lisinopril	C		
				Nitrofurantoin	C		
				Metronidazole	C		
				Simvastatin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/07/99ISR Number: 3277400-9Report Type:Expedited (15-DaCompany Report #A0086682  
 Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Level Above	Health	Lamictal Tablet	PS		ORAL
25 MG/ AT		Therapeutic	Professional				
NIGHT ORAL		Tremor	Other	Wellbutrin (Formulation Unknown)	SS		
150 MG				Valproic Acid (Formulation Unknown)	SS		ORAL
2000 MG/AT							
NIGHT/ORAL				Olanzapine	C		
				Paroxetine	C		
				Hydrochloride	C		
				Clonazepam	C		
				Liothyronine Sodium	C		
				Thyroxine Sodium	C		

Date:06/07/99ISR Number: 3277401-0Report Type:Expedited (15-DaCompany Report #A0093717  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Foreign	Zyban Tablet - Zyban	PS		ORAL
150 MG/TWICE		Difficulty In Walking	Consumer				
PER DAY ORAL		Myalgia Pain					

Date:06/09/99ISR Number: 3278626-0Report Type:Direct Company Report #  
 Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged	Completed Suicide Disturbance In Attention Judgement Impaired Memory Impairment	Wellbutrin - Dose And Type Unknown	PS
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Date:06/10/99ISR Number: 3280027-6Report Type:Direct Company Report #  
Age:52 YR Gender:Male I/FU:I

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	Joint Swelling Rash Pruritic Urticaria	Health Professional	Zyban Prednisone Allegra Hydroxyzine Doxycycline	PS C C C C		

Date:06/10/99ISR Number: 3280754-0Report Type:Expedited (15-DaCompany Report #A0094450  
Age:47 YR Gender:Male I/FU:I

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL Initial or Prolonged	Aortic Injury Road Traffic Accident	Consumer	Zyban Tablet - Zyban	PS		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/10/99ISR Number: 3280771-0Report Type:Expedited (15-DaCompany Report #A0093619

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Anomaly Of External Ear Congenital Cleft Lip And Palate	Study Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL
ORAL		Complications Of Maternal Exposure To Therapeutic Drugs Congenital Diaphragmatic Hernia Foetal Growth Retardation Hemivertebra Hypoplastic Left Heart Syndrome Limb Malformation Microcephaly Premature Baby					

Date:06/10/99ISR Number: 3280791-6Report Type:Expedited (15-DaCompany Report #A0093970

Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG/TWICE PER DAY/ORAL		Aneurysm	Foreign Consumer	Zyban Tablet - Zyban	PS		ORAL

Date:06/10/99ISR Number: 3281160-5Report Type:Direct

Age:57 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG PO		Tongue Oedema		Glaxo Wellcome - Wellbutrin Prinivil Hctz Premarin	PS C C C	Glaxo Wellcome	ORAL

Date:06/11/99ISR Number: 3281087-9Report Type:Direct  
Age:49 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150 MG PO BID	Dermatitis		Zyban (Bupropion)	PS		ORAL
Initial or Prolonged	Hypersensitivity Pruritus Rash Erythematous		Septra	SS		

Date:06/11/99ISR Number: 3281245-3Report Type:Direct  
Age:16 YR Gender:Female I/FU:I

Company Report #

Outcome  
Death  
Life-Threatening  
Hospitalization -  
Initial or Prolonged  
Disability  
Required  
Intervention to  
Prevent Permanent

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Auricular Swelling	Foreign	Zyban Tablet - Zyban	PS		ORAL
150 MG /		Dermatitis	Health				
TWICE PER DAY		Dyspnoea	Professional				
/ ORAL		Eyelid Oedema					
		Oedema					
		Oedema Peripheral					
		Pruritus					
		Pyrexia					
		Rash Papular					
		Speech Disorder					
		Urticaria					

Date:06/14/99ISR Number: 3282754-3Report Type:Expedited (15-DaCompany Report #A0090688  
Age:33 YR Gender:Female I/FU:F

Outcome	PT
Disability	Amnesia
	Aphasia
	Cerebrovascular Accident
	Choking
	Cognitive Deterioration
	Condition Aggravated

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Hemiplegia Hypoaesthesia Medication Error	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Poverty Of Speech	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

Date:06/14/99ISR Number: 3336395-XReport Type:Periodic Company Report #WAES 98060990  
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arrhythmia Chest Pain	Consumer	Tab Losartan Potassium Unk	PS		ORAL
PO 150 MG/DAILY/PO		Drug Interaction Palpitations		Zyban 150 Mg	SS		ORAL
				Composition Unspecified Nitroglycerin	C C		

Date:06/15/99ISR Number: 3284190-2Report Type:Direct Company Report #  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 100MG BID		Convulsion Fall		Bupropriion Glaxo Wellcome	PS	Glaxo Wellcome	
		Head Injury		Chloral Hydrate Divalproex Ibuprofen	C C C		

Date:06/16/99ISR Number: 3284272-5Report Type:Expedited (15-DaCompany Report #A0079132  
 Age:16 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Health	Wellbutrin			
Life-Threatening		Completed Suicide	Professional	Tablet-Controlled			
Hospitalization -		Convulsion	Company	Release	PS		ORAL
150 MG / SEE							
Initial or Prolonged		Loss Of Consciousness	Representative				
TEXT / ORAL							
Other		Overdose		Paroxetine			
				Hydrochloride	C		

Date:06/16/99ISR Number: 3284303-2Report Type:Expedited (15-DaCompany Report #A0094003  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Dry Mouth	Foreign	Zyban Tablet - Zyban	PS		ORAL
150 MG /							
TWICE PER DAY		Feeling Of Despair	Health				
/ ORAL		Headache	Professional				
		Myopia					
		Vision Blurred					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/16/99ISR Number: 3284304-4Report Type:Expedited (15-DaCompany Report #A0090360

Age:46 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - Initial or Prolonged 150 MG / PER Other DAY / ORAL		Health Professional Company  Representative	Wellbutrin Tablet-Controlled Release  Prempro (Formulation Unknown)	PS   SS		ORAL   ORAL
PER DAY /						
ORAL						

Date:06/16/99ISR Number: 3284305-6Report Type:Expedited (15-DaCompany Report #A0089648

Age:75 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 100 MG / PER DAY / ORAL		Health Professional Other	Wellbutrin (Formulation Unknown)	PS		ORAL
International Normalised Ratio Decreased			Mirtazapine Tablet	SS		ORAL
60 MG / PER DAY / ORAL						
			Warfarin Sodium Atenolol Beclomethasone Dipropion Clonazepam Docusate Sodium Terazosin Hydrochloride Lisinopril Nitrofurantoin Metronidazole Simvastatin	C C C C C C C C C C C C		

Date:06/16/99ISR Number: 3284791-1Report Type:Expedited (15-DaCompany Report #A0087945  
Age:35 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150 MG / PER	Anxiety	Health	Zyban Tablet-Zyban	PS		ORAL
Initial or Prolonged DAY / ORAL	Convulsion	Professional				
Disability	Epistaxis		Claritin-D	C		
	Loss Of Consciousness		Nicotine	C		
	Mouth Haemorrhage					
	Nasal Congestion					

Date:06/17/99ISR Number: 3286264-9Report Type:Expedited (15-DaCompany Report #A0094827  
Age:45 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other 150 MG /	Arthralgia	Foreign	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY	Dyspnoea	Consumer				
/	Oedema					
	Serum Sickness					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/18/99ISR Number: 3286403-XReport Type:Expedited (15-DaCompany Report #A0091793

Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Aggression	Foreign	Wellbutrin			
Hospitalization -		Depression	Health	Tablet-Controlled			
Initial or Prolonged		Paranoia	Professional	Release	PS		ORAL
100 MG / PER		Physical Assault					
Other		Psychotic Disorder		Clavulin	C		
DAY / ORAL		Suicidal Ideation					

Date:06/21/99ISR Number: 3288383-XReport Type:Expedited (15-DaCompany Report #A0092189

Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Asthma	Foreign	Zyban Tablet-	PS	Zyban	ORAL
150 MG /		Burning Sensation	Health				
Initial or Prolonged		Drug Hypersensitivity	Professional				
TWICE PER DAY		Eyelid Oedema					
/ ORAL		Facial Palsy					
		Pruritus					
		Urticaria					

Date:06/21/99ISR Number: 3288502-5Report Type:Expedited (15-DaCompany Report #1999-06-0585

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Interaction	Study	Wellbutrin Tablets	PS		ORAL
ORAL		Grand Mal Convulsion	Consumer	Intron A (Interferon			
			Other	Alfa-2b Recombina			
				Injectable)	SS		
5 MU TIW				Rebetol (Ribavirin)			

1200 MG DAILY

Capsules

SS

ORAL

ORAL

Date:06/21/99ISR Number: 3288509-8Report Type:Expedited (15-DaCompany Report #A0093320

Age:54 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Agitation Cerebrovascular Accident Crying	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/TWICE PER DAY/ORAL	Emotional Distress Facial Palsy Hypoaesthesia Irritability Medication Error Mental Disorder Transient Ischaemic Attack		Azithromycin Ranitidine Hydrochloride Nasonex	C C C C		

Date:06/23/99ISR Number: 3289562-8Report Type:Direct

Company Report #

Age:75 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 37.5MG PO TID	Grand Mal Convulsion	Health Professional	Wellbutrin 75 Mg (Glaxo- Wellcome) Prilosec	PS C	Glaxo-Wellcome	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Cardizem Cd	C
Lasix	C
Kcl	C
Acetaminophen	C
Darvocet	C
Buspar	C
Oxistat	C
Zostrix	C
Mentax	C

Date:06/23/99ISR Number: 3289788-3Report Type:Expedited (15-DaCompany Report #A0095039  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Foreign	Zyban Tablet - Zyban	PS		ORAL
Other		Dysphonia	Consumer				
150 MG /		Fall					
TWICE PER DAY		Head Injury					
/ ORAL		Muscle Rigidity					
		Nausea					
		Nervousness					

Date:06/24/99ISR Number: 3293878-9Report Type:Periodic Company Report #A001-002-003251  
 Age:85 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Cerebrovascular Accident	Consumer	Aricept (Donepezil)	PS		ORAL
5MG (1 IN		Hypertension	Health				
Initial or Prolonged			Professional	Wellbutrin			
1D), PER ORAL				(Bupropion Hcl)			
				(Amfebutamone			
				Hydrochloride)	SS		
NA				Aspirin			
				(Acetylsalicylic			
				Acid)	C		

Atenolol (Atenolol) C  
 Diltiazem  
 (Diltiazem) C  
 Timoptic (Timolol  
 Maleate) C  
 Digoxin (Digoxin) C

Date:06/25/99ISR Number: 3291868-3Report Type:Expedited (15-DaCompany Report #A0094919  
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Metastases To Lung	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/TWICE PER DAY/ORAL	5 DAY			Omeprazole Alprazolam Dilantin	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/25/99ISR Number: 3291906-8Report Type:Expedited (15-DaCompany Report #A0094798

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY		Paranoia	Foreign Consumer	Zyban Tablet-Zyban	PS		ORAL

Date:06/25/99ISR Number: 3291926-3Report Type:Expedited (15-DaCompany Report #A0079132

Age:16 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Life-Threatening Hospitalization - 150 MG / SEE Initial or Prolonged TEXT Disability Other		Brain Damage Cardiac Arrest Convulsion Loss Of Consciousness Overdose Tachycardia	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release Paroxetine Hydrochloride	PS C		ORAL

Date:06/25/99ISR Number: 3291929-9Report Type:Expedited (15-DaCompany Report #B0067691

Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG/ UNK/ ORAL		Medication Error Road Traffic Accident	Foreign Health Professional	Zyban Tablet - Zyban	PS		ORAL

Date:06/28/99ISR Number: 3292685-0Report Type:Expedited (15-DaCompany Report #A0085805

Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health	Wellbutrin			

150 MG TWICE	Professional Company	Tablet-Controlled Release	PS	ORAL
PER DAY ORAL 1 MON	Representative	Tamoxifen	C	

Date:06/28/99ISR Number: 3292762-4Report Type:Expedited (15-DaCompany Report #A0094985  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Alkaline Phosphatase Increased Blood Bilirubin Increased	Foreign Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / PER DAY / ORAL		Blood Creatinine Increased Blood Lactate Dehydrogenase Increased Gamma-Glutamyltransferase Increased		Cyclosporin	C		

Date:06/29/99ISR Number: 3294540-9Report Type:Expedited (15-DaCompany Report #A0095026  
 Age:51 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Anaphylactic Shock Asthenia Heart Rate Increased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Hypersensitivity Hypotension Palpitations	Report Source	Product	Role	Manufacturer	Route
150 MG /		Skin Disorder	Consumer	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Swelling					

Date:06/29/99ISR Number: 3294543-4Report Type:Expedited (15-DaCompany Report #A0094393  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Condition Aggravated	Health	Zyban Tablet - Zyban	PS		
3 WK							
Initial or Prolonged		Hallucination Psychotic Disorder	Professional Company Representative	Haloperidol	C		

Date:06/30/99ISR Number: 3294734-2Report Type:Direct Company Report #  
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Chest Pain		Bupropion Sr 100 Mg	PS		ORAL
100 MG OD PO							
Initial or Prolonged		Dermatitis Gastrooesophageal Reflux Disease Muscle Disorder Rash Pruritic Urticaria		Ginkgo Biloba Nizatidine Estropipate Vitamin E Multi-Vit	C C C C C		

Date:06/30/99ISR Number: 3295013-XReport Type:Expedited (15-DaCompany Report #A0084631  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Tablet -			

150 MG /	Depressed Level Of	Professional	Controlled Release	PS	ORAL
TWICE PER DAY	Consciousness				
			Oral Contraceptive	C	
			Fluoxetine		
			Hydrochloride	C	
			Amitriptyline	C	
			Sumatriptan		
			Succinate	C	
			Fiorinal	C	
			Fexofenadine		
			Hydrochlorid	C	
			Fluticasone		
			Propionate	C	

Date:06/30/99ISR Number: 3295035-9Report Type:Expedited (15-DaCompany Report #A0089368  
Age:43 YR Gender:Male I/FU:F

Outcome	PT
Death	Agitation
Hospitalization -	Alcohol Interaction
Initial or Prolonged	Atrioventricular Block
Other	Blood Alcohol Increased
	Cardiac Arrest
	Completed Suicide
	Convulsion



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Intentional Misuse Loss Of Consciousness Tachycardia	Report Source	Product	Role	Manufacturer	Route
200 TABLET/ SINGLE DOSE/ ORAL			Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL

Ethanol (Formulation  
Unknown) SS

Date:06/30/99ISR Number: 3295039-6Report Type:Expedited (15-DaCompany Report #A0085804  
Age:55 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG/ TWICE PER DAY/ ORAL		Myocardial Infarction	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL

Sildenafil Citrate C

Date:06/30/99ISR Number: 3295040-2Report Type:Expedited (15-DaCompany Report #A0093402  
Age:47 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG/ PER DAY/ ORAL		Liver Disorder Metastases To Liver	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

Tamoxifen C  
Paroxetine  
Hydrochloride C

Date:06/30/99ISR Number: 3295341-8Report Type:Expedited (15-DaCompany Report #A0095803  
Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional Company Representative	Zyban Tablet - Zyban	PS	Zyban	

Date:06/30/99ISR Number: 3295342-XReport Type:Expedited (15-DaCompany Report #A0095677  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG		Abdominal Distension	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
Initial or Prolonged		Insomnia					

Date:07/01/99ISR Number: 3294898-0Report Type:Direct Company Report #  
Age: Gender:Female I/FU:I

Outcome	PT
Death	Anomaly Of External Ear Congenital Cleft Lip And Palate

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Complications Of Maternal Exposure To Therapeutic Drugs	Report Source	Product	Role	Manufacturer	Route
		Congenital Diaphragmatic Hernia Congenital Hand Malformation Congenital Osteodystrophy Congenital Spinal Cord Anomaly Foetal Growth Retardation Hypoplastic Left Heart Syndrome Microcephaly Multiple Congenital Abnormalities Premature Baby		Wellbutrin	PS		

Date:07/01/99ISR Number: 3297085-5Report Type:Expedited (15-DaCompany Report #A0094919  
Age:45 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG /		Convulsion Metastases To Central Nervous System	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
	TWICE PER DAY	5 YR			Omeprazole	C		

Date:07/01/99ISR Number: 3341444-9Report Type:Periodic Company Report #USA008973  
Age:38 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	10 MG	OD	Agitation Confusional State	Health Professional	Meridia	PS		ORAL
	PO							

150 MG Gait Disturbance Other Zyban SS ORAL  
 UNK PO Restlessness  
 Date:07/01/99ISR Number: 3346622-0Report Type:Periodic Company Report #USA008215  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Consumer	Meridia	PS		ORAL
10 MG OD PO		Central Nervous System Stimulation	Other	Wellbutrin	SS		
		Feeling Of Relaxation		Lithium	C		
		Therapeutic Response		Serzone	C		
		Unexpected		Synthroid	C		

Date:07/06/99ISR Number: 3297829-2Report Type:Expedited (15-DaCompany Report #A0095664  
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bronchospasm	Health	Zyban Tablet - Zyban	PS		ORAL
Hospitalization - 150 MG/TWICE		Dermatitis	Professional				
Initial or Prolonged PER DAY ORAL		Loss Of Consciousness	Company Representative	Nitrofurantoin (Formulation			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Unknown)	SS
Clonazepam	C
Hydroxyzine	
Hydrochloride	C
Oxybutynin	
Hydrochloride	C
Thyroxine Sodium	C
Phenazopyridine Hcl	C

Date:07/06/99ISR Number: 3297832-2Report Type:Expedited (15-DaCompany Report #A0095671  
 Age:37 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 100 MG/TWICE PER DAY ORAL	Convulsion Drug Level Below Therapeutic Fall Head Injury	Health Professional	Wellbutrin (Formulation Unknown) Chloral Hydrate Semisodium Valproate Ibuprofen	PS C C C		ORAL

Date:07/06/99ISR Number: 3297998-4Report Type:Expedited (15-DaCompany Report #A0095916  
 Age:80 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/DAY / Initial or Prolonged UNK / ORAL 3 WK	Abdominal Pain Constipation Faecaloma Ileus Paralytic Vomiting	Literature	Wellbutrin Tablet Lithium Salt Diltiazem Ranitidine Hydrochloride Aspirin Calcium Salt	PS C C C C C C		ORAL

Date:07/06/99ISR Number: 3298001-2Report Type:Expedited (15-DaCompany Report #A0095916  
 Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/DAY / Initial or Prolonged UNK / ORAL	3 WK	Abdominal Pain Constipation Faecaloma Ileus Paralytic Vomiting	Literature	Wellbutrin Tablet  Lithium Salt Diltiazem Ranitidine Hydrochloride Aspirin Calcium Salt	PS  C C C C C		ORAL

Date:07/06/99ISR Number: 3298373-9Report Type:Expedited (15-DaCompany Report #A0095889  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG /  TWICE PER DAY  / ORAL		Convulsion Fatigue Hypoglycaemia Tremor	Foreign  Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/06/99ISR Number: 3300658-4Report Type:Periodic Company Report #107251  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1MG 4 PER DAY ORAL		Balance Disorder Hypoaesthesia Motion Sickness	Consumer	Klonopin	PS		ORAL
ORAL				Zyban (Bupropion Hydrochloride) Premarin	SS C		ORAL

Date:07/08/99ISR Number: 3299256-0Report Type:Expedited (15-DaCompany Report #A0095816  
 Age:6 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 37.5 MG Initial or Prolonged ALTERNATE DAYS ORAL		Abnormal Behaviour Candidiasis Dysphemia Kidney Infection Pharyngitis Streptococcal Tongue Disorder	Consumer	Wellbutrin Tablet	PS		ORAL

Date:07/12/99ISR Number: 3301398-8Report Type:Direct Company Report #  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 100MG SR BID Prevent Permanent PO Impairment/Damage		Dermatitis Pruritus		Bupropion 100mg Sr Tab Gabapentin Multivitamin/Mineral s	PS C C		ORAL

Date:07/12/99ISR Number: 3301440-4Report Type:Expedited (15-DaCompany Report #A0095928

Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Blood Creatine	Health	Wellbutrin			
Initial or Prolonged	Phosphokinase Increased	Professional	(Formulation			
Other	Leukopenia		Unknown)	PS		ORAL
ORAL	Neuroleptic Malignant Syndrome		Dexamphetamine Sulphate			
	Pyrexia		(Formulation			
	Sepsis		Unknown)	SS		
	Serotonin Syndrome					
	Urinary Tract Infection					

Date:07/12/99ISR Number: 3302470-9Report Type:Expedited (15-DaCompany Report #A0096258

Age:42 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Brain Oedema
Initial or Prolonged	Chest Pain
	Coma
	Diabetes Mellitus
	Inadequate Control
	Difficulty In Walking
	Loss Of Consciousness
	Oedema Peripheral

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vomiting

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Consumer	Zyban	PS		ORAL
			Gabapentin	C		
			Troglitazone	C		
			Amlodipine	C		
			Int./Long-Acting			
			Insulin	C		

Date:07/14/99ISR Number: 3303994-0Report Type:Expedited (15-DaCompany Report #A0095916  
Age:80 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG DAY ORAL	3 WK	Abdominal Pain	Literature	Wellbutrin Tablet	PS		ORAL
Initial or Prolonged		Blood Creatinine		Lithium Salt	C		
Increased		Blood Urea Increased		Diltiazem	C		
Constipation		Drug Interaction		Ranitidine			
Drug Interaction		Faecaloma		Hydrochloride	C		
Faecaloma		Ileus Paralytic		Aspirin	C		
Ileus Paralytic		Vomiting		Calcium Salt	C		
Vomiting							

Date:07/14/99ISR Number: 3304022-3Report Type:Expedited (15-DaCompany Report #A0077001  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 150 MG/TWICE PER DAY/ ORAL		Arthralgia	Health Professional	Wellbutrin Tablet- Controlled Release	PS		ORAL
Initial or Prolonged Disability Other		Difficulty In Walking	Company Representative				
Difficulty		Dyspnoea Exertional					
Other		Erythema Multiforme					

Eyelid Oedema  
Face Oedema  
Fall  
Fatigue  
Joint Stiffness  
Migraine  
Pain  
Pain In Extremity  
Proteinuria  
Pruritus  
Serum Sickness  
Stevens-Johnson Syndrome  
Throat Tightness  
Urticaria  
Vomiting

Date:07/14/99ISR Number: 3304023-5Report Type:Expedited (15-DaCompany Report #A0073901  
Age:29 YR Gender:Female I/FU:F

Outcome  
Life-Threatening  
Hospitalization -  
Initial or Prolonged  
Disability

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Arthralgia Decreased Appetite Dyspnoea Exertional	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
PER DAY/ ORAL		Erythema Multiforme	Representative				
		Face Oedema Fall Insomnia Joint Stiffness Migraine Oral Pain Orbital Oedema Pain In Extremity Proteinuria Rash Erythematous Serum Sickness Stevens-Johnson Syndrome Throat Tightness Urticaria Vomiting White Blood Cell Count Increased					

Date:07/15/99ISR Number: 3305168-6Report Type:Expedited (15-DaCompany Report #A0093619

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Life-Threatening Hospitalization - ORAL Initial or Prolonged Congenital Anomaly		Acrochordon Bone Development Abnormal Cleft Lip  Cleft Palate Complications Of Maternal Exposure To Therapeutic Drugs Diaphragmatic Hernia Foetal Growth Retardation Hemivertebra Hypoplastic Left Heart	Study Health Professional	Wellbutrin (Formulation Unknown)	PS		ORAL

Syndrome  
Limb Malformation  
Microcephaly  
Premature Baby

Date:07/15/99ISR Number: 3305172-8Report Type:Expedited (15-DaCompany Report #A0095664  
Age:57 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Anaphylactic Reaction
Initial or Prolonged	Blood Creatine Increased
	Bronchospasm
	Circulatory Collapse
	Depressed Level Of
	Consciousness
	Dizziness
	Dysuria
	Faeces Discoloured
	Hallucination

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
150 MG /		Hypotension Personality Change Rash Maculo-Papular	Health	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		White Blood Cell Count Increased	Professional Company				
/ ORAL		White Blood Cells Urine Positive	Representative	Nitrofurantoin Clonazepam Hydroxyzine Hydrochloride Oxybutynin Hydrochloride Thyroxine Sodium Phenazopyridine Hcl	SS C C C C C C		

Date:07/15/99ISR Number: 3305177-7Report Type:Expedited (15-DaCompany Report #A0093422  
Age:49 YR Gender:Female I/FU:I

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
150 MG /		Disability Joint Swelling	Foreign Health	Zyban Tablet - Zyban	PS		ORAL
TWICE PER DAY		Rash Erythematous	Professional				
/ ORAL		Rash Generalised Rash Papular Red Blood Cell Sedimentation Rate Increased Rheumatoid Arthritis Rheumatoid Factor Positive Serum Sickness Urticaria White Blood Cell Count Increased	Company Representative	Nicotine	C		

Date:07/16/99ISR Number: 3305591-XReport Type:Expedited (15-DaCompany Report #A0096492  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ TWICE		Dysphagia	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL
Initial or Prolonged PER DAY /		Dyspnoea					
ORAL		Embolism					
		Oedema Peripheral Swelling		Paracetamol	C		

Date:07/16/99ISR Number: 3305593-3Report Type:Expedited (15-DaCompany Report #A0096292  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /PER DAY / ORAL	2 DAY			Nefazodone Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/19/99ISR Number: 3305577-5Report Type:Direct  
 Age:21 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG QD PO	Blood Pressure Increased		Zyban 150mg Glaxo	PS	Glaxo	ORAL
		Drug Withdrawal Syndrome					
		Headache					
		Insomnia					

Date:07/20/99ISR Number: 3307410-4Report Type:Expedited (15-DaCompany Report #A0093420  
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG /	Cardiomegaly	Health	Zyban Tablet - Zyban	PS		ORAL
Initial or Prolonged	TWICE PER DAY	Chest Discomfort	Professional				
		Chest Pain					
		Dyspnoea					
		Hyperhidrosis					
		Pain In Jaw					
		Proctalgia					
		Pruritus					
		Rash Macular					
		Throat Tightness					
		Urethral Pain					
		Urticaria					
		White Blood Cell Count					
		Increased					

Date:07/20/99ISR Number: 3307424-4Report Type:Expedited (15-DaCompany Report #A0095964  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG/TWICE	Apnoea	Consumer	Zyban Tablet-Zyban	PS		ORAL
Initial or Prolonged	PER DAY/ORAL	Dysarthria					

ORAL	Feeling Hot	Ethanol Liquid	SS	ORAL
	Headache	Fluoxetine		
	Movement Disorder	Hydrochloride	C	
	Nausea	Aspirin	C	
	Syncope			
	Tremor			
	Urinary Incontinence			

Date:07/21/99ISR Number: 3308227-7Report Type:Expedited (15-DaCompany Report #A0090688  
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Amnesia	Consumer	Wellbutrin			
		Aphasia		Tablet-Controlled			
		Cerebral Artery Occlusion		Release	PS		ORAL
150 MG /		Cerebrovascular Accident					
TWICE PER DAY		Choking					
/ ORAL		Craniotomy					
		Hemiplegia					
		Hypoaesthesia					
		Mental Impairment					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/22/99ISR Number: 3308995-4Report Type:Expedited (15-DaCompany Report #A0094985  
 Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Bilirubin Increased Blood Creatinine Increased	Foreign Health Professional	Wellbutrin Tablet-Controlled Release			
150 MCG PER DAY ORAL		Liver Function Test  Abnormal Thrombocythaemia			PS		ORAL
				Cyclosporin Clonidine Dapsone Metoclopramide Acyclovir Mycophanolate Mofetil Omeprazole Sodium Bicarbonate Salbutamol Sulphate Fluticasone Propionate Prednisone Insulin	C C C C C C C C C C C C C C		

Date:07/22/99ISR Number: 3308996-6Report Type:Expedited (15-DaCompany Report #A0096983  
 Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Headache Petit Mal Epilepsy	Foreign Consumer	Wellbutrin Tablet-Controlled Release			
ORAL					PS		ORAL
				Dilantin Semisodium Valproate Ibuprofen	C C C		

Date:07/22/99ISR Number: 3309971-8Report Type:Direct Company Report #  
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urticaria		Bupropion (Zyban)	PS		ORAL
150 MG PO BID							

Date:07/22/99ISR Number: 3309972-XReport Type:Direct Company Report #  
 Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pruritus		Bupropion (Zyban)	PS		ORAL
150 MG PO BID		Urticaria					

Date:07/23/99ISR Number: 3310515-5Report Type:Direct Company Report #  
 Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG BID		Aphasia		Wellbutrin 100mg Tab	PS	Glaxo Wellcome	
Initial or Prolonged		Convulsion		Aspirin	C		
		Monoparesis		Cycrohaptachine	C		
				Thiothiarie	C		
				Licarb	C		
				Dovusate	C		
				Calcium Carbonate	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Multivitamin C

Date:07/27/99ISR Number: 3311855-6Report Type:Expedited (15-DaCompany Report #A0097480

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Study	Zyban Tablet - Zyban	PS	Zyban	ORAL
ORAL		Complications Of Maternal Exposure To Therapeutic Drugs	Health Professional				

Date:07/27/99ISR Number: 3311863-5Report Type:Expedited (15-DaCompany Report #A0090907

Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dermatitis	Health	Zyban Tablet - Zyban	PS	Zyban	ORAL
150 MG /							
Initial or Prolonged		Dysphagia	Professional				
TWICE PER DAY							
/ ORAL		Dyspnoea					
		Face Oedema					
		Tongue Oedema					

Date:07/27/99ISR Number: 3311864-7Report Type:Expedited (15-DaCompany Report #A0094393

Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Hallucination	Health	Zyban Tablet - Zyban	PS	Zyban	ORAL
ORAL	3 WK						
Initial or Prolonged		Schizophrenia	Professional	Amantadine	C		
		Suicidal Ideation	Company Representative	Haloperidol	C		

Date:07/27/99ISR Number: 3311865-9Report Type:Expedited (15-DaCompany Report #A0097494  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY  / ORAL		Aggression  Confusional State  Depression  Eating Disorder Insomnia Mental Impairment Paranoia Personality Change Psychiatric Symptom	Consumer	Zyban Tablet - Zyban	PS	Zyban	ORAL

Date:07/27/99ISR Number: 3311866-0Report Type:Expedited (15-DaCompany Report #A0097409  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death ORAL		Cerebrovascular Accident	Health  Professional Company Representative	Zyban Tablet - Zyban	PS	Zyban	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/27/99ISR Number: 3311867-2Report Type:Expedited (15-DaCompany Report #A0097410

Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Myocardial Infarction	Health	Zyban Tablet - Zyban	PS	Zyban	ORAL
ORAL			Professional Company Representative				

Date:07/27/99ISR Number: 3313893-6Report Type:Direct

Company Report #

Age:15.6 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Prolactin Increased Galactorrhoea	Health Professional	Risperadol 1.5mg Bid, Decreased 1mg Bid, Decreased 0.5mg Bid	PS		
1.5MG BID, DECREASES 1MG BID, DECREASED 0.5MG BID 75MG PO Q AM				Wellbutrin 75mg Q Am	SS		ORAL

Date:07/28/99ISR Number: 3312826-6Report Type:Expedited (15-DaCompany Report #A0097643

Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Coronary Artery Disease	Foreign	Zyban Tablet - Zyban	PS		ORAL
ORAL		Myocardial Infarction	Health Professional				

Age:28 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL	Angioneurotic Oedema  Arthralgia  Dermatitis Difficulty In Walking Ecchymosis Heart Rate Increased Malaise Oedema Peripheral Pain In Extremity Pruritus Serum Sickness Skin Discolouration Urticaria	Foreign  Health  Professional	Zyban Tablet- Zyban   Ibuprofen	PS   C		ORAL

Company Report #

Age:32 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150MG PO X 1T Required Intervention to Prevent Permanent Impairment/Damage	Dissociation  Dysphoria Self Mutilation Suicidal Ideation		Zyban 150 Mg	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/29/99ISR Number: 3313992-9Report Type:Direct  
 Age:34 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG		Face Oedema Heart Rate Increased Rash Pruritic		Wellbutrin /150 Mg /Glaxo Wellcome	PS	Glaxco Wellcome	

Date:07/29/99ISR Number: 3314836-1Report Type:Periodic  
 Age:49 YR Gender:Female I/FU:I

Company Report #A0091454

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG / TWICE PER DAY / ORAL		Convulsion	Health Professional Company Representative	Wellbutrin Tablet -Controlled Release	PS		ORAL

Date:07/29/99ISR Number: 3314841-5Report Type:Periodic  
 Age:24 YR Gender:Female I/FU:I

Company Report #A0091249

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG / TWICE PER DAY / ORAL	14 DAY	Bite Convulsion Lethargy Lip Disorder Myalgia Petit Mal Epilepsy Sedation Syncope Tremor	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release Cyclobenzaprine Hcl Fexofenadine Hydrochlorid Oral Contraceptive	PS    C C C		ORAL

Date:07/29/99ISR Number: 3314849-XReport Type:Periodic Company Report #A0090423  
Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG/ TWICE			Representative				
PER DAY /							
ORAL				Gabapentin	C		

Date:07/29/99ISR Number: 3314853-1Report Type:Periodic Company Report #A0090175  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression Suicidal Ideation	Health Professional Company	Wellbutrin Tablet -Controlled Release	PS		ORAL
UNK / THREE			Representative				
TIMES PER DAY							
/ ORAL				Lithium Salt	C		
				Risperidone	C		
				Propranolol	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/29/99ISR Number: 3314857-9Report Type:Periodic  
 Age:16 YR Gender:Female I/FU:I

Company Report #A0090056

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Condition Aggravated Convulsion Obsessive-Compulsive Disorder	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / TWICE PER DAY / ORAL				Citalopram	C		

Date:07/29/99ISR Number: 3314860-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0089931

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / TWICE PER DAY / ORAL				Zolpidem Tartrate	C		

Date:07/29/99ISR Number: 3314863-4Report Type:Periodic  
 Age:58 YR Gender:Male I/FU:I

Company Report #A0089310

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged UNK/ TWICE PER DAY / ORAL		Dizziness Malaise Pallor Tinnitus	Consumer	Wellbutrin Tablet- Controlled Release	PS		ORAL
				Quinapril			

Hydrochloride C  
Omeprazole C  
Guaiphenesin C

Date:07/29/99ISR Number: 3314866-XReport Type:Periodic Company Report #A0088986  
Age:56 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG / Other TWICE PER DAY  / ORAL	Hallucination Psychotic Disorder	Health Professional  Company  Representative	Wellbutrin Tablet -Controlled Release    Bupropion Hydrochloride	PS     C		ORAL

Date:07/29/99ISR Number: 3314870-1Report Type:Periodic Company Report #A0088918  
Age:51 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG /  TWICE PER DAY  / ORAL	Grand Mal Convulsion	Health Professional	Wellbutrin Tablet -Controlled Release   Fluoxetine Hydrochloride Risperidone	PS    C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/29/99ISR Number: 3314874-9Report Type:Periodic Company Report #A0088578  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Wellbutrin Tablet- Controlled Release	PS		ORAL
UNK / UNK /			Company				
ORAL			Representative				

Date:07/29/99ISR Number: 3315404-8Report Type:Periodic Company Report #A0087647  
 Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Convulsion Road Traffic Accident	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
Other			Company				
150 MG /			Representative				
TWICE PER DAY							
/ ORAL 8 WK				Dexamphetamine	C		

Date:07/29/99ISR Number: 3315411-5Report Type:Periodic Company Report #A0085964  
 Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Burning Sensation Dehydration Dermatitis Exfoliative	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
Other							
150 MG /							
TWICE PER DAY							
/ ORAL				Alprazolam	C		
		Dry Skin					
		Erythema Multiforme					
		Fatigue Fungal Skin Infection Infection					

Pruritus  
Psoriasis  
Rash Generalised

Date:07/29/99ISR Number: 3315418-8Report Type:Periodic Company Report #A0084031  
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Grand Mal Convulsion	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Fluoxetine Hydrochloride	C		
				Clonazepam	C		
				Quetiapine Fumarate	C		

Date:07/29/99ISR Number: 3315424-3Report Type:Periodic Company Report #A0077917  
Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Other		Grand Mal Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /			Representative				
TWICE PER DAY							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

/ ORAL

Lortab	C
Salbutamol Sulphate	C
Triamcinolone	
Acetonide	C
Testosterone	C

Date:07/29/99ISR Number: 3315431-0Report Type:Periodic  
 Age:37 YR Gender:Female I/FU:F

Company Report #A0079654

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Accidental Overdose	Health	Wellbutrin			
Initial or Prolonged	Alcohol Interaction	Professional	Tablet-Controlled			
Other	Convulsion	Company	Release	PS		ORAL

150 MG /

Representative

THREE TIMES

PER DAY ORAL

Ethanol (Formulation	
Unknown)	SS

ORAL

UNK / UNK /

ORAL

Date:07/29/99ISR Number: 3315435-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0096490

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Convulsion	Health	Wellbutrin			
		Professional	Tablet-Controlled			
		Company	Release	PS		ORAL

UNK / UNK /

Representative

ORAL

Date:07/29/99ISR Number: 3315438-3Report Type:Periodic  
 Age:11 YR Gender:Male I/FU:I

Company Report #A0096029

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Convulsion	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / PER							
DAY / ORAL 1 YR							

Date:07/29/99ISR Number: 3315442-5Report Type:Periodic Company Report #A0095870  
 Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / PER							
DAY / ORAL 3 DAY							

Date:07/29/99ISR Number: 3315446-2Report Type:Periodic Company Report #A0095616  
 Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Convulsion	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
UNK / UNK /							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/29/99ISR Number: 3315450-4Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0095478

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	Coma Suicide Attempt	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / PER DAY / ORAL						
40 MG / PER DAY / ORAL			Citalopram	SS		ORAL
			Trazodone	C		

Date:07/29/99ISR Number: 3315453-XReport Type:Periodic  
Age:35 YR Gender:Male I/FU:I

Company Report #A0094080

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	Stevens-Johnson Syndrome	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / TWICE PER DAY / ORAL						
			Paclitaxel	C		

Date:07/29/99ISR Number: 3315457-7Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #A0093073

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other	Dermatitis Pruritus	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / TWICE PER DAY						

/ ORAL

Date:07/29/99ISR Number: 3315459-0Report Type:Periodic Company Report #A0092987  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion Loss Of Consciousness	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL

150 MG / PER

DAY / ORAL

Moexipril  
Hydrochloride C  
Sertraline  
Hydrochloride C

Date:07/29/99ISR Number: 3315461-9Report Type:Periodic Company Report #A0092653  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion Overdose	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL

150 MG /

THREE TIMES

PER DAY ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/29/99ISR Number: 3315463-2Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0092326

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
UNK / UNK /			Representative				
ORAL							

Date:07/29/99ISR Number: 3315464-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0092331

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
UNK / UNK /			Representative				
ORAL							

Date:07/29/99ISR Number: 3315466-8Report Type:Periodic  
 Age:48 YR Gender:Female I/FU:I

Company Report #A0092357

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Micturition Urgency Urinary Retention	Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / PER							
DAY / ORAL							
				Mirtazepine	C		
				Thyroxine Sodium	C		
				Phentermine			
				Hydrochloride	C		
				Valaciclovir	C		

Date:07/29/99ISR Number: 3315468-1Report Type:Periodic Company Report #A0092192  
Age:21 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Grand Mal Convulsion Overdose Respiratory Failure	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
UNK / SEE		Representative				
TEXT / ORAL						

Date:07/29/99ISR Number: 3315470-XReport Type:Periodic Company Report #A0091964  
Age:25 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Other 150 MG /	Convulsion Feeling Abnormal	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
TWICE PER DAY		Representative				
/ ORAL			Medroxyprogesterone Ace. (Formulation Unknown)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/29/99ISR Number: 3315476-0Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #A0091692

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG /			Representative				
TWICE PER DAY							
/ ORAL							

Date:07/30/99ISR Number: 3315130-5Report Type:Expedited (15-DaCompany Report #A0073099  
Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ TWICE		Anxiety	Foreign	Zyban Tablet - Zyban	PS		ORAL
PER DAY /		Chest Pain	Health				
ORAL		Hypoglycaemia	Professional				
		Malaise		Glibenclamide	C		
		Palpitations		Metformin	C		

Date:07/30/99ISR Number: 3315131-7Report Type:Expedited (15-DaCompany Report #A0094919  
Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion	Health	Wellbutrin			
Initial or Prolonged		Metastases To Central Nervous System	Professional Company	Tablet-Controlled Release	PS		ORAL
Other							
150 MG /		Metastases To Lung	Representative				
TWICE PER DAY							
/ ORAL	5 DAY			Omeprazole	C		

Date:07/30/99ISR Number: 3315153-6Report Type:Expedited (15-DaCompany Report #A0088466  
Age:79 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG;	Feeling Jittery Tremor	Health Professional	Zyban Tablet - Zyban	PS		ORAL

TWICE PER

DAY; ORAL

Unknown	C
Ranitidine	
Hydrochloride	C
Thyroxine Sodium	C
Hydrochlorothiazide	C
Digoxin	C
Paracetamol	C
Potassium Salt	C

Date:07/30/99ISR Number: 3315159-7Report Type:Expedited (15-DaCompany Report #A0083735  
Age:23 YR Gender:Female I/FU:I

Outcome	PT
Life-Threatening	Anaphylactoid Reaction Arthralgia Dyspnoea Face Oedema Oedema Peripheral Pruritus

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
150 MG; TWICE		Rash Pruritic Serum Sickness Urticaria	Study Health Professional	Wellbutrin Tablet-Controlled Release	PS		ORAL
PER DAY;							
ORAL				Medroxyprogesterone Ace.	C		

Date:07/30/99ISR Number: 3315160-3Report Type:Expedited (15-DaCompany Report #A0097852  
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Blood Antidiuretic Hormone Increased	Health Professional	Wellbutrin Tablet - Controlled Release	PS		ORAL
		Hyponatraemia Pituitary Tumour	Company Representative	Methocarbamol Carisoprodol	C C		

Date:07/30/99ISR Number: 3315163-9Report Type:Expedited (15-DaCompany Report #A0091712  
Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Neuroleptic Malignant Syndrome	Health Professional	Wellbutrin (Formultion Unknown)	PS		ORAL

Date:07/30/99ISR Number: 3315165-2Report Type:Expedited (15-DaCompany Report #A0095184  
Age:85 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - ORAL	Blood Pressure Increased	Health	Wellbutrin Tablet	PS	ORAL
Initial or Prolonged 5 MG/ PER	Cerebrovascular Accident	Professional	Donepezil Hcl Tablet	SS	ORAL
DAY/ ORAL		Other			
			Aspirin	C	
			Diltiazem	C	
			Timolol Maleate	C	
			Digoxin	C	

Date:07/30/99ISR Number: 3317738-XReport Type:Direct Company Report #  
 Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 ORAL	21 DAY	Dizziness Nausea Swelling		Wellbutrin	PS		ORAL

Date:08/02/99ISR Number: 3316269-0Report Type:Expedited (15-DaCompany Report #A0097619  
 Age:28 YR Gender:Female I/FU:F

Outcome	PT
	Arthralgia
	Dermatitis
	Ecchymosis
	Heart Rate Increased
	Malaise

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Oedema Peripheral Pain In Extremity Pruritus	Foreign				
TWICE PER DAY		Serum Sickness	Health	Zyban Tablet - Zyban	PS		ORAL
/ ORAL		Skin Discolouration	Professional				
		Urticaria		Ibuprofen	C		

Date:08/02/99ISR Number: 3316525-6Report Type:Expedited (15-DaCompany Report #A0092192  
Age:21 YR Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged UNK/ SEE TEXT		Grand Mal Convulsion Intentional Misuse Respiratory Failure Suicide Attempt	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release	PS		ORAL
/ ORAL							

Date:08/02/99ISR Number: 3318345-5Report Type:Periodic Company Report #8-97286-020L  
Age:43 YR Gender:Female I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 20 MG THREE TIMES A DAY ORAL		Cardiovascular Disorder Palpitations Pulmonary Hypertension	Consumer	Pondimin	PS		ORAL
15-30 MG DAILY ORAL 30 MG DAILY ORAL				Phentermine	SS		ORAL
				Redux	SS		ORAL

10-150 MG

Wellbutrin

SS

ORAL

TWICE DAILY

ORAL

Date:08/02/99ISR Number: 3318696-4Report Type:Periodic

Company Report #8-97318-014L

Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 60-120 MG		Cardiovascular Disorder	Consumer	Pondimin	PS		ORAL
Initial or Prolonged DAILY ORAL	86 MON	Pulmonary Hypertension		Effexor (Venlafaxine)	SS		
Required Intervention to 75-225 MG				Prozac (Fluoxetine)	SS		
Prevent Permanent DAILY Impairment/Damage 20-60 MG				Wellbutrin (Bupropion)	SS		
DAILY 150-300 MG				Zoloft (Sertraline)	SS		
DAILY 50 MG DAILY				Ritalin	C		
				Cylert	C		
				Cytomel	C		
				Estraderm	C		
				Amitriptyline	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/09/99ISR Number: 3321789-9Report Type:Expedited (15-DaCompany Report #A0098119A  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cardiac Failure Cardiac Valve Disease Dizziness	Foreign Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
100 MG / TWICE PER DAY/ ORAL							

Date:08/09/99ISR Number: 3321791-7Report Type:Expedited (15-DaCompany Report #A0096005A  
 Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hallucination Insomnia Muscle Spasms Muscle Twitching	Health Professional Company Representative	Wellbutrin Tablet- Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL 3 YR							
				Cyclobenzapine Hcl (Formulation Unknown) (Cyclobenzaprine Hcl)	SS		ORAL
10 MG / TWICE TIMES PER DAY / ORAL							

Date:08/09/99ISR Number: 3321892-3Report Type:Expedited (15-DaCompany Report #A0097683A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged Other	Delirium Muscle Rigidity Muscle Rupture Neuroleptic Malignant Syndrome Pyrexia	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS	ORAL
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Date:08/09/99ISR Number: 3321896-0Report Type:Expedited (15-DaCompany Report #A0095217A  
Age:31 YR Gender:Female I/FU:I

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged  150 MG / PER  DAY / ORAL	Adrenergic Syndrome Chills Dizziness Hyperhidrosis  Hypotension  Muscle Spasms Nausea Tremor	Foreign Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)   Temazepam Buspirone Hydrochloride Venlafaxine Hydrochloride	PS    C C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/16/99ISR Number: 3326617-3Report Type:Expedited (15-DaCompany Report #A0082611A

Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amenorrhoea Convulsion Head Injury Memory Impairment	Health Professional	Wellbutrin Tablet-Controlled Release(Bupropion Hydrochloride)	PS		ORAL
1 TABLET							
TWICE DAILY							

Date:08/17/99ISR Number: 3327979-3Report Type:Expedited (15-DaCompany Report #A0098621A

Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chromaturia Convulsion Difficulty In Walking Dysphagia Feeling Abnormal Insomnia Lethargy Nausea Sedation	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:08/17/99ISR Number: 3327985-9Report Type:Expedited (15-DaCompany Report #A0098738A

Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Atrial Fibrillation Blood Pressure Increased Dizziness Fatigue Headache	Consumer	Wellbutrin Tablet - Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG /							
TWICE PER DAY							

/ ORAL	Insomnia						
	Palpitations			Verapamil		C	
				Hyzaar		C	
				Librium		C	

Date:08/17/99ISR Number: 3327992-6Report Type:Expedited (15-DaCompany Report #A0098933A  
 Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Fatigue	Foreign	Wellbutrin Tablet -			
Hospitalization -		Hepatic Failure	Health	Controlled Release			
Initial or Prolonged		Nausea	Professional	(Bupropion			
				Hydrochloride)	PS		ORAL
150 MG /							
TWICE PER DAY							

/ ORAL

Date:08/17/99ISR Number: 3331448-4Report Type:Periodic Company Report #S99-USA-01116-01  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Coma	Health	Celexa	PS		
20 MG							
Initial or Prolonged		Overdose	Professional	Wellbutrin	SS		
150 MG							
			Company	Trazodone	C		
			Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/24/99ISR Number: 3333249-XReport Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Haemoglobin Decreased White Blood Cell Count Decreased		Bupropion	PS		

Date:08/26/99ISR Number: 3334282-4Report Type:Direct  
 Age:52 YR Gender:Male I/FU:I

Company Report #

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75MG BID 6/7 INCREASED TO 75MG TID 7/9  100MG QAM INCREASED TO 200MG QAM 6/7		Abdominal Pain Upper Anorexia Chromaturia  Jaundice  Pyrexia		Buproprion 75mg Bid 6/7 Increased To 75mg Tid 7/9  Sertraline 100mg Qam Increased To 200mg Qam 6/7	PS    SS		

Date:08/27/99ISR Number: 3336302-XReport Type:Expedited (15-DaCompany Report #A0098621A  
 Age:49 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other  150 MG / TWICE PER DAY		Chromaturia Convulsion Difficulty In Walking  Dysphagia	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

/ ORAL

Feeling Abnormal  
Hypersomnia  
Insomnia  
Lethargy  
Nausea

Date:08/27/99ISR Number: 3336332-8Report Type:Expedited (15-DaCompany Report #A0052441A

Age:13 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Apallic Syndrome	Consumer	Bupropion			
Hospitalization - Initial or Prolonged		Brain Hypoxia Cardio-Respiratory Arrest Condition Aggravated		Hydrochloride Tablet (Bupropion Hydrochloride)	PS		ORAL
3600 MG / SINGLE DOSE		Convulsion					
		Iatrogenic Injury Intentional Misuse Mydriasis Suicidal Ideation		Semisodium Valproate	C		

Date:08/30/99ISR Number: 3337368-3Report Type:Expedited (15-DaCompany Report #A0092653A

Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Grand Mal Convulsion	Health Professional	Wellbutrin Tablet-Controlled			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

150 MG /	Release (Bupropion Hydrochloride)	PS	ORAL
THREE TIMES			
PER DA / ORAL	Methylphenidate Hcl	C	

Date:08/31/99ISR Number: 3338187-4Report Type:Expedited (15-DaCompany Report #A0095803A  
 Age:15 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional Company Representative	Bupropion Hydrochloride Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150MG TWICE							
PER DAY ORAL							

Date:08/31/99ISR Number: 3338188-6Report Type:Expedited (15-DaCompany Report #A0095916A  
 Age:80 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Disability Required		Abdominal Pain Blood Creatinine Increased Blood Urea Increased	Literature Health Professional	Bupropion Hydrochloride Tablet (Bupropion Hydrochloride)	PS		ORAL
75MG DAY							
Intervention to TWICE PER DAY Prevent Permanent ORAL 3 WK Impairment/Damage		Constipation Drug Interaction Faecaloma Ileus Paralytic Nausea Vomiting		Lithium Salt Diltiazem Ranitidine Hydrochloride Aspirin Calcium Salt	C C C C C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other Required 100 MG, TWICE Intervention to A DAY, ORAL; Prevent Permanent UNSPECIFIED, Impairment/Damage UNKNOWN, ORAL;		Anaemia Drug Interaction Haemoglobin Decreased Thrombocytopenia	Foreign Health Professional	Sandimmune Neoral (Cyclosporine)  Adalat (Nifedipine)  Wellbutrin (Amfebutamone Hydrochloride)  Paxil (Paroxetine Hydrochloride) Norvasc (Amlodipine Besilate) Flomax (Morniflumate)	PS  SS  SS  C C C		ORAL  ORAL  ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/01/99ISR Number: 3339102-XReport Type:Expedited (15-DaCompany Report #A0098119A  
 Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Failure	Foreign	Wellbutrin			
		Cardiac Murmur	Health	Tablet-Controlled			
		Cardiac Valve Disease	Professional	Release (Bupropion			
		Condition Aggravated		Hydrochloride)	PS		ORAL
100 MG TWICE							
		Dizziness					
PER DAY ORAL							

Date:09/01/99ISR Number: 3339162-6Report Type:Expedited (15-DaCompany Report #A0099506A  
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion	Health	Wellbutrin			
Initial or Prolonged		Feeling Abnormal	Professional	Tablet-Controlled			
		Overdose	Company	Release (Bupropion			
		Vaginal Candidiasis	Representative	Hydrochloride)	PS		ORAL
150 MG / SEE							
TEXT/ ORAL				Oral Contraceptive	C		

Date:09/01/99ISR Number: 3339166-3Report Type:Expedited (15-DaCompany Report #A0077179A  
 Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Atrial Fibrillation	Consumer	Zyban Tablet - Zyban			
Initial or Prolonged		Influenza Like Illness		(Bupropion			
		Rift Valley Fever		Hydrochloride)	PS		ORAL
PER DAY /							
ORAL							

Date:09/01/99ISR Number: 3341480-2Report Type:Periodic Company Report #A0086021A  
 Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required 150 MG		Dermatitis Dyspnoea Face Oedema	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
Intervention to TWICE PER DAY Prevent Permanent ORAL Impairment/Damage		Oral Mucosal Eruption  Tongue Oedema  Urticaria		Claritin-D Conjugated Estrogens Medroxyprogesterone Ace	C C C		

Date:09/01/99ISR Number: 3341484-XReport Type:Periodic  
Age:38 YR Gender:Male I/FU:F

Company Report #A0086000A

Outcome	PT
Life-Threatening Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage	Anaphylactic Reaction Bronchospasm Dermatitis Dyspnoea Eye Disorder Face Oedema Pharyngeal Oedema Pruritus Respiratory Distress Swelling

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Tongue Oedema Urticaria				
Dose	Duration		Report Source	Product	Role	Manufacturer
			Health Professional	Bupropion Hydrochloride Tablet - Zyban (Bupropion Hydrochloride)	PS	
150 MG						ORAL
TWICE PER DAY						
ORAL						

Date:09/01/99ISR Number: 3341491-7Report Type:Periodic Company Report #A0081646A  
 Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
Other		Insomnia Pruritus Urticaria					
150 MG							
TWIC PER DAY							
ORAL							

Date:09/01/99ISR Number: 3341495-4Report Type:Periodic Company Report #A0098261A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health Professional Company Representative	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
Other		Convulsion					

Date:09/01/99ISR Number: 3341499-1Report Type:Periodic Company Report #A0098165A  
 Age:51 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Amnesia Blood Pressure Increased Cerebrovascular Accident	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
	150 MG		Headache					
		TWICE PER DAY						
ORAL	6	WK			Amlodipine Metoprolol Succinate Valsartan	C C C		

Date:09/01/99ISR Number: 3341502-9Report Type:Periodic Company Report #A0097853A  
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Grand Mal Convulsion	Health Professional Company	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
	150 MG			Representative				
ORAL								

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/01/99ISR Number: 3341505-4Report Type:Periodic  
Age:62 YR Gender:Male I/FU:I

Company Report #A0097223A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG TWICE PER DAY ORAL	Hepatitis	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Maalox	C
Aspirin	C
Atenolol	C
Felodipine	C
Ranitidine	
Hydrochloride	C
Losartan Potassium	C

Date:09/01/99ISR Number: 3341508-XReport Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #A0096229A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150 MG TWICE PER DAY ORAL	Drug Ineffective Photosensitivity Reaction	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Amitriptyline Hcl	C
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Date:09/01/99ISR Number: 3341511-XReport Type:Periodic  
Age:33 YR Gender:Male I/FU:I

Company Report #A0082332A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Convulsion Fall	Foreign Consumer	Zyban Tablet - Zyban (Bupropion			



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) PS ORAL

1 TABLET  
 TWICE PER DAY  
 ORAL

Date:09/01/99ISR Number: 3341536-4Report Type:Periodic Company Report #A0095428A  
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Bite Convulsion Feeling Cold	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG		Haemorrhage					
TWICE PER DAY		Loss Of Consciousness					
ORAL		Tongue Disorder Tongue Haematoma					

Date:09/01/99ISR Number: 3341541-8Report Type:Periodic Company Report #A0094675A  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Stevens-Johnson Syndrome	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG							
TWICE PER DAY							
ORAL							

Date:09/01/99ISR Number: 3341545-5Report Type:Periodic Company Report #A0093712A  
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged	Arthralgia Chest Pain Drug Hypersensitivity	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS	ORAL
150 MG	Joint Swelling				
TWICE PER DAY	Rash Pruritic				
ORAL	Swelling Urticaria				

Date:09/01/99ISR Number: 3341550-9Report Type:Periodic Company Report #A0093374A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Malaise	Consumer	Bupropion Hydrochloride Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Date:09/01/99ISR Number: 3341551-0Report Type:Periodic Company Report #A0093077A  
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other 150 MG		Convulsion	Health Professional Company  Representative	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
TWICE PER DAY				Fluticasone			
ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Propionate C  
 Combivent C  
 Lorazepam C

Date:09/01/99ISR Number: 3341552-2Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0092866A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Hyperhidrosis Nausea Nervousness	Consumer	Bupropion Hydrochloride Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG	PER						
DAY	ORAL	Tremor					

Date:09/01/99ISR Number: 3341558-3Report Type:Periodic  
 Age:34 YR Gender:Male I/FU:I

Company Report #A0092517A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Dermatitis Headache	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG							
TWICE PER DAY		Myalgia					
ORAL		Pain		Diclofenac Sodium	C		

Date:09/01/99ISR Number: 3341564-9Report Type:Periodic  
 Age:39 YR Gender:Male I/FU:I

Company Report #A0092396A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Bupropion Hydrochloride Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
1 TABLET							

TWICE PER DAY

ORAL

Multivitamin C

Date:09/02/99ISR Number: 3339597-1Report Type:Expedited (15-DaCompany Report #A0098933A

Age:26 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Fatigue	Foreign	Wellbutrin			
Hospitalization -		Hepatic Failure	Health	Tablet-Controlled			
Initial or Prolonged		Jaundice	Professional	Release (Bupropion			
150 MG /		Nausea		Hydrochloride)	PS		ORAL

TWICE PER DAY

/ ORAL

Salbutamol Sulphate C  
 Beclomethasone  
 Dipropion C  
 Venlafaxine  
 Hydrochloride C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/07/99ISR Number: 3342133-7Report Type:Expedited (15-DaCompany Report #A0100405A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Lymphoedema Oedema Peripheral Weight Increased	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS	Zyban	ORAL

ORAL

Date:09/07/99ISR Number: 3342135-0Report Type:Expedited (15-DaCompany Report #A0100230A

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Complications Of Maternal Exposure To Therapeutic Drugs Congenital Anomaly	Study Health Professional Company	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		ORAL

ORAL

Representative

Date:09/08/99ISR Number: 3343029-7Report Type:Direct

Age:51 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG BID		Dizziness  Dry Mouth Dysgeusia Headache Nausea		Bupropion (Zyban)	PS		

Date:09/08/99ISR Number: 3343719-6Report Type:Direct

Age:37 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis Bullous Hypoaesthesia		Bupropion 150 Mg (Zyban) - Glaxo			

150MG PO QD X  
 3D THEN 150MG  
 BID

Date:09/09/99ISR Number: 3343508-2Report Type:Direct Company Report #  
 Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Bupropion	PS		ORAL
Other		Depression	Professional				
150MG BID		Drug Ineffective					
ORAL	2 MON			Carbamazepine (Tegretol)	C		
				Oxycodone/Acetaminop hen	C		

Date:09/09/99ISR Number: 3343536-7Report Type:Direct Company Report #  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropoin Sr	PS		
Other		Medication Residue		Zolpidem	C		
100MG TID							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/09/99ISR Number: 3344246-2Report Type:Expedited (15-DaCompany Report #A0100352A

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 75 MG/SEE TEXT/ORAL	Jaundice Liver Function Test Abnormal	Health Professional	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL

Date:09/11/99ISR Number: 3345423-7Report Type:Direct

Company Report #

Age:46 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG SR BID Initial or Prolonged	Agitation Insomnia Irritability Mania Thinking Abnormal		Bupropion	PS		

Date:09/13/99ISR Number: 3345500-0Report Type:Expedited (15-DaCompany Report #A0100543A

Age:63 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Cerebrovascular Accident	Consumer	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		

Date:09/16/99ISR Number: 3347915-3Report Type:Direct

Company Report #

Age:21 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG BID PO 14 DAY	Grand Mal Convulsion		Zyban	PS		ORAL

Initial or Prolonged  
Other

B.C. Pill

C

Date:09/16/99ISR Number: 3349491-8Report Type:Expedited (15-DaCompany Report #A0097683A  
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Confusional State Delirium	Health Professional Company	Welbutrin Tablet-Controlled Release (Bupropion Hydrochloride)			
Required 150 MG / Intervention to TWICE PER DAY Prevent Permanent / ORAL Impairment/Damage		Muscle Atrophy Muscle Rigidity  Neuroleptic Malignant Syndrome  Pyrexia	Representative		PS		ORAL
				Semisodium Valproate Librium Buspirone Hydrochloride Diltiazem Hydrochloride Conjugated Estrogens Thyroxine Sodium Hydrochlorothiazide Omeprazole	C C C C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/20/99ISR Number: 3352405-8Report Type:Expedited (15-DaCompany Report #A0099452A  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State Feeling Abnormal Pancreatic Carcinoma	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /		Suicidal Ideation					
TWICE PER DAY		Tremor					
/ ORAL							

Date:09/20/99ISR Number: 3352406-XReport Type:Expedited (15-DaCompany Report #A0099702A  
 Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization - Initial or Prolonged		Dizziness Dyspnoea Fatigue	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /		Hypophosphataemia					
Disability TWICE PER DAY		Mutism					
/ ORAL		Paraesthesia		Phenoxyethylpenicil lin Nicotine	C C		

Date:09/22/99ISR Number: 3354160-4Report Type:Direct Company Report #  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated		Wellbutrin Sr-B/W	PS		ORAL
150MG PO BID		Crying		Premarin	C		
		Depression		Fosamax	C		
				Beconase	C		
				Allegra	C		
				Cycrin	C		

Date:09/22/99ISR Number: 3354173-2Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia		Wellbutrin Sr	PS		ORAL
150 BID ORAL		Cardiomyopathy Sudden Death		Tamoxifen	SS		

Date:09/22/99ISR Number: 3354177-XReport Type:Direct  
Age:54 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction		Welbutrin Sr 100bid	PS		ORAL
150 BID ORAL		Ventricle Rupture		Viagra -Prn Last Dose 2-4d Prior To Event	SS		ORAL

ORAL

Date:09/22/99ISR Number: 3355502-6Report Type:Direct  
Age:26 YR Gender:Male I/FU:I

Company Report #

Outcome	PT
Other	Asthenia Creatine Phosphokinase



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Decreased  
Myalgia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Q AM X	3 DAYS THE		Zyban Glaxo Wellcome	PS	Glaxo Wellcome	
150MG BID						

Date:09/23/99ISR Number: 3355813-4Report Type:Direct Company Report #  
Age:26 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia		Zyban Glaxo Wellcome	PS	Glaxo Wellcome	
150MG Q AM X	3 DAYS THEN	Creatine Phosphokinase					
150 MG BID		Decreased					
		Myalgia					

Date:09/23/99ISR Number: 3356835-XReport Type:Expedited (15-DaCompany Report #A0096492A  
Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization - Initial or Prolonged	150 MG/ TWICE	Cardiac Disorder Dysphagia Dyspnoea	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
Disability PER DAY/ ORAL Required		Embolism					
Intervention to Prevent Permanent Impairment/Damage		Eye Disorder Oedema Peripheral Pulmonary Embolism Swelling Thrombosis		Paracetamol	C		

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Psychotic Disorder Suicidal Ideation	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloridie)	PS		

Date:09/24/99ISR Number: 3356978-0Report Type:Direct

Company Report #

Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Influenza Like Illness Syncope		Zyban	PS		

Date:09/27/99ISR Number: 3357338-9Report Type:Direct

Company Report #

Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100MG PO UP TO TID		Burning Sensation Glossodynia	Health Professional	Wellbutrin 100mg Glaxo/Wellcome	PS	Glaxo/Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/27/99ISR Number: 3358988-6Report Type:Expedited (15-DaCompany Report #A0078462A

Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG / PER DAY / ORAL	Drug Interaction Fear Feeling Abnormal Hallucination Paraesthesia	Consumer	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
			Pain Medication (Formulation Unknown) (Pain Medication) Muscle Relaxant (Formulation Unknown) (Muscle Relaxant) Ventolin (Formulation Unknown) (Albuterol Sulfate) Lithium Salt Insulin	SS  C  C C C		

Date:09/27/99ISR Number: 3361311-4Report Type:Periodic

Company Report #A001-002-003251

Age:85 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - SEE IMAGE Initial or Prolonged	Cerebrovascular Accident Hypertension	Consumer	Aricept (Donepezil)	PS		ORAL
		Health Professional	Wellbutrin (Bupropion Hcl) (Amfebutamone Hydrochloride) Aspirin Atenolol Timoptic Digoxin Diltiazem	SS  C C C C C		

Date:09/28/99ISR Number: 3358827-3Report Type:Direct  
Age:65 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG PO Q Initial or Prolonged DAY		Depressed Level Of Consciousness		Bupropion Sr	PS		ORAL
		Hypertension Hypertensive Encephalopathy Speech Disorder		Solumedrol Theodur	C C		

Date:09/28/99ISR Number: 3366643-1Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #USA009619

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Consumer	Meridia	PS		ORAL
10 MG OD		Nausea		Wellbutrin	SS		ORAL
100 MG BID		Vision Blurred		Toprol Xl	SS		ORAL
100 MG OD				Ogen	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/29/99ISR Number: 3359705-6Report Type:Direct  
Age:31 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
4	MON						
		Angioneurotic Oedema	Health	Wellbutrin	PS		
		Urticaria	Professional				

Date:09/29/99ISR Number: 3361004-3Report Type:Expedited (15-DaCompany Report #A0096492A  
Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening							
Hospitalization -							
Initial or Prolonged							
150 MG /							
Disability							
TWICE PER DAY							
Required							
/ ORAL							
Intervention to							
Prevent Permanent							
Impairment/Damage							
		Cardiac Disorder	Health	Zyban Tablet - Zyban			
		Dysphagia	Professional	(Bupropion			
		Dyspnoea		Hydrochloride)	PS		ORAL
		Embolism					
		Eye Disorder					
		Oedema Peripheral		Paracetamol	C		
		Pulmonary Embolism					
		Swelling					

Date:09/30/99ISR Number: 3360922-XReport Type:Direct  
Age:55 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
ONE BID							
		Dysphagia	Health	Zyban 150mg	PS		
		Face Oedema	Professional	Desowen Lotion	C		
		Oedema Peripheral					
		Periorbital Oedema					
		Scrotal Swelling					
		Urticaria					

Date:10/01/99ISR Number: 3361364-3Report Type:Direct  
Age:45 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG PO QD ; Initial or Prolonged 150MG PO BID		Chest Pain		Bupropion Sr 150mg	PS		ORAL
		Post Procedural					
		Complication		Imdur	C		
		Wound Infection		Lopressor	C		
		Wound Secretion		Lipitor	C		
				Bupropion	C		
				Eco Asa	C		

Date:10/01/99ISR Number: 3363020-4Report Type:Expedited (15-DaCompany Report #A0072356A  
Age:45 YR Gender:Female I/FU:I

Outcome	PT
Other	Akathisia
	Clonic Convulsion
	Diaphragmatic Paralysis
	Diarrhoea
	Dyskinesia
	Dyspnoea
	Dystonia
	Fatigue
	Nervous System Disorder
	Respiratory Alkalosis
	Tardive Dyskinesia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
100 MG / THREE TIMES PER DAY; ORAL		Health Professional Company Representative	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL

Triamcinolone Acetonide	C
Melatonin	C

Date:10/01/99ISR Number: 3363022-8Report Type:Expedited (15-DaCompany Report #A0102044A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Complications Of Maternal Exposure To Therapeutic Drugs	Foreign Study Health Professional Company Representative	Zyban Tablet -Zyban (Bupropion Hydrochloride)	PS		ORAL

150 MG / UNK  
/ ORAL

Date:10/01/99ISR Number: 3363025-3Report Type:Expedited (15-DaCompany Report #A0101960A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged		Cardiac Arrest	Foreign Health Professional	Zyban Tablet -Zyban (Bupropion Hydrochloride)	PS		ORAL

150 MG / PER  
Required  
DAY / ORAL  
Intervention to  
Prevent Permanent

Metoprolol	C
Glibenclamide	C

Impairment/Damage

Metformin

C

Aspirin

C

Date:10/04/99ISR Number: 3364083-2Report Type:Expedited (15-DaCompany Report #A0095928A

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Blood Creatine Phosphokinase Increased Hyperpyrexia Leukopenia	Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)			
ORAL		Neuroleptic Malignant Syndrome Sepsis Serotonin Syndrome Urinary Tract Infection		Dexamphetamine Sulphate (Formulation Unknown) (Dextroamphetamine	PS		ORAL
					SS		

Date:10/04/99ISR Number: 3364087-XReport Type:Expedited (15-DaCompany Report #A0055937A

Age:55 YR Gender:Female I/FU:I

Outcome	PT
Disability	Autonomic Nervous System Imbalance Blood Glucose Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Triglycerides Increased					
		Bradycardia	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150MG PER DAY		Dizziness					
		Haematuria					
		Headache					
ORAL		Hyperhidrosis					
		Hypertension		Conjugated Estrogens	C		
		Hypoaesthesia		Vitamin	C		
		Hypomania		Asspirin	C		
		Labile Blood Pressure		Melatonin	C		
		Mental Impairment					
		Mitral Valve Incompetence					
		Muscle Twitching					
		Musculoskeletal Stiffness					
		Panic Attack					
		Paraesthesia					
		Renal Disorder					
		Tachycardia					

Date:10/04/99ISR Number: 3364114-XReport Type:Expedited (15-DaCompany Report #A0099553A  
 Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dizziness	Health Professional Company	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150MG ORAL		Feeling Abnormal	Representative	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	SS		ORAL
ORAL		Tachycardia					

Date:10/04/99ISR Number: 3364118-7Report Type:Expedited (15-DaCompany Report #A0101583A  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged Other 150MG SEE	Grand Mal Convulsion Hypokalaemia Loss Of Consciousness  Overdose  Sinus Tachycardia Vomiting	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS	ORAL
TEXT ORAL					

Date:10/04/99ISR Number: 3364122-9Report Type:Expedited (15-DaCompany Report #A0101990A

Age: Gender:Female I/FU:I

Outcome Dose Hospitalization - Initial or Prolonged  150MG TWICE  PER DAY ORAL	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dry Mouth Oesophageal Disorder Sensation Of Foreign Body  Speech Disorder  Stomatitis Throat Irritation Tongue Oedema	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/05/99ISR Number: 3364637-3Report Type:Expedited (15-DaCompany Report #A0101636A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Arthralgia Chills Difficulty In Walking	Literature Health Professional	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
300 MG/DAY/ORAL	3 WK	Fatigue Leukocytosis Movement Disorder Myalgia Oedema Peripheral Pyrexia Serum Sickness Tenderness Urticaria					

Date:10/05/99ISR Number: 3364748-2Report Type:Expedited (15-DaCompany Report #A0102079A  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Hypothyroidism Oedema Peripheral Tinnitus	Foreign Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE PER DAY/ORAL		Weight Increased					

Date:10/05/99ISR Number: 3364980-8Report Type:Expedited (15-DaCompany Report #A0102184A  
Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cardiac Failure Clonic Convulsion Decreased Appetite Depression	Consumer	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG /							

TWICE PER DAY Grand Mal Convulsion

Headache

/ ORAL

Respiratory Failure  
Suicidal Ideation  
Weight Decreased

Fluoxetine  
Hydrochloride C  
Ferrous Sulfate C  
Demulen C

Date:10/06/99ISR Number: 3366190-7Report Type:Expedited (15-DaCompany Report #A0073385A

Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthralgia Delirium Dermatitis Malaise	Literature Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Phydrochloride)	PS		ORAL
150MG ORAL		Myalgia Pyrexia Serum Sickness					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/06/99ISR Number: 3366218-4Report Type:Expedited (15-DaCompany Report #A0102058A  
Age:65 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150MG PER DAY ORAL	Confusional State Depressed Level Of Consciousness Hypertension Hypertensive Encephalopathy Speech Disorder Vision Blurred	Study Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
			Me-Prednisolone Na Succ Theophylline Diltiazem Lisionpril Salbutamol Sulphate Ipratropium Bromide Procainamide Hcl Aspirin Isosorbide Dinitrate Metoprolol Tartrate Vitamin E Clopidogrel Bisulphate	C C C C C C C C C C C C		

Date:10/06/99ISR Number: 3366220-2Report Type:Expedited (15-DaCompany Report #A0102413A  
Age:64 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG TWICE PER DAY ORAL	Cerebrovascular Accident Transient Ischaemic Attack	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:10/06/99ISR Number: 3369484-4Report Type:Periodic Company Report #99USA10317  
Age:22 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Death	Overdose	Health Professional Other	Lopressor Tablet (Metoprolol Tartrate)	PS	ORAL
DAILY, ORAL			Acetaminophen Tablet (Paracetamol)	SS	ORAL
ORAL			Fluoxetine Capsule 20 Mg (Fluoxetine)	SS	ORAL
20 MG, DAILY, ORAL			Wellbutrin Slow Release Tablet 100 Mg (Amfebutamone Hydrochloride)	SS	ORAL
200 MG, DAILY, ORAL					

Date:10/07/99ISR Number: 3366394-3Report Type:Direct  
Age:45 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG PO BID Initial or Prolonged	Chest Pain Post Procedural Complication Wound Infection Wound Secretion		Bupropion Sr 150mg Imdur Lopressor Lipitor Eloasa	PS C C C C		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/07/99ISR Number: 3366399-2Report Type:Direct  
Age:65 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 PO Q DAY		Hypertensive		Bupropion Sr	PS		ORAL
Initial or Prolonged		Encephalopathy		Solumedrol Theodur	C C		

Date:10/07/99ISR Number: 3366423-7Report Type:Direct  
Age:29 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 1 BID ORAL	10 DAY	Clonic Convulsion	Health	Wellbutrin Sr 150 Mg	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage		Confusional State Convulsion Fall Head Injury Loss Of Consciousness Syncope	Professional				

Date:10/08/99ISR Number: 3368492-7Report Type:Expedited (15-DaCompany Report #A0073385A  
Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abnormal Behaviour Agitation Arthralgia Chest Pain	Literature Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER DAY / ORAL		Confusional State Delirium Disorientation Drug Hypersensitivity Malaise Myalgia Pyrexia Rash Morbilliform		Thiothixene Lithium Salt Venlafaxine Hydrochloride Thyroxine Sodium Haloperidol Lorazepam	C C C C C C		

Red Blood Cell  
Sedimentation Rate  
Increased  
Restlessness  
Serum Sickness

Trazodone C  
Olanzapine C

Date:10/08/99ISR Number: 3369009-3Report Type:Expedited (15-DaCompany Report #PRIUSA1999006684  
Age:35 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Self-Injury	Literature Health	Haldol (Unspecified) (Haloperidol)	PS		ORAL
ORAL			Professional	Aspirin (Acetylsalicylic Acid)	SS		ORAL
ORAL				Bupropion (Amfebutamone)	SS		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/08/99ISR Number: 3369040-8Report Type:Expedited (15-DaCompany Report #A0102079A  
Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness	Foreign	Zyban Tablet-Zyban			
		Face Oedema	Health	(Bupropion			
		Hypothyroidism	Professional	Hydrochloride)	PS		ORAL
150 MG/TWICE							
		Oedema Peripheral					
PER DAY/ ORAL		Tinnitus					
		Weight Increased					

Date:10/08/99ISR Number: 3369041-XReport Type:Expedited (15-DaCompany Report #A0102535A  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Abdominal Pain Lower	Foreign	Zyban Tablet-Zyban			
Initial or Prolonged		Aphonia	Consumer	(Bupropion			
		Decreased Appetite		Hydrochloride)	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL							

Date:10/08/99ISR Number: 3369051-2Report Type:Expedited (15-DaCompany Report #A0102360A  
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anuria	Consumer	Wellbutrin			
Initial or Prolonged		Back Pain		(Formulation			
		Bladder Disorder		Unknown) (Bupropion			
		Oedema		Hydrochloride)	PS		ORAL
150 MG/ORAL							
		Renal Disorder		Fluoxetine			
		Renal Hypertrophy		Hydrochloride			
				(Formulation			
				Unknown) (Fluoxetine			
				Hydrochloride)	SS		

Date:10/09/99ISR Number: 3367993-5Report Type:Direct  
Age:49 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Facial Palsy		Zyban 150mg	PS		
BID							

Date:10/12/99ISR Number: 3371405-5Report Type:Expedited (15-DaCompany Report #9942002  
Age:83 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Activated Partial	Health	Zoloft Tablets	PS		ORAL
Required		Thromboplastin Time	Professional				
100.00 MG		Prolonged					
Intervention to		Anxiety		Bupropion	SS		ORAL
TOTAL;DAILY;O		Drug Ineffective		Nitropatch	C		
Prevent Permanent		Drug Interaction		Nitrex	C		
RAL		Feeling Abnormal		Levothyroxine	C		
Impairment/Damage		Haemorrhage		Lansoprazole	C		
ORAL		Suicidal Ideation		Alprazolam	C		
				Coumadin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/13/99ISR Number: 3371607-8Report Type:Expedited (15-DaCompany Report #A0102566A

Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Aggression Coma Grand Mal Convulsion Intentional Misuse	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/SEE TEXT/ORAL		Lethargy Suicide Attempt		Gabapentin (Formulation Unknown) (Gabapentin)	SS		ORAL
SEE TEXT/ORAL							

Date:10/13/99ISR Number: 3371770-9Report Type:Expedited (15-DaCompany Report #A0097643A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death ORAL		Myocardial Infarction	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
				Enalapril Maleate Nicotine	C C		

Date:10/14/99ISR Number: 3371013-6Report Type:Direct

Company Report #

Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG QD PO 3 DAY Initial or Prolonged 150 MG BID PO		Chest Pain	Health Professional	Bupropion Sr Bupropion Sr	PS SS		ORAL ORAL
				Colace Ec Asa Lipitor Plavix	C C C C		

Lopressor C  
Ntg Sl C

Date:10/15/99ISR Number: 3373503-9Report Type:Expedited (15-DaCompany Report #A0102803A  
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Foreign	Zyban Tablet - Zyban			
		Drug Hypersensitivity	Health	(Bupropion			
		Dyspnoea	Professional	Hydrochloride)	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL		Laryngeal Oedema					

Date:10/15/99ISR Number: 3373873-1Report Type:Expedited (15-DaCompany Report #A0100352A  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Jaundice	Health	Wellbutrin Tablet			
Initial or Prolonged		Liver Function Test	Professional	(Bupropion			
		Abnormal		Hydrochloride)	PS		ORAL
75 MG							
SEE TEXT							
ORAL							
				Sertraline Tablet			
				(Sertraline)	SS		ORAL
200 MG							
IN THE							

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Freedom Of Information (FOI) Report

MORNING

ORAL

Date:10/18/99ISR Number: 3375713-3Report Type:Expedited (15-DaCompany Report #1999UW03215  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health	Seroquel "Zeneca"	PS	Zeneca	
		Drug Toxicity	Professional	Bupropion	SS		
		Intentional Misuse		Olanzapine	SS		
				Carbamazepine	SS		
				Pramoxine	SS		

Date:10/19/99ISR Number: 3375675-9Report Type:Expedited (15-DaCompany Report #A0086318A  
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dermatitis	Health	Zyban Tablet- Zyban			
		Drug Hypersensitivity	Professional	(Bupropion			
		Dysphagia		Hydrochloride)	PS		ORAL
150 MG/ TWICE		Eyelid Oedema					
PER DAY/ ORAL		Face Oedema		Zyban Tablet - Zyban			
		Pharyngeal Oedema		(Bupropion			
				Hydrochloride)	SS		ORAL
150 MG/							
SINGLE DOSE/							
ORAL				Lansoprazole	C		
				Lorcet	C		

Date:10/19/99ISR Number: 3375678-4Report Type:Expedited (15-DaCompany Report #A0102868A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/	ORAL			Trovafloxacin Mesylate	C		
				Morphine	C		

Date:10/19/99ISR Number: 3375747-9Report Type:Expedited (15-DaCompany Report #A0101186A  
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Disability		Pain Rash Generalised Skin Exfoliation	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /							
TWICE PER DAY							
/	ORAL						

Date:10/19/99ISR Number: 3375749-2Report Type:Expedited (15-DaCompany Report #A0102884A  
Age:45 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Blister Blood Glucose Increased

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dyspnoea Eyelid Oedema Nail Discolouration	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
TWICE PER DAY		Rash Pruritic Skin Discolouration Stevens-Johnson Syndrome					
/ ORAL		Weight Increased					

Date:10/19/99ISR Number: 3375982-XReport Type:Expedited (15-DaCompany Report #A0095816A  
Age:6 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Candidiasis Dysphemia Kidney Infection	Consumer	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
37.5 MG /		Migraine					
ALTERNATE		Pharyngitis Streptococcal					
DAYS / ORAL		Tongue Spasm					

Date:10/19/99ISR Number: 3375984-3Report Type:Expedited (15-DaCompany Report #A0102783A  
Age:13 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Bronchospasm Dermatitis Dyspnoea Oedema Pyrexia Stevens-Johnson Syndrome Urticaria Vomiting	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		

Date:10/19/99ISR Number: 3375986-7Report Type:Expedited (15-DaCompany Report #A0102623A  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrchloride)	PS		ORAL
150 MG /TWICE PER DAY/ ORAL							
40 MG/ ORAL				Fluoxetine Hydrochloride Capsule (Fluoxetine Hydrochloride)	SS		ORAL

Date:10/19/99ISR Number: 3375992-2Report Type:Expedited (15-DaCompany Report #A0080880A  
Age: Gender:Female I/FU:I

Outcome	PT
Disability	Discomfort Dizziness Drug Hypersensitivity Dry Mouth Feeling Abnormal Feeling Jittery

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Illogical Thinking Insomnia Muscle Rigidity	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE PER DAY / ORAL		Muscle Spasms Myalgia Nervousness  Panic Disorder Without Agoraphobia  Tetany Vision Blurred Visual Disturbance		Magnesium Salt	C		

Date:10/19/99ISR Number: 3375994-6Report Type:Expedited (15-DaCompany Report #A0080063A  
Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Discomfort Dizziness Drug Hypersensitivity Dry Mouth Fatigue	Health Professional	Bupropion Hydrochloride Tablet- Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL		Feeling Jittery  Illogical Thinking Insomnia Muscle Rigidity Muscle Spasms Myalgia Nervousness Panic Reaction Tetany Vision Blurred Visual Disturbance		Multivitamin	C		

Date:10/19/99ISR Number: 3376106-5Report Type:Expedited (15-DaCompany Report #A0103180A  
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cardiac Arrest Dyspnoea Tongue Oedema	Consumer	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		
100 MG / THREE TIMES PER DAY		Tracheal Oedema Tracheostomy		Blood Pressure Medication	C		

Date:10/19/99ISR Number: 3376114-4Report Type:Expedited (15-DaCompany Report #A0095964A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dysarthria Feeling Hot Headache	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY / ORAL		Monoparesis Nausea		Ethanol Liquid (Alcohol)	SS		ORAL
ORAL		Respiratory Arrest Syncope		Fluoxetine Hydrochloride	C		
		Tremor Urinary Incontinence					

Freedom Of Information (FOI) Report

Aspirin C

Date:10/20/99ISR Number: 3376492-6Report Type:Expedited (15-DaCompany Report #A0103160A  
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Lung Disorder Pneumonia	Foreign Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG							
TWICE PER DAY							
ORAL							

Nifedipine C

Date:10/20/99ISR Number: 3376495-1Report Type:Expedited (15-DaCompany Report #A0097683A  
 Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other Required		Confusional State Delirium Muscle Disorder Muscle Rigidity	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG							
Intervention to TWICE PER DAY Prevent Permanent ORAL Impairment/Damage		Neuroleptic Malignant Syndrome Pyrexia		Semisodium Valproate Librium Buspirone Hydrochloride Diltiazem Hydrochloride Conjugated Estrogens Thyroxine Sodium Hydrochlorothiazide Omeprazole Lisinopril	C C C C C C C C C C		

Date:10/22/99ISR Number: 3377668-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #990083.01

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Consumer	Furosemide Tablets			
20 MG QD,		Muscle Spasms		20 Mg Mylan	PS	Mylan	ORAL
ORAL		Paraesthesia					
				Wellbutrin 75 Mg			
75 MG QD,				Glaxo Wellcome	SS	Glaxo Wellcome	ORAL
ORAL	1	MON					

Date:10/22/99ISR Number: 3378767-3Report Type:Direct  
Age:49 YR Gender:Female I/FU:I

Company Report #

Outcome  
Life-Threatening  
Hospitalization -  
Initial or Prolonged  
Required  
Intervention to  
Prevent Permanent

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Freedom Of Information (FOI) Report

Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG BID		Convulsion	Health	Bupropion	PS		
APPROXIMATELY		Muscle Rigidity	Professional				
2 DAYS	2 DAY	Nausea					
		Pyrexia					
		Sedation					

Date:10/25/99ISR Number: 3381320-9Report Type:Expedited (15-DaCompany Report #A0093402A  
 Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Breast Cancer Recurrent	Health	Wellbutrin			
Other		Metastases To Liver	Professional	Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER				Tamoxifen	C		
DAY / ORAL				Paroxetine Hydrochloride	C		

Date:10/25/99ISR Number: 3381565-8Report Type:Expedited (15-DaCompany Report #A0101636A  
 Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Literature	Wellbutrin Tablet			
Other		Chills	Health	(Bupropion			
300 MG/DAY /		Difficulty In Walking	Professional	Hydrochloride)	PS		ORAL
ORAL		Fatigue					
		Leukocytosis					
		Myalgia					
		Oedema Peripheral					

Pyrexia  
 Serum Sickness  
 Tenderness  
 Urticaria

Date:10/25/99ISR Number: 3381679-2Report Type:Expedited (15-DaCompany Report #A0086318A  
 Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged		Dermatitis	Health Professional	Bupropion Hydrochloride Tablet			
Other		Dysphagia		-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / Required		Face Oedema					
TWICE PER DAY Intervention to / ORAL		Pharyngeal Oedema					
Prevent Permanent Impairment/Damage				Bupropion Hydrochloride Tablet			
150 MG / SINGLE DOSE / ORAL				-Zyban (Bupropion Hydrochloride)	SS		ORAL
				Lansoprazole	C		
				Lorcet	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/25/99ISR Number: 3382299-6Report Type:Expedited (15-DaCompany Report #A0068941A  
Age:54 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/ ORAL	Conversion Disorder Feeling Abnormal Hallucination Tremor	Consumer	Zyban Tablet- Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:10/25/99ISR Number: 3382918-4Report Type:Expedited (15-DaCompany Report #A0077179A  
Age:61 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Atrial Fibrillation Influenza Lung Neoplasm Malignant Pyrexia	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:10/26/99ISR Number: 3381635-4Report Type:Direct Company Report #  
Age:23 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG BID Required P.O. Intervention to Prevent Permanent Impairment/Damage	Abdominal Pain Arthralgia Bronchospasm Forced Expiratory Volume Abnormal Hypersensitivity Influenza Like Illness Malaise Pruritus Urticaria		Zyban (Bupropion)	PS		ORAL

Date:10/27/99ISR Number: 3382458-2Report Type:Direct Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Balance Disorder	Health	Wellbutrin 75mg	PS		ORAL
75MG PO BID		Feeling Jittery	Professional	Cardizem Cd	C		
				Erythromycin	C		
				Hydroxyzine	C		
				Zocor	C		

Date:10/27/99ISR Number: 3383444-9Report Type:Expedited (15-DaCompany Report #A0103571A  
Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness	Health	Wellbutrin	PS		ORAL
100 MG FOUR		Dizziness	Professional				
TIMES PER DAY		Nausea					
ORAL							

Date:10/28/99ISR Number: 3384829-7Report Type:Expedited (15-DaCompany Report #A0103729A  
Age:31 YR Gender:Female I/FU:I

Outcome	PT
Other	Convulsion
	Tachycardia



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG / UNK		Foreign Consumer Other	Ceftin (Cefuroxime Axetil) Zyban	PS SS		ORAL
/ ORAL			Ibuprofen Amoxicillin Trihydrate	SS SS		

Date:10/29/99ISR Number: 3384931-XReport Type:Direct  
Age:20 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG ORAL Initial or Prolonged		Convulsion		Wellbutrin-Glaxo Luvox	PS C	Glaxo	ORAL

Date:10/29/99ISR Number: 3386131-6Report Type:Periodic  
Age:21 YR Gender:Female I/FU:I

Company Report #A0102154A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion Overdose	Consumer	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

300 MG/TWICE

PER DAY/ ORAL

Date:10/29/99ISR Number: 3386132-8Report Type:Periodic  
Age:18 YR Gender:Female I/FU:I

Company Report #A0102376A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Convulsion Health Professional Wellbutrin  
Tablet-Controlled  
Company Release (Bupropion  
Representative Hydrochloride) PS ORAL

Date:10/29/99ISR Number: 3386133-XReport Type:Periodic Company Report #A0087281A  
Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Other		Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:10/29/99ISR Number: 3386135-3Report Type:Periodic Company Report #A0089931A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Cancer Chemotherapy  
 (Formulation  
 Unknown) (Cancer  
 Chemotherapy) SS  
 Zolpidem Tartrate C

Date:10/29/99ISR Number: 3386138-9Report Type:Periodic  
 Age:48 YR Gender:Female I/FU:F

Company Report #A0095478A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Coma Suicide Attempt	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ PER DAY/ ORAL				Citalopram (Formulation Unknown) (Citalopram)	SS		
20 MG/ PER DAY/				Trazodone	C		

Date:10/29/99ISR Number: 3386140-7Report Type:Periodic  
 Age:11 YR Gender:Male I/FU:F

Company Report #A0096029A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ PER DAY/ ORAL	1 YR						

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Head Injury	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Cyclobenzaprine Hcl	C		
				Dilantin	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Syncope	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
250 MG /							
TWICE PER DAY							
/ ORAL							
				Cisapride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/29/99ISR Number: 3386149-3Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #A0096370A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - Initial or Prolonged Other 150 MG / Required TWICE PER DAY Intervention to / ORAL Prevent Permanent Impairment/Damage	Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL			Tramadol Hydrochloride (Formulation Unknown) (Tramadol Hydrochloride)	SS		
			Venlafaxine Hydrochloride	C		

Date:10/29/99ISR Number: 3386152-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0096390A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Other 150 MG / TWICE PER DAY / ORAL	Convulsion	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
			Paroxetine Hydrochloride	C		
			Darvocet-N	C		
			Zolpidem Tartrate	C		

Date:10/29/99ISR Number: 3386155-9Report Type:Periodic Company Report #A0096971A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Amnesia Convulsion Migraine	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER DAY / ORAL	4	YR		Sertraline Hydrochloride	C		

Date:10/29/99ISR Number: 3386159-6Report Type:Periodic Company Report #A0097592A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion Drug Ineffective Laceration	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / SEE TEXT / ORAL	4	MON					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/29/99ISR Number: 3386161-4Report Type:Periodic  
Age:29 YR Gender:Male I/FU:I

Company Report #A0097691A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Depression Fatigue	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
200 MG /							
TWICE PER DAY							
/ ORAL							

Date:10/29/99ISR Number: 3386164-XReport Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #A0098230A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis Pruritus Rash Papular Stevens-Johnson Syndrome Urticaria	Health Professional	Wellbutrin Tablet-Controlled Release 150 Mg (Bupropion Hydrochloride)	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:10/29/99ISR Number: 3386169-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0098258A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged Other		Convulsion	Health Professional Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		

Date:10/29/99ISR Number: 3386172-9Report Type:Periodic Company Report #A0098464A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Health	Wellbutin Tablet			
Other		Grand Mal Convulsion	Professional	(Bupropion Hydrochloride)	PS		ORAL
400 MG / ORAL				Citalopram Hydrobromide	C		

Date:10/29/99ISR Number: 3386180-8Report Type:Periodic Company Report #A0098531A  
Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Health	Wellbutrin			
Other		Dysgeusia	Professional	Tablet-Controlled			
		Eye Rolling	Company	Release (Bupropion			
		Movement Disorder	Representative	Hydrochloride)	PS		ORAL
150 MG /		Muscle Twitching					
TWICE PER DAY		Posturing					
/ ORAL	10 DAY	Sedation		Lorazepam	C		
		Speech Disorder		Semisodium Valproate	C		
		Vision Blurred		Diphenhydramine Hcl	C		
				Cyclobenzaprine Hcl	C		
				Excedrin	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/29/99ISR Number: 3386185-7Report Type:Periodic  
Age:19 YR Gender:Female I/FU:I

Company Report #A0099382A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER DAY / ORAL				Azithromycin Doxycycline Oral Contraceptive	C C C		

Date:10/29/99ISR Number: 3386191-2Report Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #A0099922A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Face Oedema Oedema Peripheral Periorbital Oedema	Consumer	Wellbutrin Tablet-Controlled Release	PS		ORAL
150 MG / TWICE PER DAY / ORAL		Pruritus Skin Disorder		Multivitamin	C		
		Swelling Urticaria					

Date:10/29/99ISR Number: 3386197-3Report Type:Periodic  
Age:34 YR Gender:Male I/FU:I

Company Report #A0100210A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Anaphylactic Reaction Angioneurotic Oedema Arthralgia Dermatitis	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Bydrochloride)	PS		ORAL
150 MG /							

Dyspnoea  
Pain  
TWICE PER DAY  
/ ORAL

Date:10/29/99ISR Number: 3386201-2Report Type:Periodic Company Report #A0100421A  
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Convulsion Disorientation Sedation	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride) Semisodium Valproate	PS C		

Date:10/29/99ISR Number: 3386206-1Report Type:Periodic Company Report #A0100585A  
Age:23 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

150 MG /

TWICE PER DAY  
/ ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/29/99ISR Number: 3386210-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0101446A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL				Citalopram Hydrobromide	C		

Date:10/29/99ISR Number: 3386213-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0101495A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY / ORAL	10 DAY						

Date:10/29/99ISR Number: 3386215-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0101936A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Control Release (Bupropion Hydrochloride)	PS		

Date:11/01/99ISR Number: 3386411-4Report Type:Direct  
Age:43 YR Gender:Male I/FU:I

Company Report #

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 PO QD ->		Convulsion	Health	Wellbutrin 150 Sr	PS		ORAL
BID	30	DAY	Extrapyramidal Disorder	Professional				
10MG	30	DAY	Muscle Twitching		Zyprexa 10mg	SS		
			Neuroleptic Malignant Syndrome		Clonazepam	C		
					Prolixin	C		
					Cogentin	C		
					Celexa	C		

Date:11/01/99ISR Number: 3386412-6Report Type:Direct Company Report #  
Age:30 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150AM 75PM	30 DAY	Medication Error	Health	Wellbutrin Sr 150mg	PS		
				Professional	Tramadol 50mg Po	SS		ORAL
50 Q ? PO		2 DAY			Zyprexa	C		

Date:11/01/99ISR Number: 3387206-8Report Type:Expedited (15-DaCompany Report #A0103746A  
Age:24 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Arthralgia Chills

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE	PER DAY/ ORAL 16 DAY	Difficulty In Walking Dyspepsia Face Oedema Lymphadenitis Myalgia Oedema Oesophageal Disorder Pruritus Pyrexia Rash Erythematous Serum Sickness Synovitis Urticaria White Blood Cell Count Increased	Foreign Literature Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:11/01/99ISR Number: 3387287-1Report Type:Expedited (15-DaCompany Report #A0103905A  
Age: Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG / ORAL	7 WK	Completed Suicide	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:11/01/99ISR Number: 3387291-3Report Type:Expedited (15-DaCompany Report #A0103615A  
Age:49 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged	150 MG /TWICE	PER DAY/ORAL	Abnormal Dreams Angioneurotic Oedema Dermatitis Dyspnoea Eyelid Oedema Face Oedema Hypertension	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)  Nabumetone	PS  C		ORAL

Nausea  
 Oral Intake Reduced  
 Pain  
 Pruritus  
 Psychotic Disorder  
 Urticaria

Date:11/01/99ISR Number: 3387292-5Report Type:Expedited (15-DaCompany Report #A0102535A

Age:60 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Disability 150 MG/TWICE PER DAY/ORAL	Abdominal Pain Lower Aphonia Blood Sodium Decreased Decreased Appetite	Foreign Health Professional	Zyban Tablet- Zyban (Bupropion Hydrochloride)	PS		ORAL
	Influenza Pyrexia Vomiting		Metformin Aspirin Diltiazem Hydrochloride Cerivastatin Ramipril	C C C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/99ISR Number: 3388598-6Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG QD X3 Initial or Prolonged THEN BID	Erythema Multiforme  Lymphadenopathy  Stevens-Johnson Syndrome		Zyban 150mg	PS		

Date:11/03/99ISR Number: 3389230-8Report Type:Expedited (15-DaCompany Report #A0102044A  
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Congenital Anomaly  150 MG /  TWICE PER DAY  / ORAL	Complications Of Maternal Exposure To Therapeutic Drugs  Foetal Disorder  Gastroschisis	Foreign Study Health  Professional  Company  Representative	Zyban Tablet- Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:11/03/99ISR Number: 3389236-9Report Type:Expedited (15-DaCompany Report #A0104285A  
Age:18 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged  ORAL	Burning Sensation Decreased Activity Dehydration  Muscle Spasms Rash Generalised Rash Pruritic	Foreign Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:11/04/99ISR Number: 3388895-4Report Type:Expedited (15-DaCompany Report #A0104362A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin			
		Drug Interaction	Professional	Tablet-Controlled			
		Urinary Tract Infection	Company	Release (Bupropion			
			Representative	Hydrochloride)	PS		ORAL

SEE TEXT/

ORAL

Bactrim (Formulation Unknown)			
(Sulfamethoxazole/Trimetho)	SS		
Fluoxetine			
Hydrochloride	C		

Date:11/04/99ISR Number: 3388898-XReport Type:Expedited (15-DaCompany Report #A0104365A  
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dizziness	Foreign	Zyban Tablet - Zyban			
Initial or Prolonged		Headache	Consumer	(Bupropion			
		Tinnitus		Hydrochloride)	PS		ORAL

150 MG/ TWICE

PER DAY/ ORAL

Atenolol	C		
Pravastatin Sodium	C		
Nitroglycerin	C		
Salbutamol Sulphate	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lorazepam

C

Date:11/05/99ISR Number: 3390175-8Report Type:Expedited (15-DaCompany Report #A0098418A

Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage	150 MG/TWICE	Abnormal Behaviour Belligerence Convulsion Fall	Health Professional Company Representative	Wellbutrin Tablet -Controlled Release (Bupropion Hydrochlride)	PS		ORAL
PER DAY/ORAL		Haemorrhage					
ORAL		Head Injury Upper Respiratory Tract Infection		Pseudoephedrine (Formulation Unknown) (Pseudoephedrine)	SS		ORAL
ORAL				Dextromethorphan (Formulation Unknown) (Dextromethorphan)	SS		ORAL

Date:11/05/99ISR Number: 3390176-XReport Type:Expedited (15-DaCompany Report #A0102803A

Age:25 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other	150 MG/TWICE	Arthralgia Dermatitis Drug Hypersensitivity	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
PER DAY/ ORAL		Dyspnoea					
		Laryngeal Oedema					

Date:11/08/99ISR Number: 3390236-3Report Type:Direct

Age:44 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 6GM PO X 1		Clonic Convulsion	Health	Bupropion	PS		ORAL
Initial or Prolonged DOSE Required		Coma	Professional				
Intervention to Prevent Permanent Impairment/Damage		Overdose Tachycardia		Citalopram	C		

Date:11/08/99ISR Number: 3390751-2Report Type:Expedited (15-DaCompany Report #9942002  
Age:83 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG TOTAL		Activated Partial	Health	Zoloft Tablets	PS		ORAL
Initial or Prolonged DAILY ORAL Required		Thromboplastin Time	Professional				
ORAL Intervention to Prevent Permanent Impairment/Damage		Prolonged		Bupropion	SS		ORAL
		Anxiety		Nitropatch	C		
		Bleeding Time Prolonged		Nitrex	C		
		Depressed Mood		Levothyroxine	C		
		Drug Ineffective		Lansoprazole	C		
		Ear Haemorrhage		Alprazolam	C		
		Fatigue		Coumadin	C		
		Feeling Abnormal		Vitamin B12	C		
		Sleep Disorder		Vitamin E	C		
		Suicidal Ideation		Vitamin C	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/08/99ISR Number: 3391243-7Report Type:Expedited (15-DaCompany Report #A0104721A  
Age:77 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150MG ORAL	Hyperkalaemia	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
25MG TWICE PER DAY			Spiroonolactone (Formulation Unknown) (Spironolactone)	SS		
			Frusemide	C		

Date:11/08/99ISR Number: 3391335-2Report Type:Expedited (15-DaCompany Report #A0098556A  
Age:35 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to Prevent Permanent Impairment/Damage 150MG ORAL	Dysphagia Dyspnoea Oesophagitis	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
			Omeprazole	C		
			Ranitidine Hydrochloride	C		

Date:11/09/99ISR Number: 3391436-9Report Type:Direct  
Age:14 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150MG BID PO Required Intervention to Prevent Permanent	Coma Confusional State Convulsion Muscle Rigidity Paranoia		Welbutrin (Bupropion)	PS		ORAL
			Antihistamin	C		

Impairment/Damage

Sensory Loss  
Speech Disorder  
Vomiting

Date:11/10/99ISR Number: 3391808-2Report Type:Direct  
Age:36 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin Sr 150mg	PS		
150MG 2X		Urticaria					
DAILY							

Date:11/12/99ISR Number: 3396181-1Report Type:Expedited (15-DaCompany Report #A0104815A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Atrioventricular Block	Health	Wellbutrin			
Other		Complete	Professional	Tablet-Controlled			
		Dyspnoea		Release (Bupropion	PS		ORAL
				Hydrochloride)			
1 TABLET /							
PER DAY /							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/12/99ISR Number: 3396185-9Report Type:Expedited (15-DaCompany Report #B0072373A  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Agitation Alopecia Dermatitis Atopic Dermatitis Exfoliative Dry Mouth Extrasystoles Face Oedema Flushing Gastrooesophageal Reflux Disease Hypertension Muscle Rigidity Muscle Spasms Palpitations Paraesthesia Photosensitivity Reaction Pruritus Tachycardia Tremor Urinary Retention Urticaria	Foreign Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		

Date:11/12/99ISR Number: 3396816-3Report Type:Expedited (15-DaCompany Report #A0098933A  
Age:26 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Life-Threatening Hospitalization - Initial or Prolonged 150 MG/TWICE Disability PER DAY/ ORAL		Ascites Blood Bilirubin Increased Coagulopathy Depression Epididymitis Fatigue Hepatic Cirrhosis Hepatic Enzyme Increased Hepatic Failure Hepatic Fibrosis Hepatic Necrosis	Foreign Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
				Salbutamol Sulphate	C		
				Beclomethasone			
				Dipropion.	C		
				Venlafaxine Hydrochloride	C		
				Fluticasone			

Hepatitis  
International Normalised  
Ratio Increased  
Jaundice  
Nausea  
Petechiae  
Pleural Effusion  
Prothrombin Time  
Prolonged  
Pulmonary Congestion  
Renal Failure  
Splenomegaly

Propionate

C

Date:11/12/99ISR Number: 3396937-5Report Type:Expedited (15-DaCompany Report #A0104511A  
Age:46 YR Gender:Male I/FU:I

Outcome	PT	Report Source
Other	Complex Partial Seizures	Health
	Drug Interaction	Professional

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Company Representative	Product	Role	Manufacturer	Route
150 MG /			Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
TWICE PER DAY						
/ ORAL			Ritonavir (Formulation Unknown) (Ritonavir) Anti-Viral Medication	SS  C		

Date:11/12/99ISR Number: 3397224-1Report Type:Expedited (15-DaCompany Report #A0102868A  
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional Company Representative	Wellbutrin Tablet- Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ ORAL				Trovafloxacin Mesylate Morphine	C C		

Date:11/12/99ISR Number: 3397374-XReport Type:Expedited (15-DaCompany Report #A0103746A  
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthralgia Asthma Chills	Foreign Literature Health	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /							

TWICE PER DAY

Dyspepsia

Professional

/ ORAL 16 DAY

Face Oedema

Lymphadenopathy

Movement Disorder

Myalgia

Odynophagia

Oedema Peripheral

Pain In Extremity

Periorbital Oedema

Pruritus

Pyrexia

Rash Erythematous

Serum Sickness

Synovitis

Urticaria

White Blood Cell Count

Increased

Date:11/15/99ISR Number: 3397214-9Report Type:Expedited (15-DaCompany Report #A0102889A

Age:23 YR Gender:Male I/FU:I

Outcome PT

Other Abdominal Pain

Anaphylactic Reaction

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Arthralgia Bronchospasm Dyspnoea	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /		Malaise Peak Expiratory Flow Rate Decreased					
TWICE PER DAY		Pharyngolaryngeal Pain					
/ ORAL		Pruritus					
		Pulmonary Function Test Abnormal Throat Tightness Urticaria					

Date:11/15/99ISR Number: 3397218-6Report Type:Expedited (15-DaCompany Report #A0105273A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Depressed Level Of Consciousness Renal Failure	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
7 DAY		Rhabdomyolysis		Ethanol (Formulation Unknown) (Alcohol)	SS		

Date:11/15/99ISR Number: 3397912-7Report Type:Expedited (15-DaCompany Report #A0104240A  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Required		Aggression Mania Suicidal Ideation	Study Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
400 MG Intervention to PER DAY Prevent Permanent ORAL							

Impairment/Damage

Paracetamol

C

Date:11/16/99ISR Number: 3397464-1Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropion Sr	PS		
Other		Confusional State		Ranitidine	C		
100MG TID				Naproxene	C		
				Darvocet	C		
				Zolpidem	C		

Date:11/17/99ISR Number: 3399752-1Report Type:Expedited (15-DaCompany Report #A0105253A  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
Hospitalization - Initial or Prolonged		Amnesia Convulsion Loss Of Consciousness	Consumer				
150 MG/ TWICE PER DAY/ ORAL		Respiratory Arrest		Aspirin	C		
				Sertraline Hydrochloride	C		
				Atorvastatin Calcium	C		

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Freedom Of Information (FOI) Report

Cetirizine  
Hydrochloride C

Date:11/18/99ISR Number: 3402656-9Report Type:Expedited (15-DaCompany Report #A0062864A  
Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abortion Spontaneous Pregnancy	Health Professional	Bupropion Hydrochloride Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE PER DAY/ORAL							

Date:11/18/99ISR Number: 3402657-0Report Type:Expedited (15-DaCompany Report #A0062610A  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abortion Spontaneous Pregnancy	Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE PER DAY/ORAL							

Date:11/19/99ISR Number: 3403049-0Report Type:Expedited (15-DaCompany Report #A0105423A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Disability		Hypoaesthesia Multiple Sclerosis	Foreign Health Professional	Bupropion Hydrochloride Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE PER DAY/ ORAL							

Date:11/19/99ISR Number: 3403933-8Report Type:Expedited (15-DaCompany Report #A0104285A

Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required	150 MG / Intervention to TWICE PER DAY Prevent Permanent Impairment/Damage	Burning Sensation Decreased Activity Dehydration  Dermatitis  Muscle Spasms Pruritus Urticaria Vomiting	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:11/22/99ISR Number: 3403652-8Report Type:Direct

Company Report #

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
TITRATED FROM 6/2-6/25 150MG Q AM , NOON & PM 150MG Q HS		Amnesia  Drug Interaction  Loss Of Consciousness  Tonic Convulsion		Wellbutrin Sr    Clozapine 150mg  Effexor Xr	PS    SS  C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/22/99ISR Number: 3408491-XReport Type:Periodic Company Report #S99-USA-01718-01  
 Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
40 MG QD PO		Drug Interaction	Health	Celexa	PS		ORAL
		Pruritus	Professional	Wellbutrin	SS		
		Urticaria		Oral Contraceptives	C		

Date:11/23/99ISR Number: 3409304-2Report Type:Expedited (15-DaCompany Report #A0105636A  
 Age:17 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged Other		Abdominal Pain Blood Creatine Phosphokinase Increased Grand Mal Convulsion	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
SEE TEXT		Hallucination, Visual					
ORAL		Intentional Misuse Mental Impairment Sedation Sinus Tachycardia Suicide Attempt					

Date:11/23/99ISR Number: 3409306-6Report Type:Expedited (15-DaCompany Report #A0105554A  
 Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Abdominal Pain Upper Agranulocytosis Fatigue Influenza	Consumer	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
THREE TIMES		Libido Decreased					
PER DAY		Lung Disorder					
ORAL		Malaise					

Pneumonia  
Pyrexia  
Suicidal Ideation  
Vomiting  
White Blood Cell Count  
Decreased

Date:11/24/99ISR Number: 3406829-0Report Type:Direct  
Age:39 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG SR BID Initial or Prolonged	Bite Convulsion Cyanosis Salivary Hypersecretion Tremor		Wellbutrin Sr 150mg Ortho-Tricyclen	PS C		

Date:11/24/99ISR Number: 3410232-7Report Type:Expedited (15-DaCompany Report #A0102413A  
Age:64 YR Gender:Male I/FU:F

Outcome	PT
Other	Cerebral Infarction Cerebrovascular Accident Chest Pain

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dysarthria Headache Hemiparesis	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
TWICE PER DAY		Hyperhidrosis Transient Ischaemic Attack					
/ ORAL		Vertigo		Sinemet	C		

Date:11/26/99ISR Number: 3408907-9Report Type:Expedited (15-DaCompany Report #A0105636A  
Age:17 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Abdominal Pain Blood Creatine Phosphokinase Increased Grand Mal Convulsion	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
SEE TEXT /		Hallucination, Visual					
ORAL		Intentional Misuse Mental Impairment Sedation Sinus Tachycardia Suicide Attempt					

Date:11/26/99ISR Number: 3408919-5Report Type:Expedited (15-DaCompany Report #A0105877A  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Confusional State Cyanosis Dyspnoea	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
TWICE PER DAY		Eye Rolling					

Movement Disorder

/ ORAL

Sedation

Sumatriptan	
Succinate	C
Propranolol	
Hydrochloride	C
Salbutamol Sulphate	C
Triquilar	C

Date:11/26/99ISR Number: 3408922-5Report Type:Expedited (15-DaCompany Report #A0105879A

Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthritis		Zyban Tablet - Zyban			
		Diarrhoea		(Bupropion			
		Dizziness		Hydrochloride)	PS		ORAL
150 MG /		Erythema Multiforme					
TWICE PER DAY		Leukocytosis					
/ ORAL	15	DAY					
		Nausea					
		Neutrophilia					
		Oedema					
		Pruritus					
		Serum Sickness					
		Tachycardia					
		Tongue Disorder					
		Urticaria					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/99ISR Number: 3411023-3Report Type:Direct  
Age:42 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Face Oedema	Health	Zyban (Glaxo)	PS		ORAL
150MG BID PO		Urticaria	Professional	Synthroid	C		

Date:11/30/99ISR Number: 3411804-6Report Type:Expedited (15-DaCompany Report #A0098738A  
Age:72 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aneurysm	Consumer	Wellbutrin			
Hospitalization -		Arterial Rupture		Tablet-Controlled			
Initial or Prolonged		Atrial Fibrillation		Release (Bupropion			
Other		Dizziness		Hydrochloride)	PS		ORAL
150 MG		Fatigue					
TWICE PER DAY		Headache					
		Hypertension					
ORAL		Insomnia		Verapamil	C		
		Metastases To Liver		Hyzaar	C		
		Nephrolithiasis		Librium	C		
		Palpitations					
		Pneumonia					

Date:11/30/99ISR Number: 3411839-3Report Type:Expedited (15-DaCompany Report #A0104511A  
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Complex Partial Seizures	Health	Wellbutrin			
Other		Drug Interaction	Professional	Tablet-Controlled			
			Company	Release (Bupropion			
			Representative	Hydrochloride)	PS		ORAL
150 MG			Other				
TWICE PER DAY							

ORAL

Ritonavir Solution  
(Ritonavir)

SS

ORAL

ORAL

Anti-Viral  
Medication

C

Date:11/30/99ISR Number: 3411840-XReport Type:Expedited (15-DaCompany Report #A0105888A

Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other		Myocardial Infarction	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS	Zyban	ORAL
150 MG							
PER DAY							
ORAL	2 DAY						

Date:11/30/99ISR Number: 3411841-1Report Type:Expedited (15-DaCompany Report #A0105871A

Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aspiration Overdose	Foreign	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride) Nortriptyline	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Nortriptyline)  
 (Formulation Unkown) SS  
 Cocaine (Cocaine)  
 (Formulation  
 Unknown) SS

Date:11/30/99ISR Number: 3411842-3Report Type:Expedited (15-DaCompany Report #A0105850A  
 Age:17 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Overdose	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Date:11/30/99ISR Number: 3414970-1Report Type:Periodic Company Report #A0089382A  
 Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Discomfort Chest Pain Ecchymosis	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							
Hypersensitivity							
Rash Erythematous Rash Maculo-Papular Urticaria							

Date:11/30/99ISR Number: 3414971-3Report Type:Periodic Company Report #A0082332A  
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Fall Head Injury	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE							

PER DAY/ ORAL	Laceration					
	Loss Of Consciousness			Percocet		C
				Alprazolam		C
				Diamorphine		
				Hydrochloride		C

Date:11/30/99ISR Number: 3414972-5Report Type:Periodic Company Report #A0070364A  
 Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema	Health	Zyban Tablet - Zyban			
		Dermatitis	Professional	(Bupropion			
		Hypersensitivity		Hydrochloride)	PS		ORAL
150 MG/ TWICE		Pruritus					
PER DAY/ ORAL		Swelling					
		Urticaria					

Date:11/30/99ISR Number: 3414973-7Report Type:Periodic Company Report #A0101092A  
 Age: Gender:Female I/FU:I

Outcome	PT
Other	Bladder Disorder
	Convulsion

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Faecal Incontinence Feeling Abnormal Hyperhidrosis					
150 MG/ ORAL		Muscle Spasms Musculoskeletal Disorder Tremor	Health Professional Company  Representative	Zyban Tablet - Zyban (Bupropion Hydrochloride)  Fluoxetine Hydrochloride	PS  C		ORAL

Date:11/30/99ISR Number: 3414974-9Report Type:Periodic Company Report #A0094675A  
Age:39 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Rash Generalised Stevens-Johnson Syndrome	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
	150 MG/ TWICE PER DAY/ ORAL				Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	SS		ORAL
	ORAL				Thyroxine Sodium	C		

Date:11/30/99ISR Number: 3414975-0Report Type:Periodic Company Report #A0102353A  
Age:73 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Fatigue Flushing Malaise	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
	150 MG/ PER DAY/ ORAL		Myocardial Infarction  Nausea					

Date:11/30/99ISR Number: 3414976-2Report Type:Periodic  
Age:18 YR Gender:Male I/FU:I

Company Report #A0101445A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG/ ORAL	Urticaria	Health Professional Company Representative	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:11/30/99ISR Number: 3414977-4Report Type:Periodic  
Age:18 YR Gender:Male I/FU:I

Company Report #A0101335A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Required 1 TABLET/ Intervention to TWICE PER Prevent Permanent DAY/ ORAL 11 DAY Impairment/Damage	Dermatitis	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/99ISR Number: 3414978-6Report Type:Periodic  
Age:21 YR Gender:Female I/FU:I

Company Report #A0101180A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG/ PER DAY/ ORAL	Grand Mal Convulsion	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
			Oral Contraceptive	C		

Date:11/30/99ISR Number: 3414979-8Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #A0100132A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/ TWICE PER DAY/ ORAL	Convulsion Insomnia	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
			Ortho-Novum	C		

Date:11/30/99ISR Number: 3414980-4Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0100142A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL	Agitation Delusion Dry Mouth Insomnia Mania Mood Altered Paranoia	Consumer	Zyban Tablet - Zyban	PS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Grand Mal Convulsion Memory Impairment	Consumer	Zyban Tablet -Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/	ORAL			Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	SS		ORAL
150 MG/	PER						
DAY/	ORAL						

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Amnesia Circulatory Collapse Convulsion	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/	ORAL	Disturbance In Attention Hypoaesthesia					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/02/99ISR Number: 3412862-5Report Type:Expedited (15-DaCompany Report #A0090299A  
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain Chest Pain Pericarditis	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
100 MG / PER DAY	7 DAY	Vomiting					

Date:12/02/99ISR Number: 3412864-9Report Type:Expedited (15-DaCompany Report #A0090004A  
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardiac Arrest Chest Discomfort Dyspnoea	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG		Pulmonary Embolism		Lorazepam Cefuroxime Axetil Combivent Budesonide Tetracycline Theophylline	C C C C C C		

Date:12/02/99ISR Number: 3412921-7Report Type:Expedited (15-DaCompany Report #A0103729A  
Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Sinus Tachycardia Suicide Attempt Tremor	Foreign	Ceftin (Formulation Unknown) (Cefuroxime Axetil) Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS SS		
150 MG ORAL				Ibuprofen (Formulation Unknown) (Ibuprofen)	SS		

Amoxicillin  
Trihydrate  
(Formulation  
Unknown)  
(Amoxicillin SS

Date:12/06/99ISR Number: 3414110-9Report Type:Expedited (15-DaCompany Report #A0105850A  
Age:17 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Intentional Misuse Suicidal Ideation Suicide Attempt	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupriopn Hydrochloride)	PS		ORAL

SEE TEXT

Date:12/06/99ISR Number: 3414153-5Report Type:Expedited (15-DaCompany Report #A0105840A  
Age:40 YR Gender:Female I/FU:I

Outcome	PT	Report Source
Hospitalization - Initial or Prolonged Other	Cerebrovascular Accident	Health Professional Company

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Representative

Dose	Duration	Product	Role	Manufacturer	Route
ORAL		Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		ORAL
		Sertraline Hydrochloride	C		
		Mirtazapine	C		

Date:12/06/99ISR Number: 3414174-2Report Type:Expedited (15-DaCompany Report #A0105839A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cerebrovascular Accident	Health Professional Company Representative	Wellbutrin (Bupropion Hydrochloride) (Formulation Unknown)	PS		ORAL
ORAL	1 WK			Fluoxetine Hydrochloride	C		

Date:12/06/99ISR Number: 3414176-6Report Type:Expedited (15-DaCompany Report #A0093402A  
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Breast Cancer Recurrent Metastases To Liver	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG							
PER DAY							
ORAL				Tamoxifen Paroxetine	C		

Date:12/06/99ISR Number: 3414177-8Report Type:Expedited (15-DaCompany Report #A0106235A

Age:39 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Abnormal Dreams Anaphylactic Shock Blood Pressure Decreased	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG PER DAY ORAL	Dermatitis Face Oedema Flushing Loss Of Consciousness Peripheral Coldness Rash Erythematous Skin Exfoliation Tongue Oedema		Clarithromycin	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/06/99ISR Number: 3414178-XReport Type:Expedited (15-DaCompany Report #A0106475A

Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthropathy Discomfort Swelling	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG		Urticaria					
TWICE PER DAY							
ORAL							

Date:12/06/99ISR Number: 3414181-XReport Type:Expedited (15-DaCompany Report #A0106383A

Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation Confusional State Disorientation	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG		Disturbance In Attention					
TWICE PER DAY		Flight Of Ideas					
ORAL		Headache Memory Impairment Mood Swings Nausea Priapism Rash Pruritic Tenderness		Salbutamol Sulphate	C		

Date:12/07/99ISR Number: 3415134-8Report Type:Expedited (15-DaCompany Report #A0106452A

Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hypoglycaemia Loss Of Consciousness	Foreign Consumer	Zyban Tablet - Zyban (Bupropion			

150 MG /  
TWICE PER DAY  
/ ORAL

Road Traffic Accident

Hydrochloride)

PS

ORAL

Date:12/07/99ISR Number: 3415135-XReport Type:Expedited (15-DaCompany Report #A0093420A  
Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Cardiomegaly Chest Discomfort Chest Pain	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY / ORAL	Drug Hypersensitivity Dyspnoea Hyperhidrosis Hypoaesthesia Pain In Jaw Photosensitivity Reaction Proctalgia Pruritus Rash Erythematous Throat Tightness Urethral Pain Urticaria White Blood Cell Count Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/08/99ISR Number: 3416792-4Report Type:Expedited (15-DaCompany Report #B0072373A

Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Agitation Alopecia Chills Dermatitis Atopic Dermatitis Exfoliative Extrasystoles Face Oedema Flushing Gastrooesophageal Reflux Disease Hypersensitivity Hypertension Muscle Spasms Palpitations Paraesthesia Photosensitivity Reaction Rash Erythematous Tachycardia Tremor Urinary Retention Urticaria	Foreign Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride) Thyroxine Sodium	PS C		

Date:12/08/99ISR Number: 3416795-XReport Type:Expedited (15-DaCompany Report #A0106237A

Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Chills Dermatitis Hypersensitivity Localised Oedema  Oedema Peripheral  Pruritus  Urticaria	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY / ORAL							

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Dose Duration 400.000 MG PO BID	Coma Complex Partial Seizures Drug Interaction	Health Professional Other	Abbott - Norvir Soft Gelatin Capsules Wellbutrin	PS SS	Abbott	ORAL ORAL
150.000 MG PO BID			Fortovase Zerit Epivir	C C C		

Outcome  
Death  
Hospitalization -  
Initial or Prolonged  
Required  
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Intervention to  
Prevent Permanent  
Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 TABLET, 30	IN 1 TIME(S),	Agitation Cardiac Arrest	Literature Health	Haldol (Tablet) (Haloperidol)	PS		ORAL
		Completed Suicide	Professional				
		Convulsion					
1 TABLET, 500	IN 1 TIME(S),	Intentional Misuse Tachycardia		Aspirin (Acetylsalicylic Acid)	SS		ORAL
		Toxicologic Test Abnormal					
		Ventricular Fibrillation					
100 MG, 30 IN	1 TIME(S),	Vomiting		Bupropion (Amfebutamone)	SS		ORAL

Date: 12/10/99  
ISR Number: 3419037-4  
Report Type: Expedited (15-DaCompany Report #A0104885A)  
Age: 19 YR  
Gender: Female  
I/FU: I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG / TWICE PER DAY / ORAL	Arthralgia Burning Sensation Erythema	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
		Erythema Multiforme					
		Lymphadenopathy					
		Malaise Pain Pruritus Pyrexia Serum Sickness					

Date:12/13/99ISR Number: 3420202-0Report Type:Expedited (15-DaCompany Report #A0106978A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness	Health Professional	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
100 MG/ THREE							
TIMES PER							
DAY/ ORAL							
200 MG/ THREE				Carbamazepine Tablet (Carbamazepaine)	SS		ORAL
TIMES PER							
DAY/ ORAL				Oestradiol	C		

Date:12/13/99ISR Number: 3420204-4Report Type:Expedited (15-DaCompany Report #A0055937A  
Age:55 YR Gender:Female I/FU:F

Outcome	PT
Disability	Asthenia Blood Glucose Increased Chills Dizziness Drug Hypersensitivity Headache Hyperhidrosis Hypertension

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL		Hypoaesthesia Mania Mental Impairment Mitral Valve Incompetence Muscle Twitching Musculoskeletal Stiffness Nervousness Paraesthesia Renal Impairment Tongue Oedema Tremor Urinary Incontinence		Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride) Conjugated Estrogens Vitamin Aspirin Melatonin	PS C C C C		ORAL

Date:12/13/99ISR Number: 3420692-3Report Type:Expedited (15-DaCompany Report #A0106200A  
Age:54 YR Gender:Female I/FU:I

Dose	Duration	Outcome	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL	2 WK	Hospitalization - Initial or Prolonged	Chest Pain Dysgeusia Dyspepsia Sinusitis	Consumer	Wellbutrin Tablets - Controlled Release (Bupropion Hydrochloride) Gabapentin Oxybutynin Baclofen Frusemide Rofecoxib	PS C C C C C		ORAL

Date:12/13/99ISR Number: 3421072-7Report Type:Expedited (15-DaCompany Report #A0073385A  
Age:45 YR Gender:Male I/FU:F

Dose	Duration	Outcome	PT	Report Source	Product	Role	Manufacturer	Route
		Hospitalization - Initial or Prolonged	Abnormal Behaviour Agitation	Literature Health	Wellbutrin Tablet-Controlled			

150 MG / PER	Arthralgia	Professional	Release (Bupropion	PS	ORAL
	Chest Pain		Hydrochloride)		
DAY / ORAL	Confusional State				
	Delirium		Thiothixene	C	
	Disorientation		Lithium Salt	C	
	Drug Hypersensitivity		Venlafaxine		
	Malaise		Hydrochloride	C	
	Myalgia		Thyroxine Sodium	C	
	Pyrexia		Haloperidol	C	
	Rash Morbilliform		Lorazepam	C	
	Rash Pruritic		Trazodone	C	
	Red Blood Cell		Olanzapine	C	
	Sedimentation Rate				
	Increased				
	Restlessness				
	Serum Sickness				

Date:12/14/99ISR Number: 3420932-0Report Type:Direct  
 Age:24 YR Gender:Male I/FU:I

Company Report #

Outcome PT  
 Hospitalization - Abdominal Pain  
 Initial or Prolonged Diarrhoea

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Hallucination Vomiting	Report Source	Product	Role	Manufacturer	Route
				Bupropion	PS		

Date:12/14/99ISR Number: 3421148-4Report Type:Expedited (15-DaCompany Report #  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG PO QD; Initial or Prolonged 150 MG PO BID		Anxiety Chest Pain		Bupropion Sr	PS		ORAL
		Dizziness		Ecotrin	C		
		Dyspnoea		Plavix	C		
		Hyperhidrosis		Atenolol	C		
		Hypotension		Imdur	C		
				Lipitor	C		
				Folate	C		
				Vitamin C	C		
				Vitamin E	C		

Date:12/14/99ISR Number: 3421152-6Report Type:Expedited (15-DaCompany Report #  
 Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG PO QD Initial or Prolonged 300MG PO QD		Cardiac Failure Congestive		Bupropion Sr	PS		ORAL
		Dyspnoea Exertional		Flagyl	C		
		Myocardial Infarction		Catapril	C		
		Orthopnoea		Prilosec	C		
		Oxygen Saturation Decreased		Atenolol	C		
				Zocor	C		
				Asa	C		
				Ntg	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abnormal Behaviour	Health	Meridia	PS		ORAL
15 MG UNK PO							
Life-Threatening		Bradycardia	Professional	Wellbutrin Sr	SS		ORAL
150 MG UNK PO							
Hospitalization -		Coma		Klonopin	SS		
Initial or Prolonged		Completed Suicide		Potassium	SS		
Required		Grand Mal Convulsion		Lasix	SS		
Intervention to		Hypoglycaemia		Alcohol	SS		
Prevent Permanent		Lung Disorder					
Impairment/Damage		Sinus Tachycardia					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Convulsion		Zyban 150 Glaxo			
Initial or Prolonged		Medication Error		Wellcome	PS	Glaxo Wellcome	
1 T BID							
				Celexa Forest	SS	Forest	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/16/99ISR Number: 3423545-XReport Type:Expedited (15-DaCompany Report #A0106621A  
Age:81 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aspiration Choking Muscle Rigidity Rhinorrhoea	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
100 MG / PER DAY / ORAL		Sepsis Vomiting		Donepezil Hcl Doxazosin Mesylate Vitamin E Glucosamine	C C C C		

Date:12/16/99ISR Number: 3423550-3Report Type:Expedited (15-DaCompany Report #A0104511A  
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other Required Intervention to 150 MG / Prevent Permanent TWICE PER DAY Impairment/Damage / ORAL		Complex Partial Seizures Drug Interaction	Health Professional Company Representative Other	Wellbutrin Tablet-Controlled Release (Bupropoin Hydrochloride)	PS		ORAL
400 MG / TWICE PLER DAY / ORAL				Ritonavir Solution (Ritonavir)	SS		ORAL
				Saquinavir Stavudine Lamivudine Atovaquone Loperamide Hydrochloride	C C C C C		

Date:12/16/99ISR Number: 3423555-2Report Type:Expedited (15-DaCompany Report #A0107071A  
Age:78 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Atrophy Confusional State Convulsion	Health Professional	Zyban Tablet -Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER DAY / ORAL	Delusion Disorientation Drug Interaction Miosis Muscle Rigidity Urinary Tract Infection Visual Disturbance		Hormones Multivitamin Citalopram Hydrobromide (Formulation Unknown) (Citalopram Hydrobromide)	C C I		
20 MG / UNK / UNKNOWN						

Date:12/16/99ISR Number: 3423560-6Report Type:Expedited (15-DaCompany Report #A0068941A  
Age:52 YR Gender:Male I/FU:F

Outcome	PT
Other	Agitation Conversion Disorder Dizziness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Feeling Abnormal Hallucination Hallucination, Auditory	Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / UNK		Nausea Nephrolithiasis Pain					
/ ORAL		Tremor Urinary Tract Infection					

Date:12/17/99ISR Number: 3425009-6Report Type:Expedited (15-DaCompany Report #A0107117A  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
Other		Insomnia Nausea					
150 MG/TWICE		Tremor					
PER DAY/ORAL		Visual Acuity Reduced					

Date:12/20/99ISR Number: 3425894-8Report Type:Expedited (15-DaCompany Report #A0100230A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Complications Of Maternal Exposure To Therapeutic Drugs	Study Health Professional Company	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		ORAL
Congenital Anomaly		Congenital Anomaly					
UNK / UNK /		Heart Disease Congenital	Representative				
ORAL							

Date:12/20/99ISR Number: 3425935-8Report Type:Expedited (15-DaCompany Report #A0104240A  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Required	400 MG / PER DAY/ ORAL	Aggression Child Abuse Mania Suicidal Ideation	Study Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage				Paracetamol	C		

Date:12/20/99ISR Number: 3426738-0Report Type:Expedited (15-DaCompany Report #A0080414A  
Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Disability	150 MG/TWICE PER DAY/ORAL	Arthralgia Joint Swelling Oedema Serum Sickness Urticaria	Foreign Health Professional	Bupropion Hydrochloride Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
				Lansoprazole Cisapride	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/21/99ISR Number: 3426699-4Report Type:Direct  
Age:42 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 2 DAILY Initial or Prolonged	Pruritus Urticaria		Zyban 150mg Glaxo	PS	Glaxo	

Date:12/21/99ISR Number: 3426765-3Report Type:Expedited (15-DaCompany Report #A0105273A  
Age:44 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - Initial or Prolonged 7 DAY Disability	Anaemia Blood Creatine Phosphokinase Increased Blood Creatinine Increased Candidiasis Confusional State Crush Injury Delirium Depressed Level Of Consciousness Disorientation Herpes Zoster Hyperphosphataemia Hypoaesthesia Hypocalcaemia Hypovolaemia Liver Function Test Abnormal Myocarditis Neuropathy Peripheral Paraesthesia Renal Failure Acute Rhabdomyolysis	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride) Ethanol (Formulation Unknown) (Alcohol)	PS SS		

Date:12/23/99ISR Number: 3430073-4Report Type:Expedited (15-DaCompany Report #990809-LX254  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required QHS PO		Dysphonia	Health Professional	Ambien	PS		ORAL
Intervention to 900.000 MG QD		Stereotypy	Professional	Lithium Carbonate	SS		ORAL
Prevent Permanent PO			Company				
Impairment/Damage 150.000 MG			Representative	Wellbutrin	SS		ORAL
TID PO				Other Cns Drugs Celecoxib	C C		

Date:12/23/99ISR Number: 3430277-0Report Type:Periodic Company Report #A0092326A  
Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Health Professional Company	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
SEE TEXT/ORAL			Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/23/99ISR Number: 3430279-4Report Type:Periodic  
Age:20 YR Gender:Female I/FU:F

Company Report #A0092331A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
SEE TEXT/ ORAL			Representative				

Date:12/23/99ISR Number: 3430281-2Report Type:Periodic  
Age:30 YR Gender:Female I/FU:F

Company Report #A0069375A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Grand Mal Convulsion Pain	Health Professional	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
Other		Soft Tissue Injury					
75 MG/ TWICE PER DAY/ ORAL		Tongue Disorder		Sertraline Hydrochloride	C		

Date:12/23/99ISR Number: 3430283-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0099384A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Overdose	Health Professional Company	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL			Representative				

Date:12/23/99ISR Number: 3430285-XReport Type:Periodic  
Age:8 YR Gender:Male I/FU:I

Company Report #A0097837A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Health Professional Company Representative	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		

Date:12/23/99ISR Number: 3430286-1Report Type:Periodic Company Report #A0097641A  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other Required 75 MG/ ORAL		Drug Interaction Grand Mal Convulsion Lacrimation Increased	Health Professional Company Representative	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage 30 MG/ ORAL		Productive Cough		Paroxetine Hydrochloride Tablet (Paroxetine Hydrochloride)	SS		ORAL
				Perphenazine	C		

Date:12/23/99ISR Number: 3430288-5Report Type:Periodic Company Report #A0096336A  
 Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source
Other	Convulsion Overdose	Health Professional

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Company Representative	Product	Role	Manufacturer	Route
			Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		
Date:12/23/99ISR Number: 3430290-3Report Type:Periodic Age:70 YR Gender:Female I/FU:I			Company Report #A0096393A			

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Grand Mal Convulsion	Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		
				Glibenclamide	C		
				Doxepin	C		
				Benazepril Hydrochloride	C		
				Diltiazem Hydrochloride	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other Required Intervention to 112.5 MG/ Prevent Permanent THREE TIMES Impairment/Damage PER ORAL		Convulsion	Health Professional Company Representative	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
				Salbutamol Sulphate	C		
				Prednisone	C		
				Clonidine	C		

Date:12/23/99ISR Number: 3430295-2Report Type:Periodic  
Age:15 YR Gender:Male I/FU:I

Company Report #A0094046A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion Overdose	Health Professional	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Date:12/23/99ISR Number: 3430298-8Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #A0093758A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Myocardial Infarction	Health Professional Other	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Omeprazole	C
Loratadine	C
Lorazepam	C
Conjugated Estrogens	C



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/23/99ISR Number: 3430301-5Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0091695A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - Initial or Prolonged SINGLE	Duration Abnormal Behaviour Delusion Hallucination Suicide Attempt	Health Professional	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
DOSE/ORAL			Phentermine Tablet (Phentermine)	SS		ORAL
37.5 MG/ORAL			Doxylamine Succinate Ketoprofen Erythromycin	C C C		

Date:12/23/99ISR Number: 3430303-9Report Type:Periodic  
Age:15 YR Gender:Male I/FU:I

Company Report #A0091099A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Duration Convulsion	Health Professional Company Representative	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
100 MG/ THREE TIMES PER DA ORAL			Methylphenidate Hcl Guanfacine Hydrochloride Methylphenidate Hcl	C C C C		

Date:12/23/99ISR Number: 3430306-4Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0090199A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Duration Stevens-Johnson Syndrome	Health Professional	Wellbutrin (Formulation			

ORAL  
Company Representative      Unknown) (Bupropion Hydrochloride)      PS      ORAL

Date:12/23/99ISR Number: 3430308-8Report Type:Periodic      Company Report #A0087645A  
Age:20 YR      Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Health Professional	Wellbutrin (Formulation Unknown) Lithium Salt	PS C		

Date:12/23/99ISR Number: 3430311-8Report Type:Periodic      Company Report #A0086778A  
Age:      Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Medication Error	Health Professional Company Representative	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		ORAL

100 MG/ PER

DAY/ ORAL

Wellbutrin  
Tablet-Controlled  
Release (Bupropion

FDA - Adverse Event Reporting System (AERS)

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100 MG/  
 SINGLE DOSE/  
 ORAL

Hydrochloride)      SS      ORAL

Date:12/23/99ISR Number: 3430315-5Report Type:Periodic      Company Report #A0084030A  
 Age:57 YR    Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Tongue Oedema	Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		ORAL
Other Required Intervention to Prevent Permanent Impairment/Damage				Conjugated Estrogens	C		

Date:12/23/99ISR Number: 3430318-0Report Type:Periodic      Company Report #A0083663A  
 Age:13 YR    Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Health Professional Company	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
			Representative	Methylphenidate Hcl	C		
				Erythromycin	C		
				Tretinoin	C		
				Melatonin	C		

Date:12/23/99ISR Number: 3430321-0Report Type:Periodic      Company Report #A0083479A  
 Age:46 YR    Gender:Male      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Grand Mal Convulsion	Health Professional	Wellbutrin (Formulation Unknown) (Bupropion			

ORAL  
Hydrochloride) PS ORAL  
Neuroleptic C

Date:12/23/99ISR Number: 3430324-6Report Type:Periodic Company Report #A0082070A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Agitation Chills Delusion Hearing Impaired	Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		ORAL
ORAL	3 WK	Insomnia Pyrexia Tremor Weight Decreased					

Date:12/27/99ISR Number: 3430560-9Report Type:Direct Company Report #  
Age:57 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Amnesia Coma Confusional State

FDA - Adverse Event Reporting System (AERS)

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Dose	Duration	Convulsion Muscle Rigidity Tremor	Report Source	Product	Role	Manufacturer	Route
150MG QD OR BID? PO				Bupropion	PS		ORAL

Date:12/27/99ISR Number: 3433476-7Report Type:Expedited (15-DaCompany Report #A0107781A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG / TWICE PER DAY / ORAL	Convulsion Drug Interaction	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
				Citalopram Hydrobromide (Citalopram Hydrobromide)	SS		

Date:12/27/99ISR Number: 3433482-2Report Type:Expedited (15-DaCompany Report #A0104499A  
Age:74 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged	150 MG / Required TWICE PER DAY Intervention to / ORAL	Asthenia Balance Disorder Cerebral Ischaemia Cerebrovascular Accident Confusional State	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
Prevent Permanent Impairment/Damage		Convulsion Dizziness Fall		Aspirin Atorvastatin Calcium	C C		

Headache  
Hemiplegia  
Paralysis  
Tremor

Date:12/28/99ISR Number: 3432984-2Report Type:Expedited (15-DaCompany Report #A0104783A

Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Disability 150 MG/TWICE PER DAY/ORAL	Antibody Test Positive Blindness Brain Oedema Chest Pain Dermatitis Difficulty In Walking Discomfort Optic Neuropathy Pruritus Serum Sickness Syncope Urticaria	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)  Ranitidine Hydrochloride Metronidazole Temazepam	PS   C C C		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/29/99ISR Number: 3435167-5Report Type:Expedited (15-DaCompany Report #001-0945-991279

Age:16 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1)PER ORAL;2) Other 3000 MG (AS ONE DOSE), PER ORAL; 3) PER ORAL	Grand Mal Convulsion Intentional Misuse Lethargy Loss Of Consciousness Suicide Attempt	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1) PER ORAL; 2) 4500 MG (AS ONE DOSE), PER ORAL; 3) PER			Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL

Date:01/03/00ISR Number: 3435491-6Report Type:Direct

Company Report #

Age:8 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 75MG BID X 1 Initial or Prolonged WEEK 1 WK 150MG BID X 2 DAYS	Convulsion		Wellbutrin 75mg  Wellbutrin Sr 150mg	PS  SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Disability 150 MG / TWICE PER DAY /ORAL		Blindness Brain Oedema Chest Pain Difficulty In Walking Optic Nerve Disorder Pruritus Serum Sickness Skin Disorder Syncope Urticaria	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)  Ranitidine Hydrochloride Metronidazole Temazepam	PS    C C C		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other TRANSDERMAL TTD 150 MG/ BID/ QD	1 PATCH/ QD/	Application Site Reaction	Consumer	Habitrol-Nicotine Transdermal 21mg-Nvch  Zyban-Bupropion Hcl 150mg-Glaxo  Antihypertensive	PS  SS C	Nvch  Glaxo	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/10/00ISR Number: 3441735-7Report Type:Expedited (15-DaCompany Report #A0108484A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG ORAL		Aggression Anxiety	Foreign Consumer	Zyban (Bupropion Hydrochloride)	PS		ORAL
		Decreased Activity Dermatitis Exfoliative Dysgeusia Dysphagia Face Oedema Fatigue Gingival Ulceration Halitosis Insomnia Lip Disorder Nail Avulsion Oedema Oral Intake Reduced Oral Pain Pain Pruritus Rash Erythematous Salivary Hypersecretion					

Date:01/10/00ISR Number: 3442243-XReport Type:Expedited (15-DaCompany Report #B0074965A

Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1		Laryngospasm Rash Pruritic	Foreign Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
TABLET/TWICE							
PER DAY/ORAL							

Date:01/12/00ISR Number: 3443168-6Report Type:Direct

Company Report #

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Disorientation		Zyban (Bupropion Sr)			
		Eye Pain		150 Mg	PS		ORAL
150MG PO QD		Headache					
		Neck Pain					
		Tremor					

Date:01/12/00ISR Number: 3443574-XReport Type:Expedited (15-DaCompany Report #224296  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion	Health	Bactrim			
Initial or Prolonged		Drug Interaction	Professional	(Sulfamethoxazole /			
		Loss Of Consciousness	Other	Trimethoprim)	PS		ORAL
1 DOSE FORM 2		Medication Error					
PER DAY ORAL							

				Wellbutrin			
				(Bupropion			
				Hydrochloride) 200			
				Mg	SS		ORAL
150 MG 2 PER							

				Prozac (Fluoxetine			
--	--	--	--	--------------------	--	--	--

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C

Date:01/12/00ISR Number: 3443588-XReport Type:Expedited (15-DaCompany Report #A0104783A

Age:44 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Disability 150 MG/ TWICE PER DAY/ ORAL	Autoimmune Disorder Brain Oedema Chest Pain Difficulty In Walking	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
	Optic Nerve Disorder Pruritus Serum Sickness Syncope Urticaria Visual Acuity Reduced		Ranitidine Hydrochloride Metronidazole Temazepam	C C C		

Date:01/12/00ISR Number: 3443591-XReport Type:Expedited (15-DaCompany Report #A0108167A

Age:44 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Disability 150 MG/ TWICE PER DAY/ ORAL	Blindness Brain Oedema Chest Pain Difficulty In Walking	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
	Discomfort Optic Nerve Disorder Pruritus Serum Sickness Syncope Urticaria		Ranitidine Hydrochloride Metronidazole Temazepam	C C C		

Date:01/12/00ISR Number: 3443593-3Report Type:Expedited (15-DaCompany Report #A0108666A

Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dehydration Dermatitis Exfoliative Hyperglycaemia	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL		Oral Mucosal Exfoliation Stevens-Johnson Syndrome		Glibenclamide Diltiazem Hydrochloride Loxapine Aspirin Losartan Potassium Conjugated Estrogens Thyroxine Sodium	C C C C C C C		

Date:01/12/00ISR Number: 3443761-0Report Type:Expedited (15-DaCompany Report #A0108441A  
Age:10 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Laboratory Test Abnormal Overdose	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/13/00ISR Number: 3443467-8Report Type:Direct  
 Age:37 YR Gender:Female I/FU:I

Company Report #USP 52773

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin Sr	PS	Catalytica Pharmaceuticals/Glax o Wellcome, Inc.	
				Wellbutrin	SS	Catalytica Pharmaceuticals/Glax o Wellcome, Inc.	

Date:01/14/00ISR Number: 3443753-1Report Type:Direct  
 Age:6 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema Multiforme	Health	Wellbutrin 75mg Glx	PS	Glx	
Other		Oedema Peripheral	Professional				
37.5MG TO		Synovitis					
START THEN							
75MG BID							
AM/NOON							

Date:01/14/00ISR Number: 3444512-6Report Type:Expedited (15-DaCompany Report #A0102623A  
 Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Completed Suicide Intentional Misuse	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		
Death			Other				
OPHTHALMIC	150MG SEE			Fluoxetine Hydrochloride Capsule (Fluoxetine Hydochloride)	SS		ORAL

20MG SEE TEXT

ORAL

Venlafaxine  
Hydrochloride Tablet  
(Venlafaxine  
Hydrochloride)

SS

ORAL

37.5MG SEE

TEXT ORAL

Date:01/18/00ISR Number: 3444741-1Report Type:Direct  
Age:52 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Dyspepsia Erythema		Zyban 150mg Glaxo Wellcome	PS	Glaxo Wellcome	ORAL
1T PO QD X 3		Erythema Multiforme					
DAYS THEN 1T		Flatulence					
PO BID		Mucosal Inflammation		Lipitor	C		
		Rash Pruritic		Premarin	C		
		Urticaria		Vitamin E	C		
		Vaginal Infection		Multivitamins	C		
				Stool Softener	C		

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150 MG/TWICE	Erythema Multiforme	Hydrochloride)	PS	ORAL
PER DAY/ORAL	Hyperglycaemia			
	Stevens-Johnson Syndrome	Glibenclamide	C	
		Diltiazem		
		Hydrochloride	C	
		Loxapine	C	
		Aspirin	C	
		Losartan Potassium	C	
		Conjugated Estrogens	C	
		Thyroxine Sodium	C	
		Metformin	C	

Date:01/18/00ISR Number: 3445757-1Report Type:Expedited (15-DaCompany Report #A0109272A  
Age:40 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Asthma	Foreign	Zyban Tablet - Zyban			
Initial or Prolonged	Pruritus	Health	(Bupropion			
	Urticaria	Professional	Hydrochloride)	PS		ORAL
150 MG /						
UNKNOWN /						
ORAL						



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/19/00ISR Number: 3445319-6Report Type:Expedited (15-DaCompany Report #A0108878A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pharyngeal Oedema Tongue Oedema	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL							

Date:01/20/00ISR Number: 3445588-2Report Type:Expedited (15-DaCompany Report #A0109223A

Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthritis	Health	Zyban	PS		ORAL
150 MG/ TWICE							
PER DAY /		Immobile	Professional				
ORAL		Urticaria					

Date:01/24/00ISR Number: 3446533-6Report Type:Expedited (15-DaCompany Report #A0108853A

Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization - Initial or Prolonged		Anaphylactic Reaction Dysphagia Face Oedema Pruritus	Foreign Health Professional	Bupropion Hydrochloride Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /							
TWICE PER DAY		Shock					
/ ORAL		Swelling					
		Urticaria		Fosinopril Sodium Hydrochlorothiazide Medroxyprogesterone Ace.	C C C		

Amlodipine

C

Date:01/24/00ISR Number: 3446625-1Report Type:Expedited (15-DaCompany Report #B0075853A

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Hyperventilation	Foreign	Zyban Tablet-Zyban			
Initial or Prolonged	Malaise	Health	(Bupropion			
	Nausea	Professional	Hydrochloride)	PS		

Date:01/24/00ISR Number: 3446789-XReport Type:Expedited (15-DaCompany Report #A0108087A

Age:39 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Amnesia	Consumer	Zyban Tablet-Zyban			
Initial or Prolonged	Convulsion		(Bupropion			
	Dysphemia		Hydrochloride)	PS		ORAL
150 MG/TWICE	Electroencephalogram					
PER DAY/ORAL	Abnormal		Blood Pressure			
	Sedation		Medication	C		
	Tremor		Hormones	C		

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Date:01/24/00ISR Number: 3446792-XReport Type:Expedited (15-DaCompany Report #A0109400A  
Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Amnesia Cognitive Disorder Convulsion	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE DAILY/ORAL		Dizziness Fall Head Injury Tremor					

Date:01/24/00ISR Number: 3446794-3Report Type:Expedited (15-DaCompany Report #A0109220A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Pressure Decreased Bundle Branch Block Left Drug Level Above Therapeutic	Literature Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		ORAL
4.5 G/ SINGLE DOSE/ORAL		Drug Toxicity Electrocardiogram Qrs Complex Prolonged Electrocardiogram Qt Prolonged Heart Rate Increased Overdose Tachycardia		Paroxetine Paracetamol	C C		

Date:01/24/00ISR Number: 3446801-8Report Type:Expedited (15-DaCompany Report #A0109367A  
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Diabetes Mellitus Non-Insulin-Dependent	Health Professional Other	Wellbutrin (Formulation Unknown) (Bupropion			

ORAL				Hydrochloride)	PS	ORAL
				Risperidone (Formulation Unknown) (Risperidone)	SS	ORAL

Date:01/24/00ISR Number: 3447110-3Report Type:Expedited (15-DaCompany Report #A0109619A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abnormal Behaviour Burns Third Degree Mania	Health Professional Company Representative	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:01/24/00ISR Number: 3447480-6Report Type:Expedited (15-DaCompany Report #A0106235A  
 Age:39 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Anaphylactic Shock Face Oedema Flushing

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Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Hypotension Loss Of Consciousness Peripheral Coldness Rash Erythematous Skin Exfoliation Tongue Oedema	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
				Clarithromycin	C		

Date:01/27/00ISR Number: 3447358-8Report Type:Expedited (15-DaCompany Report #2000001499-1  
Age:47 YR Gender:Male I/FU:I

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
2.0 DAILY TRANSDERMAL	21 MILLIGRAMS	Amnesia Cerebrovascular Accident Myocardial Infarction Sinusitis	Consumer	Nicoderm Cq 21 Mg	PS	Smithkline Beecham Consumer Healthcare	
150 MILLIGRAMS 2.0 DAILY ORAL				Zyban (Bupropion Hcl) Glaxo Wellcome, Inc.	SS	Glaxo Wellcome, Inc.	ORAL

Date:01/27/00ISR Number: 3448419-XReport Type:Periodic Company Report #9944261  
Age: Gender:Male I/FU:I

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
50.00 MG		Drug Effect Decreased	Consumer	Viagra Tablets	PS		ORAL

TOTAL:PRN:ORA Drug Ineffective  
 L Drug Interaction  
 Wellbutrin SS ORAL  
 300.00 MG

TOTAL:PID:ORA  
 L  
 Norvasc C  
 Accupril C  
 Zantac C

Date:01/28/00ISR Number: 3448015-4Report Type:Expedited (15-DaCompany Report #A0103586A  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other ORAL		Coma Confusional State Convulsion  Grand Mal Convulsion Loss Of Consciousness Sedation	Foreign	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL

Date:01/28/00ISR Number: 3448016-6Report Type:Expedited (15-DaCompany Report #A0109909A  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged  150 MG /		Cerebrovascular Accident Headache Visual Acuity Reduced	Foreign Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

TWICE PER DAY  
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Freedom Of Information (FOI) Report

/ ORAL

Date:01/28/00ISR Number: 3448280-3Report Type:Expedited (15-DaCompany Report #S99-USA-02273-01  
 Age:78 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 20 MG QD PO	Brain Scan Abnormal Coma	Health Professional	Celexa (Citalopram Hydrochloride)	PS		ORAL
40 MG QD PO	Confusional State Convulsion		Celexa (Citalopram Hydrobromide)	SS		ORAL
150 MG BID PO	Decreased Interest Dermatitis Atopic		Zyban (Amfebutamone Hydrochloride)	SS		ORAL
	Disorientation		B Complex	C		
	Insomnia		Synthroid	C		
	Lethargy		Folic Acid	C		
	Memory Impairment					
	Miosis					
	Muscle Rigidity					
	Urinary Tract Infection					
	Visual Disturbance					

Date:01/31/00ISR Number: 3447850-6Report Type:Direct  
 Age:64 YR Gender:Female I/FU:I Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Clonic Convulsion Gait Disturbance Parkinson'S Disease Speech Disorder		Bupropion (Wellbutrin) 100mg Trazodone	PS C		

Date:01/31/00ISR Number: 3448785-5Report Type:Direct  
 Age:62 YR Gender:Male I/FU:I Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Required	Tremor	Health	Welbutrin 150 Mg	PS	ORAL
PO BID					
Intervention to		Professional	Rocalcetrol	C	
Prevent Permanent			Lotensin	C	
Impairment/Damage			Erythropoietin	C	
			Iron	C	
			Allopurinol	C	
			Renigel	C	
			Lipitor	C	
			Pepcid	C	

Date:02/02/00ISR Number: 3450944-2Report Type:Expedited (15-DaCompany Report #A0103729A  
Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Convulsion Intentional Misuse Sinus Tachycardia	Foreign	Zyban Tablet -Zyban (Bupropion Hydrohchloride)	PS		ORAL
150 MG/ ORAL		Suicide Attempt Tremor Vomiting		Ceftin (Formualation Unknown) (Cefuroxime Axetil) Ibuprofen (Formulation Unknown) (Ibuprofen) Amoxycillin	SS		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Trihydrate  
(Formulation  
Unknown)  
(Amoxicillin SS

Date:02/02/00ISR Number: 3451155-7Report Type:Expedited (15-DaCompany Report #A0109223A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Angiopathy Arthralgia Arthritis	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY / ORAL		Immobile Inflammation Rheumatoid Arthritis Urticaria					

Date:02/02/00ISR Number: 3451869-9Report Type:Periodic Company Report #A0101377A  
Age:13 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Amnesia Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
75 SEE TEXT / TWICE PER DAY / ORAL	6 MON			Desipramine Methylphenidate Hcl	C C		

Date:02/02/00ISR Number: 3451871-7Report Type:Periodic Company Report #A0102784A  
Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Drug Interaction	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
1 TABLET / TWICE PER DAY/ ORAL				Desipramine Hydrochloride (Formulation Unknown) (Desipramine	SS		
UNK/ UNK/ UNKNOWN							

Date:02/02/00ISR Number: 3451874-2Report Type:Periodic Company Report #A0102910A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Electroencephalogram Abnormal Weight Increased	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY / ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/02/00ISR Number: 3451876-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0103516A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY / ORAL							

Date:02/02/00ISR Number: 3451878-XReport Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0103526A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY/ ORAL							

Date:02/02/00ISR Number: 3451880-8Report Type:Periodic  
 Age:37 YR Gender:Male I/FU:I

Company Report #A0104375A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthralgia Asthenia Back Pain Chest Pain	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER DAY/ ORAL							
		Difficulty In Walking					
		Dysphagia		Ziac	C		

Fatigue  
Hypersensitivity  
Joint Swelling  
Nausea  
Oedema Peripheral  
Oral Intake Reduced  
Urticaria

Date:02/02/00ISR Number: 3451882-1Report Type:Periodic  
Age:15 YR Gender:Female I/FU:I

Company Report #A0104843A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER							
DAY / ORAL							

Date:02/02/00ISR Number: 3451885-7Report Type:Periodic  
Age:18 YR Gender:Female I/FU:I

Company Report #A0105026A

Outcome  
Hospitalization -  
Initial or Prolonged  
Other  
Required

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Intervention to Prevent Permanent Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1500 MG/		Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
SINGLE DOSE /		Intentional Misuse					
ORAL		Suicide Attempt		Ibuprofen Tablet (Ibuprefen)	SS		ORAL
50 TABLET /							
SINGLE DOSE /							
ORAL							

Date:02/02/00ISR Number: 3451887-0Report Type:Periodic Company Report #A0105971A  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:02/02/00ISR Number: 3451889-4Report Type:Periodic Company Report #A0108085A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Grand Mal Convulsion	Health Professional Company	Wellbutrin Tablet-Controlled Release (Bupropion			

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
450 MG/ PER			Representative	Hydrochloride)	PS		ORAL
DAY / ORAL							
Date:02/02/00ISR Number: 3451891-2Report Type:Periodic			Company Report #A0081121A				
Age:37 YR Gender:Female I/FU:F							
Other Required Intervention to Prevent Permanent Impairment/Damage TWICE PER DAY		Conjunctival Hyperaemia Depression Dermatitis Dermatitis Exfoliative Hypersensitivity Oral Intake Reduced	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
/ ORAL		Periorbital Oedema Sedation Suicidal Ideation		Wellbutrin Tablet (Bupropion Hydrochloride)	SS		ORAL
UNK/ UNK/		Suicide Attempt					
ORAL				Fluoxetine Hydrochloride	C		

Date:02/02/00ISR Number: 3451893-6Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #A0100421A

Outcome	PT
Hospitalization - Initial or Prolonged	Convulsion Disorientation

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sedation

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
UNK/ UNK/	UNKNOWN	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		
UNK/ UNK/	UNKNOWN		Nutritional Supplement (Formulation Unknown) (Nutritional)	SS		
			Semidodium Valproate	C		

Date:02/02/00ISR Number: 3451895-XReport Type:Periodic Company Report #A0101936A  
 Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other Required Intervention to Prevent Permanent Impairment/Damage 150 MG/ TWICE PER DAY/ ORAL		Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
				Hydrochlorothiazide	C		
				Phentermine Hydrochloride	C		
				Fluoxetine Hydrochloride	C		

Date:02/02/00ISR Number: 3451898-5Report Type:Periodic Company Report #A0102154A  
 Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Overdose	Consumer	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
300 MG /							
TWICE PER							
DAY/ ORAL							

Date:02/03/00ISR Number: 3451165-XReport Type:Expedited (15-DaCompany Report #A0103727A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Agitation Alcohol Interaction Alcohol Withdrawal	Foreign	Zyban Tablet -Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG, ORAL		Syndrome Amnesia Blood Alcohol Increased Convulsion Delirium Irritability Overdose Suicide Attempt Tachycardia		Ethanol (Formulation Unknown) (Alcohol) Lorazepam (Formulation Unknown) (Lorazepam)	SS SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/03/00ISR Number: 3451184-3Report Type:Expedited (15-DaCompany Report #A0096983A

Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Convulsion Headache Petit Mal Epilepsy	Foreign	Wellbutrin Tablet - Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL				Dilantin Semisodium Valproate Ibuprofen	C C C		

Date:02/04/00ISR Number: 3452543-5Report Type:Direct

Company Report #

Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 100 3X A DAY Prevent Permanent Impairment/Damage		Amnesia Anger Drug Dependence Fatigue Influenza Like Illness Irritability Libido Increased		Wellbutrin 100mg Glaxowellcome?	PS	Glaxowellcome	

Date:02/04/00ISR Number: 3452825-7Report Type:Expedited (15-DaCompany Report #A0107356A

Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other  150 MG/ ORAL		Chills Chromaturia Drug Interaction  Ear Haemorrhage Haematuria Haemorrhage International Normalised Ratio Increased Prothrombin Time	Consumer	Zyban Tablets - Zyban (Bupropion Hydrochloride)	PS		ORAL
				Warfarin Sodium (Formulation Unknow) (Warfarin Sodium)	SS		

Prolonged  
Pyrexia  
Renal Colic  
Tremor

Date:02/04/00ISR Number: 3452827-0Report Type:Expedited (15-DaCompany Report #A0106892A

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage 150 MG/ TWICE PER DAY/ ORAL		Coordination Abnormal Depression Disturbance In Attention Drug Interaction Drug Level Above	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
300 MG/ AT NIGHT/ ORAL / YEARS		Therapeutic Dysphoria Increased Appetite Sedation		Imipramine Tablet (Imipramine)	SS		ORAL
				Inhaler	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/04/00ISR Number: 3452837-3Report Type:Expedited (15-DaCompany Report #A0110172A  
 Age:30 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/ TWICE PER DAY/ ORAL	Dry Mouth Feeling Abnormal Hiatus Hernia Tremor	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:02/04/00ISR Number: 3452839-7Report Type:Expedited (15-DaCompany Report #B0076434A  
 Age:50 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/ PER DAY/ ORAL	Chest Discomfort Chest Pain	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
			Aspirin	C		
			Doxycycline	C		
			Hyzaar	C		
			Beclomethasone+Salbu tamol	C		

Date:02/04/00ISR Number: 3452979-2Report Type:Expedited (15-DaCompany Report #A0105554A  
 Age:23 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged UNK/SEE TEXT/ORAL	Agranulocytosis Asthenia Bone Marrow Depression Corynebacterium Infection Dizziness Postural Ecchymosis Fatigue	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
			Fluticasone Propionate	C		

Gram Stain Positive  
 Influenza Like Illness  
 Libido Decreased  
 Lung Infiltration  
 Neutropenia  
 Pain  
 Pharyngolaryngeal Pain  
 Pleuritic Pain  
 Pneumonia  
 Pyrexia  
 Sedation  
 Staphylococcal Infection  
 Suicidal Ideation

Fexofenadine C  
 Hydrochlorid C  
 Budesonide C  
 Oral Contraceptive C

Date:02/08/00ISR Number: 3454305-1Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Calcinosis		Klonopin	PS		
		Congenital Anomaly		Paxil	SS		
		Nervous System Disorder		Depakote	SS		
				Gabapentin	SS		
				Wellbutrin	SS		

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Freedom Of Information (FOI) Report

Date:02/09/00ISR Number: 3455517-3Report Type:Expedited (15-DaCompany Report #A0110103A  
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage 150 MG / TWICE PER DAY  / ORAL		Colitis Ischaemic Rectal Haemorrhage Syncope	Study Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
				Birth Control	C		

Date:02/09/00ISR Number: 3455518-5Report Type:Expedited (15-DaCompany Report #A0110688A  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged 150 MG / THREE TIMES PER DAY ORAL		Pneumonia Renal Failure Wegener'S Granulomatosis	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
				Olanzapine	C		
				Alprazolam	C		

Date:02/09/00ISR Number: 3456962-2Report Type:Direct Company Report #  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dermatitis Difficulty In Walking Erythema Multiforme Pruritus		Wellbutrin	PS		

Date:02/11/00ISR Number: 3456737-4Report Type:Direct  
Age:36 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Wellbutrin Sr 150mg	PS		
1 TAB 2X/DAY, MID-MORNING AND 4 PM		Anger Drug Dependence Emotional Disorder Euphoric Mood Marital Problem Nightmare Personality Change Psychotic Disorder		Vitamins Tylenol	C C		

Date:02/11/00ISR Number: 3456934-8Report Type:Expedited (15-DaCompany Report #A0110477A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL		Abdominal Pain Faecaloma	Consumer	Zyban	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/11/00ISR Number: 3456936-1Report Type:Expedited (15-DaCompany Report #A0110910A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abdominal Pain Upper	Foreign	Zyban	PS		ORAL
Life-Threatening		Blood Disorder	Health				
Disability		Cardiac Arrest Influenza	Professional	Pain Medication	C		

Date:02/11/00ISR Number: 3456982-8Report Type:Expedited (15-DaCompany Report #A0110682A  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required		Amnesia Cerebrovascular Accident Loss Of Consciousness	Consumer Other	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage TOPICAL		Myocardial Infarction Sinusitis		Nicotine Patch (Nicotine)	SS		
	42MG PER DAY						

Date:02/11/00ISR Number: 3456984-1Report Type:Expedited (15-DaCompany Report #A0110609A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Erythema Multiforme Hyperlipidaemia Hypersensitivity Leukocytosis Pain Prostatic Specific Antigen Increased Purpura Pyrexia	Consumer Other	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		

Rash Pruritic  
Sinusitis  
Urticaria

Date:02/14/00ISR Number: 3457280-9Report Type:Direct  
Age:30 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin 150 Mg			
Other		Chest Pain		Glaxxo Wellcome	PS		
		Confusional State					
		Headache					
		Memory Impairment					

Date:02/14/00ISR Number: 3457648-0Report Type:Expedited (15-DaCompany Report #A0110940A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/00ISR Number: 3457649-2Report Type:Expedited (15-DaCompany Report #A0102044A

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly 150 MG / TWICE PER DAY / ORAL	Duration Complications Of Maternal Exposure To Therapeutic Drugs Gastroschisis Systemic Candida Ultrasound Antenatal Screen Abnormal	Foreign Study Health Professional Company Representative	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:02/14/00ISR Number: 3457651-0Report Type:Expedited (15-DaCompany Report #A0102884A

Age:45 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Required 150 MG / Intervention to TWICE PER DAY Prevent Permanent / ORAL Impairment/Damage	Blister Blood Glucose Increased Dyspnoea Eyelid Oedema Nail Discolouration Oedema Peripheral Rash Pruritic Stevens-Johnson Syndrome Weight Increased	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:02/14/00ISR Number: 3457655-8Report Type:Expedited (15-DaCompany Report #A0106621A

Age:81 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Hospitalization - Initial or Prolonged	Aspiration Choking Malaise	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion			

100 MG / PER	Muscle Rigidity	Hydrochloride)	PS	ORAL
DAY / ORAL	Rhinorrhoea			
	Sepsis	Donepezil Hcl	C	
	Vomiting	Doxazosin Mesylate	C	
		Vitamin E	C	
		Glucosamine	C	

Date:02/14/00ISR Number: 3457855-7Report Type:Expedited (15-DaCompany Report #A0110613A  
 Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Chest Pain Cold Sweat Hyperhidrosis	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE PER DAY /	Nausea					
ORAL			Nitroglycerin Isosorbide Dinitrate	C C		

Date:02/14/00ISR Number: 3459567-2Report Type:Periodic Company Report #001-0945-980906  
 Age:24 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged PER ORAL	Stevens-Johnson Syndrome	Health Professional	Neurontin (Gabapentin)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

300 MG (150  
 MG,BID), PER  
 ORAL  
 Zyban (Amfebutamone) SS ORAL

Date:02/14/00ISR Number: 3459611-2Report Type:Periodic Company Report #001-0945-990106  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Agitation Drug Level Above	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (DAILY), PER ORAL		Therapeutic Feeling Drunk Nausea		Lithobid (Lithium Carbonate)	SS		ORAL
PER ORAL				Zyprexa (Olanzapine)	SS		ORAL
PER ORAL				Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL
PER ORAL				Ambien (Zolpidem Tartrate)	SS		ORAL
PER ORAL				Zoloft (Sertraline Hydrochloride)	SS		ORAL
PER ORAL				Klonopin (Clonazepam)	SS		ORAL
PER ORAL				Benzotropine (Benzatropine Mesilate)	SS		ORAL
				Inderal (Propranolol Hydrochloride)	C		

Date:02/15/00ISR Number: 3457366-9Report Type:Expedited (15-DaCompany Report #A0110939A  
Age:50 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Anxiety Depression Feeling Abnormal	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER DAY / ORAL	Mental Disorder					
150 MG / TWICE PER DAY / ORAL	Mood Swings		Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	SS		ORAL

Date:02/15/00ISR Number: 3458422-1Report Type:Expedited (15-DaCompany Report #A0111455A  
Age:19 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Blood Creatine Phosphokinase Increased Chest Pain	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
UNK / UNK / ORAL	Myocarditis					
			Cannabis Mushrooms	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/15/00ISR Number: 3458451-8Report Type:Expedited (15-DaCompany Report #A0102623A

Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / SEE			Other				
TEXT / ORAL				Fluoxetine Hydrochloride Capsule (Fluoxetine Hydrochloride)	SS		ORAL
20 MG / SEE							
TEXT / ORAL				Venflaxine Hydrochloride Tablet (Venlafaxine Hydrochloride)	SS		ORAL
37.5 MG / SEE							
TEXT / ORAL							

Date:02/15/00ISR Number: 3458516-0Report Type:Periodic Company Report #M094176

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Lethargy	Health Professional	Serzone Tabs (Nefazodone Hcl)	PS		ORAL
ORAL				Wellbutrin (Bupropion Hcl)	SS		ORAL
ORAL				Risperidone	SS		ORAL

Date:02/17/00ISR Number: 3459083-8Report Type:Direct

Company Report #

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required				Zyban Sustained			
Intervention to Prevent Permanent Impairment/Damage		Colitis Ischaemic Mucous Membrane Disorder Rectal Haemorrhage Syncope		Release 150 Mg Tablet Glaxo Wellcome	PS	Glaxo Wellcome	ORAL
150MG TWICE							
DAILY ORAL				Birth Control Pills Acupril	C		

Date:02/18/00ISR Number: 3460142-4Report Type:Periodic Company Report #HQ9426222DEC1999  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health	Effexor	PS		ORAL
600 MG			Professional				
OVERDOSE							
AMOUNT, ORAL				Prozac	SS		ORAL
440 MG							
OVERDOSE							
AMOUNT, ORAL				Wellbutrin	SS		ORAL
APPROX. 12000							
MG OVERDOSE							
AMOUNT, ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/18/00ISR Number: 3460213-2Report Type:Expedited (15-DaCompany Report #HQ1070415FEB2000  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other UNKNOWN TO 75 MG DAILY	3 YR	Cervix Disorder Depression Headache Menstruation Irregular Soft Tissue Disorder	Consumer	Effexor Xr  Wellbutrin Anaprox	PS  SS C		

Date:02/22/00ISR Number: 3461030-XReport Type:Expedited (15-DaCompany Report #A0096492A  
Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged 150 MG / Disability TWICE PER DAY Required / ORAL Intervention to Prevent Permanent Impairment/Damage		Dysphagia Dyspnoea Embolism Eye Disorder Oedema Peripheral Swelling	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)  Paracetamol	PS  C		ORAL

Date:02/22/00ISR Number: 3461213-9Report Type:Expedited (15-DaCompany Report #A0112060A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other  150 MG / TWICE PER DAY		Eye Pain Intraocular Pressure Increased Open Angle Glaucoma Optic Nerve Disorder	Consumer	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

/ ORAL

Date:02/22/00ISR Number: 3461315-7Report Type:Expedited (15-DaCompany Report #A0111797A

Age:67 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Arrhythmia	Study	Zyban Tablet - Zyban			
Initial or Prolonged	Confusional State	Health	(Bupropion			
150 MG PER	Dehydration	Professional	Hydrochloride)	PS		ORAL
DAY ORAL	Depressed Level Of					
1 TABLET PER	Consciousness		Placebo Tablet	SS		ORAL
DAY ORAL	Dizziness					
	Dysarthria		Clopidogrel			
	Hypotension		Bisulphate	C		
	Ventricular Tachycardia		Atenolol	C		
			Lisinopril	C		
			Omeprazole	C		
			Aspirin	C		
			Atorvastatin Calcium	C		
			Nitroglycerin	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/22/00ISR Number: 3461318-2Report Type:Expedited (15-DaCompany Report #A0110611A

Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Ineffective Drug Interaction Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
350 MG, PER DAY, ORAL				Venlafaxine Hydrochloride (Formulation Unknown) (Venlafaxine	SS		ORAL
37.5 MG PER DAY ORAL				Hormones	C		

Date:02/23/00ISR Number: 3461662-9Report Type:Expedited (15-DaCompany Report #A0110172A

Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Chest Pain Disorientation Dry Mouth	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS	Zyban	ORAL
150 MG TWICE PER DAY ORAL		Hiatus Hernia  Malaise Myocardial Infarction Nausea Tremor					

Date:02/23/00ISR Number: 3461725-8Report Type:Expedited (15-DaCompany Report #A0111148A

Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Disability	Amnesia	Health	Wellbutrin	
	Clonic Convulsion	Professional	(Formulation	
	Dysgraphia		Unknown) (Bupropion	
100 MG / UNK	Gait Disturbance		Hydrochloride)	PS
				ORAL
/ ORAL	Muscle Rigidity			
	Parkinson'S Disease		Trazodone	C
	Skin Disorder		Estratest	C
	Speech Disorder		Thyroxine Sodium	C
			Lisinopril	C
			Flunisolide	C
			Salbutamol Sulphate	C
			Clorazepate	
			Dipotassium	C
			Salmeterol Xinafoate	C
			Depression	C
			Asthma	C

Date:02/23/00ISR Number: 3462465-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #9937752

Outcome PT  
Other Dizziness  
Headache  
Influenza  
Personality Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vomiting

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
100.00 MG		Consumer	Zoloft Tablets	PS		ORAL
TOTAL: BID:		Health				
ORAL		Professional				
300.00 MG			Wellbutrin	SS		ORAL
TOTAL: BID:						
ORAL						

Date:02/23/00ISR Number: 3462954-XReport Type:Periodic Company Report #9933726  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression	Health	Zoloft Tablets	PS		ORAL
ORAL		Hyperhidrosis	Professional	Wellbutrin	SS		ORAL
ORAL		Nausea		Xanax	C		
		Pain		Prempro	C		
		Sedation		Multivitamins	C		
		Stupor					
		Thinking Abnormal					
		Yawning					

Date:02/23/00ISR Number: 3465744-7Report Type:Periodic Company Report #9910420  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Consumer	Zoloft Tablets	PS		ORAL
50.00 MG		Drug Ineffective	Health				
TOTAL:DAILY:0							

RAL	Tremor	Professional				
ORAL	Vertigo		Wellbutrin	SS		ORAL
			Advil	C		

Date:02/23/00ISR Number: 3466594-8Report Type:Periodic Company Report #9951766  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia	Health	Zoloft Tablets	PS		ORAL
100.00 MG			Professional				
TOTAL DAILY							

ORAL				Wellbutrin	SS		
				Birth Control	SS		

Date:02/23/00ISR Number: 3467331-3Report Type:Periodic Company Report #9919213  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression	Health	Zoloft Tablets	PS		
50.00 MG			Professional				
TOTAL:DAILY							

				Zyban	SS		
				Nizoral Shampoo	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/23/00ISR Number: 3467779-7Report Type:Periodic  
 Age:52 YR Gender:Male I/FU:I

Company Report #9937968

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100.00 MG		Abdominal Pain	Health	Zoloft Tablets	PS		ORAL
Initial or Prolonged TOTAL DAILY Required ORAL		Anorexia	Professional				
Intervention to 150.00 MG		Jaundice		Bupropion	SS		ORAL
Prevent Permanent TOTAL BID Impairment/Damage ORAL		Nausea					
		Pyrexia					
		Vomiting					

Date:02/23/00ISR Number: 3468259-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #9904353

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100MG TOTAL; DAILY; ORAL		Anxiety	Consumer	Zoloft Tablets	PS		ORAL
		Tremor					
				Wellbutrin Xanax	SS C		

Date:02/23/00ISR Number: 3469832-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #9944600

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100.00 MG		Asthenia	Consumer	Zoloft Tablets	PS		ORAL
TOTAL:DAILY:0 RAL		Dizziness					
		Drug Withdrawal Syndrome					

ORAL		Thinking Abnormal		Wellbutrin	SS		ORAL
		Weight Increased		Ambien	C		

Date:02/23/00ISR Number: 3470125-6Report Type:Periodic Company Report #9948895  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Health	Zoloft Tablets	PS		ORAL
100.00 MG		Attention	Professional				
TOTAL:DAILY:0		Deficit/Hyperactivity					
RAL		Disorder		Effexor	SS		
		Drug Ineffective		Wellbutrin	SS		
		Hostility					
		Hyperhidrosis					

Date:02/23/00ISR Number: 3470470-4Report Type:Periodic Company Report #9943253  
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Prolactin Increased	Health	Zoloft Tablets	PS		ORAL
150.00 MG		Depression	Professional				
TOTAL: ORAL		Female Sexual Dysfunction		Wellbutrin	SS		ORAL
150.00 MG		Galactorrhoea					
TOTAL: DAILY:							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/23/00ISR Number: 3565792-2Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0113107A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diabetes Mellitus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Therapeutic Response					
TWICE PER DAY		Decreased					
/ ORAL				Metronidazole	C		
				Cephalexin	C		

Date:02/24/00ISR Number: 3463088-0Report Type:Expedited (15-DaCompany Report #A0110613A  
 Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain	Foreign	Bupropion			
Hospitalization -		Cold Sweat	Consumer	Hydrochloride			
Initial or Prolonged		Hyperhidrosis		Tablet-Zyban			
		Nausea		(Bupropion			
		Panic Attack		Hydrochloride)	PS		ORAL
150 MG /							
TWICE PER							
DAY/ ORAL				Nitroglycerin	C		
				Isosorbide Dinitrate	C		

Date:02/24/00ISR Number: 3463203-9Report Type:Direct  
 Age:29 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression		Bupropion Sr	PS		
Life-Threatening		Cognitive Disorder		Tetracycline	SS		
Hospitalization -		Coma		Divalpro Ex	SS		
Initial or Prolonged		Confusional State					
		Drug Level Below					
		Therapeutic					

Ecchymosis  
Fatigue  
Folliculitis  
Headache  
Hypotension  
Influenza Like Illness  
Movement Disorder  
Muscle Rigidity  
Nausea  
Oxygen Saturation  
Decreased  
Pupillary Reflex Impaired  
Respiratory Rate  
Decreased  
Skin Discolouration  
Speech Disorder  
Tachycardia  
Tremor

Date:02/25/00ISR Number: 3463091-0Report Type:Expedited (15-DaCompany Report #A0112179A  
Age:39 YR Gender:Female I/FU:I

Outcome	PT	Report Source
Hospitalization -	Cardiomyopathy	Health
Initial or Prolonged	Overdose	Professional

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Company Representative	Product	Role	Manufacturer	Route
ORAL			Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
			Mirtazapine (Formulation Unknown) (Mirtazapine)	SS		

Date:02/25/00ISR Number: 3463974-1Report Type:Expedited (15-DaCompany Report #A0112549A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cellulitis Oedema Peripheral Rash Erythematous	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ORAL	2 WK	Vasculitic Rash					

Date:02/25/00ISR Number: 3464045-0Report Type:Expedited (15-DaCompany Report #A0112039A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Bite Memory Impairment Pain	Health Professional Company	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE		Syncope	Representative				
PER DAY/ORAL		Tongue Disorder					

Date:02/28/00ISR Number: 3464397-1Report Type:Expedited (15-DaCompany Report #A0112382A  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Cyanosis Drug Interaction Respiratory Arrest	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

TWICE PER DAY

/ ORAL

25 MG / SEE

TEXT/ ORAL

Venlafaxine  
Hydrochloride Tablet  
(Venlafaxine  
Hydrochloride)

SS

ORAL

Date:02/28/00ISR Number: 3464843-3Report Type:Expedited (15-DaCompany Report #A0102623A

Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide Intentional Misuse	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

150 MG / SEE

TEXT / ORAL

Other

Fluoxetine  
Hydrochloride

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Freedom Of Information (FOI) Report

20 MG / SEE

TEXT / ORAL

Capsule (Fluoxetine  
Hydrochloride) SS

ORAL

37.5 MG / SEE

TEXT / ORAL

Venlafaxine  
Hydrochloride Tablet  
(Venlafaxine  
Hydrochloride) SS

ORAL

Date:02/28/00ISR Number: 3464858-5Report Type:Expedited (15-DaCompany Report #A0112392A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Foreign	Bupropion			
Required		Burning Sensation	Health	Hydrochloride Tablet			
Intervention to		Complement Factor	Professional	- Zyban (Buproipion			
Prevent Permanent		Increased		Hydrochloride)	PS		ORAL
150 MG /							
Impairment/Damage		Decreased Activity					
TWICE PER DAY							
		Difficulty In Walking					
/ ORAL							
		Drug Hypersensitivity		Mometasone Furoate	C		
		Dysphagia					
		Erythema Multiforme					
		Eyelid Oedema					
		Face Oedema					
		Fatigue					
		Feeling Cold					
		Insomnia					
		Oedema Peripheral					
		Pain					
		Peripheral Coldness					
		Pharyngolaryngeal Pain					
		Rash Pruritic					
		Sensation Of Heaviness					
		Skin Discolouration					
		Type Iv Hypersensitivity					
		Reaction					
		Urticaria					
		Vasculitis					

Date:02/28/00ISR Number: 3464859-7Report Type:Expedited (15-DaCompany Report #A0112549A  
Age:60 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Cellulitis	Foreign	Bupropion			
Initial or Prolonged	Gingivitis	Health	Hydrochloride Tablet			
	Influenza Like Illness	Professional	- Zyban (Bupropion			
	Lymphangiopathy		Hydrochloride)	PS		ORAL
150 MG / ORAL						
	Pruritus		Aspirin	C		
	Pyrexia					
	Swelling					
	Vasculitic Rash					

Date:02/28/00ISR Number: 3464860-3Report Type:Expedited (15-DaCompany Report #A0112179A  
Age:35 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Cardiac Disorder
Initial or Prolonged	Cardiomyopathy
	Chest Pain

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dyspnoea Electrocardiogram Qt Prolonged					
100 MG / SEE		Intentional Misuse Myocardial Ischaemia Syncope Toxicologic Test Abnormal	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
TEXT / ORAL		Tricuspid Valve Incompetence		Mirtazapine Tablet (Mirtazapine)	SS		ORAL
20 MG / SEE				Librium (Formulation Unknown) (Librium)	SS		
TEXT / ORAL							
SINGLE DOSE							

Date:02/28/00ISR Number: 3465209-2Report Type:Expedited (15-DaCompany Report #M0091-2000  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Cholesterol	Health	Remeron	PS		ORAL
7.5 - 22.5		Increased	Professional				
MG/DAY PO		Blood Triglycerides Increased		Wellbutrin Desoxyn	SS C		
		Liver Function Test Abnormal		Synthroid	C		
		Oedema Peripheral					

Date:03/01/00ISR Number: 3466568-7Report Type:Expedited (15-DaCompany Report #A005730  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Abdominal Distension	Health	Zoloft Tablets	PS		
Intervention to		Dissociative Identity	Professional	Bupropion	SS		
Prevent Permanent		Disorder					

Impairment/Damage      Hysterectomy  
                            Ill-Defined Disorder  
                            Nausea  
                            Tremor

Date:03/01/00ISR Number: 3469426-7Report Type:Periodic      Company Report #A0108483A  
Age:              Gender:Female      I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose              Duration						
Hospitalization - Initial or Prolonged	Palpitations	Health Professional Company Representative	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		

Date:03/01/00ISR Number: 3469429-2Report Type:Periodic      Company Report #A0104885A  
Age:19 YR      Gender:Female      I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Arthralgia Burning Sensation Erythema Multiforme Lymphadenopathy Malaise Pain

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Pruritus Pyrexia Serum Sickness	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/00ISR Number: 3469431-0Report Type:Periodic Company Report #A0105122A  
Age:63 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Tachycardia	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
Dose							
Life-Threatening							
150 MG / PER							
DAY/ ORAL							
				Conjugated Estrogens Aspirin	C C		

Date:03/01/00ISR Number: 3469434-6Report Type:Periodic Company Report #A0082834A  
Age:39 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Convulsion	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
Dose							
Other							
150 MG /							
TWICE PER DAY							
/ ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Health Professional Company Representative	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /							
TWICE PER							
DAY/ ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dyspnoea Throat Tightness Urticaria	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Weight Loss Medication	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3469442-5Report Type:Periodic  
Age:38 YR Gender:Male I/FU:F

Company Report #A0086000A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - Initial or Prolonged 150 MG / Required TWICE PER DAY Intervention to / ORAL Prevent Permanent Impairment/Damage	Abdominal Pain Anaphylactic Reaction Bronchospasm Conjunctivitis Dermatitis Dyspnoea Eyelid Oedema Face Oedema Headache Nausea Pharyngeal Oedema Pruritus Respiratory Distress Tongue Oedema Urticaria	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:03/01/00ISR Number: 3469452-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0110912A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Dyspnoea Hypersensitivity	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		

Date:03/01/00ISR Number: 3469455-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0110651A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other	Convulsion Drug Interaction	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL			Septra (Formulation			

Unknown)  
(Sulfamethoxazole  
Trimetho) SS

Date:03/01/00ISR Number: 3469458-9Report Type:Periodic Company Report #A0109719A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Akinesia Dystonia	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

ORAL

Date:03/01/00ISR Number: 3469461-9Report Type:Periodic Company Report #A0107904A  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Grand Mal Convulsion	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

150 MG /  
TWICE PER  
DAY/ ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3469464-4Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0107591A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL			Representative	Birth Control	C		

Date:03/01/00ISR Number: 3469466-8Report Type:Periodic  
Age:51 YR Gender:Male I/FU:I

Company Report #A0107632A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Convulsion Dry Mouth	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / ORAL							

Date:03/01/00ISR Number: 3469469-3Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #A0107283A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Zyban Tablet - Zyban (Bupriopion Hydrochloride)	PS		ORAL
Required Intervention to Prevent Permanent Impairment/Damage / ORAL				Ortho Cyclen	C		

Date:03/01/00ISR Number: 3547102-XReport Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0109325A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG /  
TWICE PER DAY  
/ ORAL

Anxiety  
Nervousness  
Psychomotor Hyperactivity

Consumer                      Zyban                      PS                      Glaxo Wellcome Inc                      ORAL

Date:03/01/00ISR Number: 3547108-0Report Type:Periodic                      Company Report #A0109434A  
Age:49 YR    Gender:Male                      I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Dysphonia					
		Rash Erythematous					
		Rash Pruritic					
		Throat Tightness					

Date:03/01/00ISR Number: 3547109-2Report Type:Periodic                      Company Report #A0109437A  
Age:26 YR    Gender:Female                      I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Rash Erythematous	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Rash Pruritic					

TWICE PER  
DAY/ ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3547112-2Report Type:Periodic Company Report #A0109438A  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL	1 MON	Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Ill-Defined Disorder					

Date:03/01/00ISR Number: 3547114-6Report Type:Periodic Company Report #A0109496A  
 Age:47 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER		Apathy					
DAY/ ORAL		Tremor					

Date:03/01/00ISR Number: 3547115-8Report Type:Periodic Company Report #A0109497A  
 Age:20 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Rash Generalised	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER		Throat Tightness					
DAY/ ORAL							

Date:03/01/00ISR Number: 3547117-1Report Type:Periodic Company Report #A0109498A  
 Age:73 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Balance Disorder	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

PER DAY/ ORAL  
Dysarthria  
Insomnia  
Speech Disorder

Date:03/01/00ISR Number: 3547120-1Report Type:Periodic Company Report #A0109499A  
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Withdrawal Syndrome	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER							
DAY/ ORAL							

Date:03/01/00ISR Number: 3547122-5Report Type:Periodic Company Report #A0109500A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Atopic	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Drug Hypersensitivity					
PER DAY/ ORAL		Face Oedema					
		Pruritus					
		Urticaria					

Date:03/01/00ISR Number: 3547169-9Report Type:Periodic Company Report #A0107721A  
Age: Gender:Female I/FU:I

Outcome	PT
	Bruxism
	Feeling Abnormal



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Date:03/01/00ISR Number: 3547176-6Report Type:Periodic Company Report #A0107834A							
Age:47 YR Gender:Male I/FU:I							
150 MG / TWICE PER DAY / ORAL		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Insomnia					
		Psychomotor Hyperactivity					
		Pyrexia					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Date:03/01/00ISR Number: 3547177-8Report Type:Periodic Company Report #A0107835A							
Age:29 YR Gender:Female I/FU:I							
150 MG / TWICE PER DAY / ORAL		Breast Tenderness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Menorrhagia					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Date:03/01/00ISR Number: 3547178-XReport Type:Periodic Company Report #A0107839A							
Age:25 YR Gender:Female I/FU:I							
150 MG / TWICE PER DAY / ORAL		Mydriasis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL



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Date:03/01/00ISR Number: 3547179-1Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0107871A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3547181-XReport Type:Periodic  
Age:51 YR Gender:Male I/FU:I

Company Report #A0107873A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							
DAY/ ORAL	3 MON			Lansoprazole	C		
				Methylphenidate Hcl	C		

Date:03/01/00ISR Number: 3547183-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0107929A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE							
PER DAY/ ORAL		Urticaria					

Date:03/01/00ISR Number: 3547185-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0107931A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Unevaluable Event	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							

Date:03/01/00ISR Number: 3547187-0Report Type:Periodic Company Report #A0107937A  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER							
DAY/ ORAL							

Date:03/01/00ISR Number: 3547190-0Report Type:Periodic Company Report #A0108071A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharyngeal Oedema	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL	2	WK					

Date:03/01/00ISR Number: 3547192-4Report Type:Periodic Company Report #A0108072A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Pruritus					
TWICE PER							
DAY/ ORAL							

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Date:03/01/00ISR Number: 3547195-XReport Type:Periodic  
 Age:65 YR Gender:Male I/FU:I

Company Report #A0108073A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY/ ORAL		Rash Erythematous	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3547198-5Report Type:Periodic  
 Age:37 YR Gender:Male I/FU:I

Company Report #A0108074A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Depression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3547201-2Report Type:Periodic  
 Age:47 YR Gender:Male I/FU:I

Company Report #A0108075A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY/ ORAL		Burning Sensation Myalgia Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3547203-6Report Type:Periodic  
 Age:56 YR Gender:Male I/FU:I

Company Report #A0108076A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

TWICE PER

DAY/ ORAL

Date:03/01/00ISR Number: 3547205-XReport Type:Periodic Company Report #A0108077A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Feelings Of Worthlessness					
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3547209-7Report Type:Periodic Company Report #A0108079A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Feelings Of Worthlessness					
TWICE PER DAY							

/ ORAL

Date:03/01/00ISR Number: 3547216-4Report Type:Periodic Company Report #A0107030A

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							

/ ORAL



150 MG /  
TWICE PER  
DAY/ ORAL  
Pruritus  
Consumer  
Zyban  
PS  
Glaxo Wellcome Inc  
ORAL

Date:03/01/00ISR Number: 3547223-1Report Type:Periodic Company Report #A0107040A  
Age:74 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL		Agitation Hyperhidrosis Insomnia Pollakiuria Vomiting	Consumer	Zyban   Atorvastatin Multivitamin	PS   C C	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3547225-5Report Type:Periodic Company Report #A0107043A  
Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Burning Sensation Pruritus Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

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Date:03/01/00ISR Number: 3547226-7Report Type:Periodic Company Report #A0107065A  
 Age:36 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER		Irritability					
DAY/ ORAL	2 WK	Weight Increased					

Date:03/01/00ISR Number: 3547228-0Report Type:Periodic Company Report #A0107067A  
 Age:64 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER		Dysgeusia					
DAY/ ORAL		Tremor					

Date:03/01/00ISR Number: 3547229-2Report Type:Periodic Company Report #A0107068A  
 Age:57 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /	ORAL	Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3547230-9Report Type:Periodic Company Report #A0107069A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Drug Effect Decreased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL			Other				

7 MG

Nicotine  
(Formulation  
Unknown)

SS

Date:03/01/00ISR Number: 3547233-4Report Type:Periodic Company Report #A0107093A  
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nightmare	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Tremor					
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3547234-6Report Type:Periodic Company Report #A0107112A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL	6 WK	Emotional Disorder		Nitrous Oxide Gas			
		Panic Reaction		(Nitrous Oxide)	SS		

Date:03/01/00ISR Number: 3547235-8Report Type:Periodic Company Report #A0107159A  
Age:41 YR Gender:Male I/FU:I

Outcome	PT
	Agitation
	Blood Pressure Increased
	Discomfort
	Dizziness
	Excitability



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		Flushing Nervousness Tachycardia	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER							
DAY/ ORAL							

Date:03/01/00ISR Number: 3547236-XReport Type:Periodic Company Report #A0107188A  
 Age:29 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Accommodation Disorder	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Dose		Dry Mouth					
150 MG/ TWICE		Tachycardia					
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3547237-1Report Type:Periodic Company Report #A0107200A  
 Age:39 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Dose		Feeling Abnormal					
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3547239-5Report Type:Periodic Company Report #A0107202A  
 Age:38 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Dose		Dysgeusia					
150 MG /							
TWICE PER DAY							

/ ORAL  
Insomnia  
Vision Blurred

Date:03/01/00ISR Number: 3547243-7Report Type:Periodic Company Report #A0107203A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cough	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Dyspnoea					
TWICE PER DAY		Pruritus					
/ ORAL		Urticaria					

Date:03/01/00ISR Number: 3547246-2Report Type:Periodic Company Report #A0107204A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/00ISR Number: 3547249-8Report Type:Periodic Company Report #A0107205A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hair Texture Abnormal	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL 1 YR		Nail Disorder					



150 MG/ TWICE  
PER DAY/ ORAL  
Dermatitis  
Nausea  
Health  
Professional  
Zyban  
PS  
Glaxo Wellcome Inc  
ORAL

Date:03/01/00ISR Number: 3547300-5Report Type:Periodic Company Report #A0109601A  
Age:27 YR Gender:Female I/FU:I

Outcome PT Report Source Product Role Manufacturer Route  
Dose Duration  
150 MG/ PER  
DAY/ ORAL  
Dermatitis Consumer Zyban PS Glaxo Wellcome Inc ORAL

Date:03/01/00ISR Number: 3547303-0Report Type:Periodic Company Report #A0109604A  
Age: Gender:Female I/FU:I

Outcome PT Report Source Product Role Manufacturer Route  
Dose Duration  
150 MG/ TWICE  
PER DAY/ ORAL  
Flushing Consumer Zyban PS Glaxo Wellcome Inc ORAL

Date:03/01/00ISR Number: 3547308-XReport Type:Periodic Company Report #A0109611A  
Age:55 YR Gender:Female I/FU:I

Outcome PT Report Source Product Role Manufacturer Route  
Dose Duration  
150 MG/ TWICE  
PER DAY/ ORAL  
Alopecia Consumer Zyban PS Glaxo Wellcome Inc ORAL  
Dry Mouth  
Insomnia

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Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3547310-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0110307A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ UNK/ORAL		Aggression Personality Change  Psychiatric Symptom	Health Professional  Company Representative	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3547312-1Report Type:Periodic  
Age:41 YR Gender:Male I/FU:I

Company Report #A0110328A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG, / TWICE PER DAY/ ORAL		Chest Pain Dyspnoea  Insomnia  Tachycardia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3547313-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0109633A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ UNK/ORAL		Drug Ineffective  Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3547315-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0110329A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective	Consumer	Bupropion			

Urticaria Hydrochloride PS Glaxo Wellcome Inc ORAL  
150 MG / UNK  
/ ORAL  
Date:03/01/00ISR Number: 3547316-9Report Type:Periodic Company Report #A0110331A  
Age: Gender:Female I/FU:I  
Outcome PT Report Source Product Role Manufacturer Route  
Dose Duration Drug Ineffective Consumer Bupropion  
Irritability Hydrochloride PS Glaxo Wellcome Inc ORAL  
150 MG /UNK /  
ORAL

Date:03/01/00ISR Number: 3547317-0Report Type:Periodic Company Report #A0110332A  
Age: Gender:Male I/FU:I  
Outcome PT Report Source Product Role Manufacturer Route  
Dose Duration Drug Ineffective Consumer Bupropion  
Irritability Hydrochloride PS Glaxo Wellcome Inc ORAL  
150 MG / UNK  
/ ORAL

Date:03/01/00ISR Number: 3547319-4Report Type:Periodic Company Report #A0110354A  
Age:55 YR Gender:Female I/FU:U  
Outcome PT  
Dysphagia  
Dyspnoea  
Pharyngeal Oedema

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		Rash Erythematous Rash Pruritic				
Dose	Duration		Report Source	Product	Role	Manufacturer
			Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc
150 MG /TWICE						ORAL
PER DAY /						
ORAL						

Date:03/01/00ISR Number: 3547321-2Report Type:Periodic Company Report #A0110371A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Consumer	Zyban Tablet	PS	Glaxo Wellcome Inc	ORAL
ORAL		Feeling Abnormal Flushing Gastrointestinal Disorder Pyrexia Urticaria					

Date:03/01/00ISR Number: 3547323-6Report Type:Periodic Company Report #A0110377A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE							
PER DAY /							
ORAL							

Date:03/01/00ISR Number: 3547326-1Report Type:Periodic Company Report #A0110378A  
Age:43 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anaphylactic Reaction Dermatitis	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ORAL							

Date:03/01/00ISR Number: 3547328-5Report Type:Periodic Company Report #A0110379A  
 Age: Gender:Unknown I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dissociation Feeling Abnormal	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / UNK		Nausea					
/ ORAL	3 DAY						

Date:03/01/00ISR Number: 3547330-3Report Type:Periodic Company Report #A0110399A  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / UNK							
/ ORAL	7 WK						



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Date:03/01/00ISR Number: 3547331-5Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #A0110419A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache Visual Disturbance	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/00ISR Number: 3547333-9Report Type:Periodic  
 Age:55 YR Gender:Male I/FU:I

Company Report #A0110562A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150MG / TWICE							
PER DAY /							
ORAL							

Date:03/01/00ISR Number: 3547335-2Report Type:Periodic  
 Age:47 YR Gender:Male I/FU:I

Company Report #A0110563A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Irritability	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/00ISR Number: 3547336-4Report Type:Periodic  
 Age:30 YR Gender:Female I/FU:I

Company Report #A0110564A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dry Mouth	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							

Date:03/01/00ISR Number: 3547338-8Report Type:Periodic Company Report #A0110565A  
 Age:41 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Accommodation Disorder Agitation	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Disturbance In Attention							
Dizziness							
Mental Impairment							
Nausea							
Tremor							
TWICE PER DAY/ ORAL							

Date:03/01/00ISR Number: 3547340-6Report Type:Periodic Company Report #A0110566A  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / UNK		Abdominal Discomfort	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
/ ORAL							



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Condition Aggravated	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL		Drug Dependence					
Date:03/01/00ISR Number: 3547366-2Report Type:Periodic Company Report #A0109635A							
Age:63 YR Gender:Male I/FU:I							
150 MG/ TWICE		Condition Aggravated	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL	1 WK	Micturition Urgency					
		Pollakiuria					
Date:03/01/00ISR Number: 3547368-6Report Type:Periodic Company Report #A0109636A							
Age:53 YR Gender:Male I/FU:I							
150 MG/ TWICE		Apathy	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL		Insomnia					
Date:03/01/00ISR Number: 3547371-6Report Type:Periodic Company Report #A0109637A							
Age:35 YR Gender:Female I/FU:I							
UNK/ SEE		Diarrhoea	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
TEXT/ ORAL		Nausea	Professional				

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Date:03/01/00ISR Number: 3547373-XReport Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #A0109640A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3547376-5Report Type:Periodic  
Age:60 YR Gender:Male I/FU:I

Company Report #A0109641A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ UNK/ ORAL		Rash Erythematous Rash Pruritic	Health Professional Company Representative	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3547379-0Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0109690A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL		Face Oedema Mouth Ulceration Oedema Peripheral Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3547383-2Report Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #A0109721A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Galactorrhoea	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL

Hypertrophy Breast

Professional

PER DAY/ ORAL

Date:03/01/00ISR Number: 3547386-8Report Type:Periodic Company Report #A0109722A

Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							

PER DAY/ ORAL

Date:03/01/00ISR Number: 3547392-3Report Type:Periodic Company Report #A0109723A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Pruritic	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/ UNK/							

ORAL

Date:03/01/00ISR Number: 3547402-3Report Type:Periodic Company Report #A0109725A

Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Paraesthesia					

PER DAY/ ORAL

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150 MG /  
Dizziness Consumer Zyban PS Glaxo Wellcome Inc ORAL  
TWICE PER DAY  
/ ORAL

Date:03/01/00ISR Number: 3547431-XReport Type:Periodic Company Report #A0108276A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Burning Sensation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Chills					
TWICE PER DAY		Dizziness					
/ ORAL		Dysphagia					
		Headache					
		Sensation Of Pressure					
		Tremor					
		Vision Blurred					

Date:03/01/00ISR Number: 3547433-3Report Type:Periodic Company Report #A0108290A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depressed Mood	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3547435-7Report Type:Periodic  
 Age:60 YR Gender:Male I/FU:I

Company Report #A0108293A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Irritability	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER DAY / ORAL							

Date:03/01/00ISR Number: 3547437-0Report Type:Periodic  
 Age:61 YR Gender:Female I/FU:I

Company Report #A0108303A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eye Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY / ORAL							

Date:03/01/00ISR Number: 3547438-2Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0108485A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Malaise	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER DAY/ ORAL							
		Sedation		Sertraline Hydrochloride Diazepam	C C		

Date:03/01/00ISR Number: 3547439-4Report Type:Periodic  
 Age:45 YR Gender:Male I/FU:I

Company Report #A0108622A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Pollakiuria Consumer Zyban PS Glaxo Wellcome Inc ORAL  
150 MG /  
TWICE PER DAY  
/ ORAL

Date:03/01/00ISR Number: 3547442-4Report Type:Periodic Company Report #A0108623A  
Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Depressed Mood					
TWICE PER DAY		Nervousness					
/ ORAL							

Date:03/01/00ISR Number: 3547445-XReport Type:Periodic Company Report #A0108624A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Nervousness					
TWICE PER DAY							
/ ORAL							

Date:03/01/00ISR Number: 3547447-3Report Type:Periodic Company Report #A0108625A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Headache					
DAY / ORAL		Insomnia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3547461-8Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #A0108626A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							

Date:03/01/00ISR Number: 3547466-7Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0108627A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
Tremor							

Date:03/01/00ISR Number: 3547469-2Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #A0108628A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
Tachycardia							

Date:03/01/00ISR Number: 3547472-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0108629A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							

/ ORAL

Date:03/01/00ISR Number: 3547475-8Report Type:Periodic Company Report #A0108630A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Nausea					
TWICE PER DAY		Urticaria					

/ ORAL

Date:03/01/00ISR Number: 3547477-1Report Type:Periodic Company Report #A0108631A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Insomnia					
TWICE PER DAY		Nervousness					
/ ORAL		Tremor					

Date:03/01/00ISR Number: 3547483-7Report Type:Periodic Company Report #A0108632A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL	2	WK					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3547486-2Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0108633A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL	WK		Professional				

Date:03/01/00ISR Number: 3548822-3Report Type:Periodic  
 Age:57 YR Gender:Female I/FU:F

Company Report #A0102578A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Drug Withdrawal Syndrome	Professional				
PER DAY ORAL		Tremor		Thyroxine Sodium	C		

Date:03/01/00ISR Number: 3548823-5Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:F

Company Report #A0102897A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
Required		Dizziness	Professional				
150 MG UNK		Erythema Multiforme	Company				
Intervention to		Fatigue	Representative				
ORAL		Pharyngolaryngeal Pain					
Prevent Permanent		Pyrexia					
Impairment/Damage							

Date:03/01/00ISR Number: 3548825-9Report Type:Periodic  
 Age:50 YR Gender:Female I/FU:F

Company Report #A0103098A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG TWICE							

PER DAY ORAL	Pruritus	Professional				
	Urticaria		Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	SS		ORAL
150 MG TWICE						
PER DAY ORAL			Lisinopril Hydrochlorothiazide	C C		

Date:03/01/00ISR Number: 3548826-0Report Type:Periodic Company Report #A0103106A  
 Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Nausea	Professional				
PER DAY ORAL		Tremor					

Date:03/01/00ISR Number: 3548831-4Report Type:Periodic Company Report #A0103108A  
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Spasms	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG TWICE		Pain In Extremity	Professional				
PER DAY ORAL		Paraesthesia Peripheral Coldness Pulse Pressure Decreased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3548832-6Report Type:Periodic  
Age:45 YR Gender:Female I/FU:F

Company Report #A0103165A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Spasms	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Paraesthesia	Professional				
PER DAY ORAL	1	MON					

Date:03/01/00ISR Number: 3548833-8Report Type:Periodic  
Age:39 YR Gender:Female I/FU:F

Company Report #A0103325A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blister	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Oedema	Professional				
DAY ORAL		Pruritus					

Date:03/01/00ISR Number: 3548834-XReport Type:Periodic  
Age:61 YR Gender:Male I/FU:F

Company Report #A0103421A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Affective Disorder	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Depression	Professional				
PER DAY ORAL		Dry Mouth					
		Dysarthria					
		Lip Dry					
		Mental Impairment					
		Nervousness					
		Tremor					

Date:03/01/00ISR Number: 3548835-1Report Type:Periodic  
Age:32 YR Gender:Female I/FU:F

Company Report #A0103653A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		African Trypanosomiasis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Aggression	Professional				
PER DAY ORAL		Depression					
		Lethargy					
		Mood Altered					
		Pain					
		Sedation					
		Weight Increased					

Date:03/01/00ISR Number: 3548837-5Report Type:Periodic Company Report #A0104078A  
Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Dissociation	Professional				
DAY ORAL		Irritability					
		Major Depression					
		Nausea					
		Pain					
		Pyrexia					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3548854-5Report Type:Periodic  
Age:53 YR Gender:Male I/FU:F

Company Report #A0104325A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperglycaemia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG TWICE			Professional				
PER DAY ORAL				Insulin	C		

Date:03/01/00ISR Number: 3548855-7Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #A0104459A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG TWICE A		Dry Mouth	Professional				
DAY ORAL		Urticaria		Esterified Estrogens	C		
				Medroxyprogesterone			
				Ace.	C		

Date:03/01/00ISR Number: 3548856-9Report Type:Periodic  
Age:39 YR Gender:Female I/FU:F

Company Report #A0104485A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG TWICE		Nausea					
PER DAY ORAL							

Date:03/01/00ISR Number: 3548858-2Report Type:Periodic  
Age:24 YR Gender:Female I/FU:F

Company Report #A0104493A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Conversion Disorder	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG PER DAY							

ORAL		Crying	Professional				
		Delusion		Medroxyprogesterone			
		Major Depression		Ace		C	
		Mental Impairment					
		Panic Disorder Without					
		Agoraphobia					

Date:03/01/00ISR Number: 3548861-2Report Type:Periodic Company Report #A0104588A  
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG PER		Dermatitis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY ORAL		Dry Skin	Professional				
		Erythema					
		Immune System Disorder					
		Pruritus					
		Upper Respiratory Tract					
		Infection					
		Urticaria					

Date:03/01/00ISR Number: 3548864-8Report Type:Periodic Company Report #A0081321A  
 Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
ORAL		Flatulence	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TRANSDERMAL		Insomnia	Other	Nicotine Patch	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3548887-9Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #A0085269A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Dose	150 MG ORAL	Convulsion	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Intervention to Prevent Permanent Impairment/Damage	PER DAY	Erythema					
		Hypersensitivity		Aspirin	C		
		Lacrimation Increased		Digoxin	C		
		Throat Tightness		Semisodium Valproate	C		
				Vitamin	C		
				Frusemide	C		
				Loracarbef	C		

Date:03/01/00ISR Number: 3548890-9Report Type:Periodic  
Age:53 YR Gender:Male I/FU:I

Company Report #A0096361A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Dose	150MG PER DAY	Diarrhoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
	ORAL	Flatulence					
		Headache		Lisinopril	C		
		Irritable Bowel Syndrome		Int/Long-Acting			
		Pyrexia		Insulin	C		
				Insulin Lispro	C		

Date:03/01/00ISR Number: 3548891-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0096591A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Dose	150 MG TWICE	Depression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
	PER DAY ORAL	Drug Withdrawal Syndrome					

Date:03/01/00ISR Number: 3548893-4Report Type:Periodic Company Report #A0096594A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Pruritus					
DAY ORAL							

Date:03/01/00ISR Number: 3548918-6Report Type:Periodic Company Report #A0105084A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /ORAL			Professional Company Representative				

Date:03/01/00ISR Number: 3548923-XReport Type:Periodic Company Report #A0105085A  
Age:44 YR Gender:Female I/FU:I

Outcome	PT
	Abdominal Pain
	Back Pain
	Chest Pain
	Condition Aggravated
	Depression
	Diarrhoea

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dry Mouth Increased Appetite Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Nausea					
/ ORAL		Pyrexia					
		Sinus Headache					
		Therapeutic Response Unexpected					

Date:03/01/00ISR Number: 3548926-5Report Type:Periodic Company Report #A0105123A  
 Age:50 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /			Tremor	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY				Other				
/ ORAL					Nicotine Patch (Nicotine)	SS		
TRANSDERMAL	21 MG/							
TRANSDERMAL								

Date:03/01/00ISR Number: 3548929-0Report Type:Periodic Company Report #A0105124A  
 Age:47 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /			Abdominal Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY			Anxiety					
/ ORAL			Disturbance In Attention					
			Tachycardia					

Date:03/01/00ISR Number: 3548931-9Report Type:Periodic Company Report #A0105307A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Decreased Appetite					
TWICE PER DAY		Diarrhoea					
/ORAL		Hyperhidrosis		Hyoscyamine Sulphate	C		
		Nausea		Mesalazine	C		
		Pollakiuria					
		Tremor					
		Viral Infection					

Date:03/01/00ISR Number: 3548933-2Report Type:Periodic Company Report #A0105374A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Nervousness					
TWICE PER		Tension					
DAY/ ORAL		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3548935-6Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #A0105375A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Dizziness Sedation	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Company Representative	Tamoxifen	C		

Date:03/01/00ISR Number: 3548938-1Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0105377A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Dry Mouth Dyspepsia Headache Insomnia Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3548941-1Report Type:Periodic  
Age:59 YR Gender:Female I/FU:I

Company Report #A0105390A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY /ORAL		Dizziness Face Oedema Flushing Hyperhidrosis Hypersensitivity Oedema Oedema Peripheral Pruritus Urticaria Weight Increased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3548944-7Report Type:Periodic Company Report #A0105392A  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia Headache	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /	ORAL						

Date:03/01/00ISR Number: 3548946-0Report Type:Periodic Company Report #A0105393A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper Decreased Activity	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 GM /							
TWICE PER DAY							
/ ORAL	8 DAY	Diarrhoea Headache					

Date:03/01/00ISR Number: 3548950-2Report Type:Periodic Company Report #A0105394A  
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity Pruritus Swelling	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER							
DAY/ ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3548953-8Report Type:Periodic Company Report #A0105396A  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Nausea					
TWICE PER							
DAY/ ORAL							

Date:03/01/00ISR Number: 3548955-1Report Type:Periodic Company Report #A0105397A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL		Malaise					

Date:03/01/00ISR Number: 3548957-5Report Type:Periodic Company Report #A0105404A  
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER							
DAY/ ORAL							

Date:03/01/00ISR Number: 3548960-5Report Type:Periodic Company Report #A0105405A  
 Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							

Insomnia

DAY / ORAL

Date:03/01/00ISR Number: 3548963-0Report Type:Periodic Company Report #A0105506A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mood Altered	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							

Date:03/01/00ISR Number: 3548965-4Report Type:Periodic Company Report #A0105507A

Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hunger	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL	40 DAY	Weight Increased	Other	Nicotine Polacrilex	C		

Date:03/01/00ISR Number: 3548969-1Report Type:Periodic Company Report #A0105510A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Unevaluable Event	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							

Date:03/01/00ISR Number: 3548972-1Report Type:Periodic Company Report #A0105511A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Libido Decreased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3548975-7Report Type:Periodic  
 Age:25 YR Gender:Male I/FU:I

Company Report #A0110730A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Syncope Visual Disturbance	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ AS DIRECTED/ ORAL 4 DAY							

Date:03/01/00ISR Number: 3548979-4Report Type:Periodic  
 Age:63 YR Gender:Male I/FU:I

Company Report #A0110842A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Residue	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE PER DAY/ ORAL							

Date:03/01/00ISR Number: 3548982-4Report Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0110850A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Abnormal Drug Ineffective	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE PER DAY/ORAL							

Date:03/01/00ISR Number: 3548985-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0110991A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Health	Bupropion			

150 MG ORAL	Drug Interaction	Professional	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
	Dysphonia	Company	Thyroxine Sodium			
	Tremor	Representative	(Levothyroxine Sodium)	SS		
15 MG			Pravastatin Sodium			
			(Pravastatin Sodium)	SS		
10 MG						

Date:03/01/00ISR Number: 3548989-7Report Type:Periodic Company Report #A0111092A  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
PER DAY /ORAL							

Date:03/01/00ISR Number: 3548992-7Report Type:Periodic Company Report #A0055184A  
 Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Feeling Abnormal	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER		Insomnia					
DAY/ORAL		Tremor		Cisapride	C		
				Loratadine	C		
				Herbal Medication	C		
				Glucophage	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3548994-0Report Type:Periodic  
Age:61 YR Gender:Male I/FU:F

Company Report #A0058731A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE		Drug Ineffective Nervousness	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ORAL	2 MON	Tremor					

Date:03/01/00ISR Number: 3548997-6Report Type:Periodic  
Age:63 YR Gender:Male I/FU:F

Company Report #A0066125A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE		Depression Drug Ineffective	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ORAL		Headache					
150 MG/TWICE		Nervousness		Zyban Tablet-Zyban (Bupropion Hydrochloride)	SS		ORAL
PER DAY/ORAL	2 WK			Etodolac Cardiovascular Medication Pain Medication Sleeping Medication	C C C C		

Date:03/01/00ISR Number: 3549000-4Report Type:Periodic  
Age:54 YR Gender:Female I/FU:F

Company Report #A0080417A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE		Depression Insomnia	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL		Restlessness					

Suicidal Ideation

Hydroxyzine  
Hydrochloride

C

Date:03/01/00ISR Number: 3549002-8Report Type:Periodic  
Age:30 YR Gender:Female I/FU:F

Company Report #A0087975A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash Papular	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Hydrocodone

C

Date:03/01/00ISR Number: 3549004-1Report Type:Periodic  
Age:33 YR Gender:Female I/FU:F

Company Report #A0096164A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia Irritability	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							
		Mood Altered Nightmare					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3549007-7Report Type:Periodic  
Age:30 YR Gender:Female I/FU:F

Company Report #A0099338A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL		Crying Derealisation  Emotional Disorder Feeling Abnormal Hyperhidrosis Insomnia Nausea Paranoia Psychotic Disorder Tremor	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3549010-7Report Type:Periodic  
Age:58 YR Gender:Female I/FU:F

Company Report #A0099558A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Decreased Appetite Dysgeusia  Insomnia  Irritability Parosmia	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3549011-9Report Type:Periodic  
Age:46 YR Gender:Female I/FU:F

Company Report #A0100148A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Dermatitis Pruritus  Swelling  Urticaria	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Constipation	Health	Bupropion			
		Hair Disorder	Professional	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
PER DAY/ORAL		Insomnia					
				Nicotine	C		
				Benazepril			
				Hydrochloride	C		
				Nifedipine	C		
				Thyroid Medication	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation	Consumer	Bupropion			
		Feeling Abnormal		Hydrochloride	PS	Glaxo Wellcome Inc	
OPHTHALMIC	150 MG/ TWICE	Ill-Defined Disorder					
PER DAY/ORAL		Insomnia					
		Malaise					
		Nausea					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3549016-8Report Type:Periodic  
 Age:39 YR Gender:Male I/FU:F

Company Report #A0101851A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/PER DAY/ORAL		Agitation Disturbance In Attention Hyperhidrosis Insomnia Nervousness	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3549017-XReport Type:Periodic  
 Age:60 YR Gender:Female I/FU:F

Company Report #A0102168A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Cough Hyperhidrosis	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3549019-3Report Type:Periodic  
 Age:33 YR Gender:Male I/FU:F

Company Report #A0102172A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ ORAL		Dizziness Emotional Disorder Flushing Nausea Personality Change Psychomotor Hyperactivity	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3549021-1Report Type:Periodic  
 Age:46 YR Gender:Male I/FU:F

Company Report #A0102506A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypertension Neck Pain	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER							
DAY/ORAL				Theophylline Salbutamol Sulphate	C C		

Date:03/01/00ISR Number: 3549120-4Report Type:Periodic Company Report #A0109768A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tachycardia Tremor	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							
DAY/ ORAL							

Date:03/01/00ISR Number: 3549121-6Report Type:Periodic Company Report #A0109910A  
 Age:67 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective Insomnia	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG, ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3549122-8Report Type:Periodic  
Age:53 YR Gender:Male I/FU:I

Company Report #A0109913A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Benign Prostatic Hyperplasia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE PER DAY/ ORAL							

Date:03/01/00ISR Number: 3549123-XReport Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #A0109919A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER DAY ORAL							

Date:03/01/00ISR Number: 3549124-1Report Type:Periodic  
Age:59 YR Gender:Male I/FU:I

Company Report #A0109921A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema Hypersensitivity	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE PER DAY							

Date:03/01/00ISR Number: 3549125-3Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #A0109922A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							

TWICE PER DAY

ORAL

Date:03/01/00ISR Number: 3549126-5Report Type:Periodic Company Report #A0109924A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction Premature Ejaculation	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL							

Date:03/01/00ISR Number: 3549127-7Report Type:Periodic Company Report #A0110045A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL							

Date:03/01/00ISR Number: 3549129-0Report Type:Periodic Company Report #A0110048A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3549130-7Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0110051A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3549131-9Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0110057A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3549132-0Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0110059A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia Tinnitus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							
DAY/ ORAL							

Date:03/01/00ISR Number: 3549133-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0110061A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL							

Malaise  
Tremor

Date:03/01/00ISR Number: 3549135-6Report Type:Periodic Company Report #A0110113A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3549136-8Report Type:Periodic Company Report #A0110114A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity Rash Generalised	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							
DAY/ ORAL							

Date:03/01/00ISR Number: 3549137-XReport Type:Periodic Company Report #A0110116A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Effect Decreased	Consumer Other	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3549139-3Report Type:Periodic  
Age:22 YR Gender:Male I/FU:I

Company Report #A0110117A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth Dyspnoea	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Hyperhidrosis					
PER DAY ORAL		Insomnia Lethargy Psychomotor Hyperactivity Tachycardia					

Date:03/01/00ISR Number: 3549140-XReport Type:Periodic  
Age:41 YR Gender:Male I/FU:I

Company Report #A0110118A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Behaviour	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							
DAY/ ORAL							

Date:03/01/00ISR Number: 3549141-1Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0110119A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Cough	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Dyspnoea					
DAY/ORAL		Nausea					

Date:03/01/00ISR Number: 3549142-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0110120A

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Suicidal Ideation	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL			Company Representative				

Date:03/01/00ISR Number: 3549297-0Report Type:Periodic Company Report #A0109065A  
 Age:23 YR Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Irritability Palpitations	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3549298-2Report Type:Periodic Company Report #A0109066A  
 Age:42 YR Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Headache Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3549299-4Report Type:Periodic  
Age:27 YR Gender:Female I/FU:I

Company Report #A0109067A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Emotional Disorder	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3549302-1Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0109068A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3549304-5Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0109110A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Panic Attack	Consumer Other	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL							

Date:03/01/00ISR Number: 3549306-9Report Type:Periodic  
Age:65 YR Gender:Male I/FU:I

Company Report #A0109131A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3549309-4Report Type:Periodic Company Report #A0109200A  
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3549311-2Report Type:Periodic Company Report #A0109201A  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							
DAY/ ORAL							

Date:03/01/00ISR Number: 3549313-6Report Type:Periodic Company Report #A0109202A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth Rash Pruritic	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL 2 WK							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3549314-8Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #A0109203A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Confusional State Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL		Nausea Tremor					

Date:03/01/00ISR Number: 3549316-1Report Type:Periodic  
Age:22 YR Gender:Female I/FU:I

Company Report #A0109204A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Back Pain Headache	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL		Insomnia					

Date:03/01/00ISR Number: 3549319-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0109205A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Dysgeusia Muscle Spasms	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL		Nausea					

Date:03/01/00ISR Number: 3549323-9Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #A0109216A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness	Consumer	Bupropion			

150 MG/ TWICE	Headache		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
	Nausea					
PER DAY/ ORAL	Neck Pain					

Date:03/01/00ISR Number: 3549325-2Report Type:Periodic Company Report #A0109292A  
 Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness	Consumer	Bupropion			
		Feeling Abnormal		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Vertigo					
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3549327-6Report Type:Periodic Company Report #A0109319A  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression	Consumer	Bupropion			
				Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE				Sertraline			
PER DAY/ORAL				Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3549329-XReport Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #A0109320A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3549331-8Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0109321A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysarthria Psychomotor Hyperactivity	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							
DAY/ ORAL							
Tachycardia							

Date:03/01/00ISR Number: 3549334-3Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #A0109322A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							
Vision Blurred							

Date:03/01/00ISR Number: 3549337-9Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #A0109323A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Hallucination, Auditory	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							

PER DAY/ ORAL  
Nausea  
Tremor

Date:03/01/00ISR Number: 3549340-9Report Type:Periodic Company Report #A0109324A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Dyspnoea	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Fatigue					
PER DAY/ ORAL		Feeling Abnormal Insomnia					

Date:03/01/00ISR Number: 3549362-8Report Type:Periodic Company Report #A0110570A  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3549365-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0110571A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disturbance In Attention Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Tremor					
TWICE PER							
DAY/ ORAL							

Date:03/01/00ISR Number: 3549369-0Report Type:Periodic  
 Age:42 YR Gender:Female I/FU:I

Company Report #A0110572A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying Dry Mouth	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Dyspepsia					
PER DAY/ ORAL		Mood Altered Psychomotor Hyperactivity Tachycardia		Diazepam Nabumetone Darvocet-N	C C C		

Date:03/01/00ISR Number: 3549370-7Report Type:Periodic  
 Age:45 YR Gender:Male I/FU:I

Company Report #A0110573A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							
DAY/ ORAL							

Date:03/01/00ISR Number: 3549372-0Report Type:Periodic  
 Age:36 YR Gender:Female I/FU:I

Company Report #A0110577A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cataract Subcapsular	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL	MON						

Date:03/01/00ISR Number: 3549374-4Report Type:Periodic Company Report #A0110654A  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Restlessness	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER DAY/ ORAL		Tachycardia					

Date:03/01/00ISR Number: 3549376-8Report Type:Periodic Company Report #A0110656A  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE PER DAY/ ORAL		Urticaria					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3549378-1Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #A0110658A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Disturbance In Attention Flat Affect Memory Impairment Sedation	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3549380-XReport Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #A0110675A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL		Dizziness Dry Mouth Irritability Lethargy	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3549381-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0110692A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ ORAL		Face Oedema Hypersensitivity Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3549383-5Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0110697A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation	Consumer	Bupropion			

150 MG/ ORAL      Constipation      Hydrochloride      PS      Glaxo Wellcome Inc      ORAL  
Dry Mouth

Date:03/01/00    ISR Number: 3549385-9    Report Type:Periodic      Company Report #A0110700A  
Age:53 YR    Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL				Nicotine Patch (Nicotine)	SS		
TRANSDERMAL	TRANSDERMAL						

Date:03/01/00    ISR Number: 3549387-2    Report Type:Periodic      Company Report #A0110701A  
Age:43 YR    Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Formication Insomnia	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							
DAY/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3549389-6Report Type:Periodic  
 Age:57 YR Gender:Female I/FU:I

Company Report #A0110704A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Chest Discomfort Chest Pain  Dizziness  Dry Mouth Dry Throat Dysphagia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3549391-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0110707A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ ORAL		Hyperventilation Paraesthesia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3549392-6Report Type:Periodic  
 Age:47 YR Gender:Male I/FU:I

Company Report #A0110710A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Constipation Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3549393-8Report Type:Periodic  
 Age:54 YR Gender:Female I/FU:I

Company Report #A0110711A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash Pruritic	Consumer	Bupropion			

Urticaria Hydrochloride PS Glaxo Wellcome Inc ORAL  
150 MG/ TWICE  
PER DAY/ ORAL

Date:03/01/00ISR Number: 3549395-1Report Type:Periodic Company Report #A0110712A  
Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

150 MG/ PER  
DAY/ ORAL

Date:03/01/00ISR Number: 3549396-3Report Type:Periodic Company Report #A0110718A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying Depression	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

150 MG/ ORAL

Date:03/01/00ISR Number: 3549398-7Report Type:Periodic Company Report #A0110719A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

150 MG/ TWICE  
PER DAY/ ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3549477-4Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #A0106915A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY), ORAL							

Date:03/01/00ISR Number: 3549480-4Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #A0106957A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL							

Date:03/01/00ISR Number: 3549482-8Report Type:Periodic  
Age:69 YR Gender:Female I/FU:I

Company Report #A0106958A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL							

Date:03/01/00ISR Number: 3549485-3Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #A0106959A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Consumer	Bupropion			

150 MG (TWICE  
PER DAY) ORAL  
Insomnia  
Nightmare  
Hydrochloride  
PS  
Glaxo Wellcome Inc  
ORAL

Date:03/01/00ISR Number: 3549488-9Report Type:Periodic Company Report #A0106960A  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

150 MG (TWICE  
PER DAY) ORAL

Date:03/01/00ISR Number: 3549493-2Report Type:Periodic Company Report #A0106961A  
Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Excitability Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

150 MG (TWICE  
PER DAY) ORAL

Theraflu  
C

Date:03/01/00ISR Number: 3549494-4Report Type:Periodic Company Report #A0106962A  
Age:49 YR Gender:Male I/FU:I

Outcome	PT
	Dermatitis Hypertension

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Tachycardia Tension Tremor	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL				Nicotine	C		

Date:03/01/00ISR Number: 3549495-6Report Type:Periodic Company Report #A0106963A  
Age:49 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome Dose	Duration	Blister Dermatitis Exfoliative	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (PER DAY) ORAL		Oedema		Methotrexate	C		
		Pain Photosensitivity Reaction Urinary Retention		Verapamil	C		

Date:03/01/00ISR Number: 3549497-XReport Type:Periodic Company Report #A0106964A  
Age:33 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome Dose	Duration	Dry Mouth Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (PER DAY) ORAL							

Date:03/01/00ISR Number: 3549502-0Report Type:Periodic Company Report #A0106965A  
Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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150 MG (TWICE  
PER DAY) ORAL

Alopecia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Dermatitis					
Pruritus					

Date:03/01/00ISR Number: 3549505-6Report Type:Periodic Company Report #A0106966A  
Age:29 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL		Anxiety Cognitive Disorder	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Emotional Disorder Fatigue Poverty Of Speech					

Date:03/01/00ISR Number: 3549516-0Report Type:Periodic Company Report #A0106967A  
Age:39 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL		Agitation Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Skin Odour Abnormal					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3549519-6Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #A0106968A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG (TWICE PER DAY) ORAL		Hypoaesthesia Mental Disorder	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3549520-2Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0106969A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG (PER DAY), ORAL		Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3549522-6Report Type:Periodic  
Age:79 YR Gender:Female I/FU:I

Company Report #A0106970A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG (TWICE PER DAY), ORAL		Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3549523-8Report Type:Periodic  
Age:68 YR Gender:Female I/FU:I

Company Report #A0106971A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache	Consumer	Bupropion			

150 MG (TWICE  
PER DAY) ORAL

Date:03/01/00ISR Number: 3549525-1Report Type:Periodic Company Report #A0106972A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort Dizziness	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (PER DAY) ORAL		Panic Disorder					

Date:03/01/00ISR Number: 3549530-5Report Type:Periodic Company Report #A0106973A  
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anger Communication Disorder	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL		Confusional State Crying Emotional Disorder Fear Hyperhidrosis Musculoskeletal Stiffness Nausea Palpitations Restlessness Thinking Abnormal					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3549532-9Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0106974A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL		Nausea	Consumer Other	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
14 MG PATCH				Nicotine Patch (Nicotine)	SS		

Date:03/01/00ISR Number: 3549535-4Report Type:Periodic  
Age:28 YR Gender:Male I/FU:I

Company Report #A0106984A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG (TWICE PER DAY) ORAL		Dry Mouth Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Claritin-D	C		

Date:03/01/00ISR Number: 3553421-3Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0106396A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Condition Aggravated Drug Dependence  Increased Appetite  Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Augmentin	C		

Date:03/01/00ISR Number: 3553424-9Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #A0106397A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain	Health	Bupropion			
150 MG /		Chills	Professional	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Flushing					
/ ORAL		Tinnitus					
		Tremor		Triamcinolone			
				Acetonide	C		
				Azelastine			
				Hydrochloride	C		
				Salbutamol Sulphate	C		
				Adrenaline	C		

Date:03/01/00ISR Number: 3553427-4Report Type:Periodic Company Report #A0106398A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Bupropion			
150 MG /				Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3553429-8Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0106399A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pain	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/00ISR Number: 3553433-XReport Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #A0106400A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Influenza Like Illness Pain In Extremity	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/00ISR Number: 3553435-3Report Type:Periodic  
Age:65 YR Gender:Male I/FU:I

Company Report #A0106401A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/00ISR Number: 3553437-7Report Type:Periodic  
Age:65 YR Gender:Male I/FU:I

Company Report #A0106402A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL	3 DAY	Drug Interaction Erythema Feeling Hot Hyperaesthesia Paraesthesia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3553441-9Report Type:Periodic Company Report #A0106453A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL	5 WK	Sleep Disorder	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3553445-6Report Type:Periodic Company Report #A0106455A  
 Age:42 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Irritability Major Depression	Consumer Other	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3553448-1Report Type:Periodic Company Report #A0106463A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY / ORAL		Galactorrhoea	Health Professional Company Representative	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3553450-XReport Type:Periodic Company Report #A0106466A  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL		Discomfort Feeling Abnormal Muscle Twitching	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3553453-5Report Type:Periodic Company Report #A0106495A  
 Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL		Dermatitis	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3553460-2Report Type:Periodic Company Report #A0106504A  
 Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL		Dermatitis	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3553463-8Report Type:Periodic Company Report #A0106505A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL			Professional				

Date:03/01/00ISR Number: 3553468-7Report Type:Periodic Company Report #A0106887A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bronchitis Nervousness	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Tachycardia					
PER DAY ORAL							

Date:03/01/00ISR Number: 3553473-0Report Type:Periodic Company Report #A0106888A  
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE			Professional				
PER DAY							
ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3553477-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0106889A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Irritability	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG	ORAL 1 WK						

Date:03/01/00ISR Number: 3553481-XReport Type:Periodic  
 Age:44 YR Gender:Female I/FU:I

Company Report #A0106890A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Pruritic Skin Exfoliation	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG	TWICE PER DAY ORAL 3 WK						

Date:03/01/00ISR Number: 3553485-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0106891A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG	ORAL						

Date:03/01/00ISR Number: 3553489-4Report Type:Periodic  
 Age:25 YR Gender:Female I/FU:I

Company Report #A0106893A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nightmare	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG	TWICE PER DAY ORAL						
				Lisinopril	C		
				Oral Contraceptive	C		

Date:03/01/00ISR Number: 3553498-5Report Type:Periodic Company Report #A0105913A  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amenorrhoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							
DAY / ORAL							

Date:03/01/00ISR Number: 3553502-4Report Type:Periodic Company Report #A0105914A  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Disturbance In Attention	Professional				
PER DAY /		Dry Mouth					
ORAL		Insomnia		Nicotine	C		

Date:03/01/00ISR Number: 3553505-XReport Type:Periodic Company Report #A0105915A  
 Age:68 YR Gender:Female I/FU:I

Outcome	PT
	Decreased Appetite
	Headache
	Insomnia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nausea

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG /		Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY						
/ ORAL						

Date:03/01/00ISR Number: 3553509-7Report Type:Periodic Company Report #A0105916A  
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Swelling	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Toothache					
TWICE PER DAY							
/ ORAL							

Date:03/01/00ISR Number: 3553513-9Report Type:Periodic Company Report #A0105917A  
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/00ISR Number: 3553516-4Report Type:Periodic Company Report #A0106201A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Excitability	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Feeling Abnormal					
Date:03/01/00ISR Number: 3553520-6Report Type:Periodic			Company Report #A0106204A				
Age:37 YR Gender:Female I/FU:I							
Dose Required		Anxiety	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Depression	Professional				
Intervention to		Drug Withdrawal Syndrome					
TWICE PER DAY							
Prevent Permanent		Insomnia		Extra Strength			
/ ORAL		Nervousness		Tylenol Pm	C		
Impairment/Damage		Palpitations		Kava	C		
		Tremor					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Date:03/01/00ISR Number: 3553524-3Report Type:Periodic	Company Report #A0106206A						
Age:46 YR Gender:Female I/FU:I							
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Headache					
TWICE PER DAY							
/ ORAL				Paracetamol	C		



150 MG / PER	Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY / ORAL	Excitability					
ORAL	Flushing		Caffeine			
	Headache		(Formulation			
	Tachycardia		Unknown) (Caffeine)	SS		ORAL

Date:03/01/00ISR Number: 3553536-XReport Type:Periodic Company Report #A0106228A  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER		Headache	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY/ ORAL		Nausea	Professional				

Date:03/01/00ISR Number: 3553539-5Report Type:Periodic Company Report #A0106230A  
 Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dermatitis	Consumer	Bupropion	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Pruritus		Hydrochloride			
/ ORAL		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3553542-5Report Type:Periodic  
Age:67 YR Gender:Female I/FU:I

Company Report #A0106232A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Dermatitis					
PER DAY/ORAL		Middle Insomnia					
		Tremor					

Date:03/01/00ISR Number: 3553545-0Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #A0106276A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Diarrhoea					
TWICE PER DAY		Haematochezia					
/ ORAL		Hyperhidrosis					
		Tremor					

Date:03/01/00ISR Number: 3553547-4Report Type:Periodic  
Age:55 YR Gender:Male I/FU:I

Company Report #A0106390A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Dizziness					
DAY / ORAL				Multivitamin	C		
				Calcium Salt	C		

Date:03/01/00ISR Number: 3553548-6Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #A0106392A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							
DAY / ORAL							
				Guaifen	C		
				Co-Trimoxazole	C		

Date:03/01/00ISR Number: 3553553-XReport Type:Periodic Company Report #A0106393A  
 Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gingival Bleeding	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/00ISR Number: 3553555-3Report Type:Periodic Company Report #A0106394A  
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3553557-7Report Type:Periodic  
Age:37 YR Gender:Male I/FU:I

Company Report #A0106395A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL	2 MON	Dermatitis Erythema Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3553563-2Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0104673A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ ORAL		Pruritus Tremor Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3553565-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0104674A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL	3 WK	Anxiety Dizziness Drug Ineffective Nervousness		Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3553570-XReport Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #A0104702A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression	Consumer	Bupropion			

150 MG/ PER	Lethargy		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
DAY/ORAL						
Date:03/01/00ISR Number: 3553574-7Report Type:Periodic			Company Report #A0104703A			
Age:45 YR	Gender:Male	I/FU:I				
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Abdominal Pain	Consumer	Bupropion		
		Nausea		Hydrochloride	PS	Glaxo Wellcome Inc
		Pain In Extremity				

Date:03/01/00ISR Number: 3553577-2Report Type:Periodic			Company Report #A0104704A			
Age:26 YR	Gender:Female	I/FU:I				
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Headache	Consumer	Bupropion		
		Insomnia		Hydrochloride	PS	Glaxo Wellcome Inc
150 MG/TWICE		Nausea				
PER DAY/ORAL		Neck Pain				
		Tinnitus				
		Weight Decreased				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3553580-2Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #A0104705A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia Nausea	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER DAY/ORAL							

Date:03/01/00ISR Number: 3553585-1Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0104742A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE PER DAY/ORAL							

Date:03/01/00ISR Number: 3553589-9Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0104778A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tachycardia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY / ORAL							

Date:03/01/00ISR Number: 3553594-2Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #A0104861A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Bupropion			

150 MG /  
TWICE PER DAY  
/ ORAL

Date:03/01/00ISR Number: 3553598-XReport Type:Periodic Company Report #A0104862A  
Age:59 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Insomnia	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

TWICE PER DAY  
/ ORAL WEEKS

Date:03/01/00ISR Number: 3553601-7Report Type:Periodic Company Report #A0104863A  
Age:33 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY /ORAL		Dermatitis Swelling	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3553605-4Report Type:Periodic  
Age:28 YR Gender:Male I/FU:I

Company Report #A0104864A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth Mood Altered Personality Change	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE PER DAY /ORAL							

Date:03/01/00ISR Number: 3553608-XReport Type:Periodic  
Age:65 YR Gender:Female I/FU:I

Company Report #A0104865A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia Nausea	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY / ORAL							

Date:03/01/00ISR Number: 3553610-8Report Type:Periodic  
Age:65 YR Gender:Female I/FU:I

Company Report #A0104884A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY / ORAL							

Date:03/01/00ISR Number: 3553614-5Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #A0105053A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Abnormal Dreams Depressed Mood Dermatitis Insomnia Pruritus Urticaria	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3553615-7Report Type:Periodic Company Report #A0105054A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Overdose Palpitations	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3553618-2Report Type:Periodic Company Report #A0105055A  
 Age:35 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3553621-2Report Type:Periodic  
Age:75 YR Gender:Female I/FU:I

Company Report #A0105057A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Increased Appetite Pain	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Asthma Medication	C		

Date:03/01/00ISR Number: 3553623-6Report Type:Periodic  
Age:65 YR Gender:Male I/FU:I

Company Report #A0105058A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Flatulence Nausea Paraesthesia Oral	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3553625-XReport Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #A0105083A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dizziness Malaise	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3553630-3Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #A0108634A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Dizziness					
DAY / ORAL		Early Morning Awakening					
		Insomnia					
		Nausea					
		Nightmare					

Date:03/01/00ISR Number: 3553631-5Report Type:Periodic Company Report #A0108785A  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diplopia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Insomnia					
DAY / ORAL		Vision Blurred					

Date:03/01/00ISR Number: 3553635-2Report Type:Periodic Company Report #A0108786A  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Tobacco Abuse					
DAY / ORAL							





150 MG / Headache Consumer Zyban PS Glaxo Wellcome Inc ORAL  
Nausea  
TWICE PER DAY  
/ ORAL

Date:03/01/00ISR Number: 3553647-9Report Type:Periodic Company Report #A0108882A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL	5 DAY					

Date:03/01/00ISR Number: 3553649-2Report Type:Periodic Company Report #A0108883A  
Age:77 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Balance Disorder	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /						

TWICE PER DAY  
/ ORAL 8 DAY

Date:03/01/00ISR Number: 3553655-8Report Type:Periodic Company Report #A0108884A  
Age:67 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Abdominal Distension	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER						
DAY / ORAL	Pain					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3553659-5Report Type:Periodic Company Report #A0108885A  
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:03/01/00ISR Number: 3553666-2Report Type:Periodic Company Report #A0108886A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Nightmare	Professional				
DAY / ORAL							

Date:03/01/00ISR Number: 3553670-4Report Type:Periodic Company Report #A0108887A  
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/00ISR Number: 3553674-1Report Type:Periodic Company Report #A0108888A  
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Pollakiuria					
TWICE PER DAY							

/ ORAL

Date:03/01/00ISR Number: 3553675-3Report Type:Periodic Company Report #A0108889A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Drug Ineffective					
TWICE PER DAY		Nervousness					
/ ORAL		Tremor					

Date:03/01/00ISR Number: 3553677-7Report Type:Periodic Company Report #A0109060A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Paraesthesia Oral	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Tongue Oedema					
DAY / ORAL				Herbal Medication	C		

Date:03/01/00ISR Number: 3553678-9Report Type:Periodic Company Report #A0109061A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Glossodynia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							

/ ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3553680-7Report Type:Periodic  
 Age:46 YR Gender:Male I/FU:I

Company Report #A0109062A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Blood Pressure Increased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Vision Blurred					
/ ORAL							

Date:03/01/00ISR Number: 3553682-0Report Type:Periodic  
 Age:42 YR Gender:Female I/FU:I

Company Report #A0109063A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							

Date:03/01/00ISR Number: 3553685-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0109064A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /TWICE		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ORAL		Tinnitus					

Date:03/01/00ISR Number: 3554700-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0108118A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Diarrhoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

TWICE PER DAY

/ ORAL

Date:03/01/00ISR Number: 3554701-8Report Type:Periodic Company Report #A0108120A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Malaise	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Nervousness					

TWICE PER DAY

/ ORAL

Date:03/01/00ISR Number: 3554702-XReport Type:Periodic Company Report #A0108233A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cognitive Deterioration	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Depressed Level Of Consciousness					
		Dizziness					
		Dry Mouth					
		Headache					
		Insomnia					

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Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3554703-1Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #A0108249A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Diarrhoea Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3554704-3Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #A0108253A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Nausea Tremor	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3554705-5Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #A0108254A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Nervousness Palpitations	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3554706-7Report Type:Periodic  
Age:34 YR Gender:Male I/FU:I

Company Report #A0108255A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

TWICE PER DAY

/ ORAL

Date:03/01/00ISR Number: 3554707-9Report Type:Periodic Company Report #A0108256A  
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

150 MG/ TWICE

PER DAY /

ORAL

Date:03/01/00ISR Number: 3554708-0Report Type:Periodic Company Report #A0108257A  
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

150 MG /

TWICE PER DAY

/ ORAL

Date:03/01/00ISR Number: 3554709-2Report Type:Periodic Company Report #A0108258A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

150 MG /

TWICE PER DAY

/ ORAL





150 MG /  
Nausea  
TWICE PER  
DAY/ ORAL

Date:03/01/00ISR Number: 3554714-6Report Type:Periodic Company Report #A0108263A  
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

150 MG /  
TWICE PER DAY  
/ ORAL

Date:03/01/00ISR Number: 3554715-8Report Type:Periodic Company Report #A0108264A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

150 MG /  
TWICE PER DAY  
/ ORAL

Date:03/01/00ISR Number: 3554716-XReport Type:Periodic Company Report #A0108265A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

150 MG /  
TWICE PER DAY  
/ ORAL



150 MG	Oligomenorrhoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
/UNK/ORAL	1 MON					
Date:03/01/00ISR Number: 3554721-3Report Type:Periodic Company Report #A0107208A						
Age:	Gender:Female	I/FU:I				
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose						Route
150 MG /		Nervousness	Consumer	Zyban	PS	Glaxo Wellcome Inc
TWICE PER DAY		Nightmare				ORAL
/ ORAL				Caffeine	C	
Date:03/01/00ISR Number: 3554722-5Report Type:Periodic Company Report #A0107211A						
Age:51 YR	Gender:Male	I/FU:I				
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose						Route
150 MG /		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc
TWICE PER DAY						ORAL
/ ORAL						
Date:03/01/00ISR Number: 3554723-7Report Type:Periodic Company Report #A0107213A						
Age:33 YR	Gender:Female	I/FU:I				
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose						Route
150 MG /		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc
TWICE PER DAY						ORAL
/ ORAL						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3554724-9Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0107223A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Dry Mouth					
/ ORAL		Insomnia					

Date:03/01/00ISR Number: 3554725-0Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0107224A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Night Sweats	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
ORA L				Ibuprofen	C		
				Birth Control	C		

Date:03/01/00ISR Number: 3554726-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0107234A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Arrhythmia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Chest Pain					
/ OROAL		Discomfort					
		Flushing					
		Insomnia					
		Nausea					
		Sensation Of Pressure					
		Vision Blurred					

Date:03/01/00ISR Number: 3554727-4Report Type:Periodic Company Report #A0107249A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mouth Ulceration	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ORAL							

Date:03/01/00ISR Number: 3554728-6Report Type:Periodic Company Report #A0107293A  
Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Dissociation					
TWICE PER DAY		Disturbance In Attention					
/ORAL		Dizziness					
		Feeling Abnormal					
		Hyperhidrosis					
		Urinary Incontinence					

Date:03/01/00ISR Number: 3554729-8Report Type:Periodic Company Report #A0107393A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Excitability					
TGWICE PER							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

DAY / ORAL

Date:03/01/00ISR Number: 3554730-4Report Type:Periodic Company Report #A0107483A  
 Age:56 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Burning Sensation Dermatitis Dermatitis Exfoliative Rash Generalised	Health Professional	Zyban	PS	Glaxo Wellcome Inc	

Date:03/01/00ISR Number: 3554731-6Report Type:Periodic Company Report #A0107494A  
 Age:52 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

TWICE PER DAY

/ ORAL

Date:03/01/00ISR Number: 3554732-8Report Type:Periodic Company Report #A0107495A  
 Age:63 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

TWICE PER DAY

/ ORAL

Date:03/01/00ISR Number: 3554733-XReport Type:Periodic Company Report #A0107496A  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Tremor					
PER DAY /							
ORAL							

Date:03/01/00ISR Number: 3554734-1Report Type:Periodic Company Report #A0107586A  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bone Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Chills					
TWICE PER DAY		Diarrhoea					
/ ORAL		Pyrexia		Nicotine Patch	SS		
TRANSDERMAL	21 MG /	UNK /					
TRANSDERMAL		Rash Erythematous					
		Sleep Disorder					
		Vomiting Projectile					

Date:03/01/00ISR Number: 3554735-3Report Type:Periodic Company Report #A0107640A  
 Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Tremor					
DAY / ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3554736-5Report Type:Periodic Company Report #A0107649A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / UNK/							
ORAL							

Date:03/01/00ISR Number: 3554737-7Report Type:Periodic Company Report #A0107717A  
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/01/00ISR Number: 3554738-9Report Type:Periodic Company Report #A0107719A  
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression Irritability	Consumer	Zyban	PS	Glaxo Wellcome Inc	

Date:03/01/00ISR Number: 3554739-0Report Type:Periodic Company Report #A0107720A  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Conjunctival Hyperaemia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
		Dermatitis					
		Eyelid Oedema					

Flonase Aqueous  
Spray (Fluticasone  
Propionate ) SS

1 SPRAY / PER

DY /

INTRANASAL

Date:03/01/00ISR Number: 3554740-7Report Type:Periodic Company Report #A0105647A  
Age:37 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE		Dissociation Disturbance In Attention	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ORAL		Hyperphagia					
		Speech Disorder		Estrogen	C		
		Urticaria		Amitriptyline Hcl	C		

Date:03/01/00ISR Number: 3554741-9Report Type:Periodic Company Report #A0105714A  
Age:28 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE		Agitation Irritability	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY /ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3554742-0Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0105720A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dissociation Feeling Abnormal	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Major Depression					
PER DAY/ORAL							

Date:03/01/00ISR Number: 3554743-2Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0105722A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER							
DAY/ORAL							

Date:03/01/00ISR Number: 3554744-4Report Type:Periodic  
Age:73 YR Gender:Female I/FU:I

Company Report #A0105723A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation Dry Mouth	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Dysgeusia					
PER DAY/ORAL		Insomnia Nausea					

Date:03/01/00ISR Number: 3554745-6Report Type:Periodic  
Age:74 YR Gender:Female I/FU:I

Company Report #A0105724A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Health	Bupropion			

150 MG/SEE  
TEXT/ORAL  
Constipation  
Drug Ineffective  
Dry Mouth  
Insomnia  
Libido Increased  
Nausea  
Tremor

Date:03/01/00ISR Number: 3554746-8Report Type:Periodic Company Report #A0105725A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Constipation	Consumer	Bupropion			
		Dyspepsia		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER		Migraine					
DAY /ORAL		Pruritus					
		Rash Erythematous					

Date:03/01/00ISR Number: 3554747-XReport Type:Periodic Company Report #A0105726A  
Age:36 YR Gender:Female I/FU:I

Outcome	PT
	Face Oedema
	Feeling Abnormal
	Feeling Hot
	Headache

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Hypoaesthesia Psychomotor Hyperactivity	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
PER DAY/ORAL							

Date:03/01/00ISR Number: 3554748-1Report Type:Periodic Company Report #A0105759A  
Age: Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Rash Erythematous Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Dose							
150 MG/TWICE							
PER DAY/ORAL							

Date:03/01/00ISR Number: 3554749-3Report Type:Periodic Company Report #A0105762A  
Age:45 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Arthralgia Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Dose							
150 MG/TWICE							
PER DAY./ORAL		Swelling					
		Urticaria					

Date:03/01/00ISR Number: 3554750-XReport Type:Periodic Company Report #A0105763A  
Age:38 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Headache	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Dose							
150 MG./TWICE							

PER DAY/ORAL

Date:03/01/00ISR Number: 3554751-1Report Type:Periodic Company Report #A0105765A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL 20 DAY							

Date:03/01/00ISR Number: 3554752-3Report Type:Periodic Company Report #A0105854A

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Apathy Dysgeusia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
PER DAY/ ORAL							
		Insomnia					
		Sedation					

Date:03/01/00ISR Number: 3554753-5Report Type:Periodic Company Report #A0105856A

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/00ISR Number: 3554754-7Report Type:Periodic  
Age:28 YR Gender:Male I/FU:I

Company Report #A0105859A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Emotional Disorder Tachycardia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/01/00ISR Number: 3554755-9Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #A0105862A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/PER DAY/ ORAL		Dermatitis Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Alesse C

Date:03/01/00ISR Number: 3554756-0Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #A0105863A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ORAL		Constipation	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Nicotine C

Date:03/01/00ISR Number: 3554757-2Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #A0105865A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Balance Disorder	Consumer	Bupropion			

150 MG/TWICE		Bone Pain		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Dizziness					
PER DAY/ORAL		Nausea					

Date:03/01/00ISR Number: 3554758-4Report Type:Periodic Company Report #A0105911A  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash Erythematous Swelling	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Urticaria					
PER DAY/ORAL				Lovastatin	C		
				Lithium Salt	C		
				Multivitamin	C		

Date:03/01/00ISR Number: 3554759-6Report Type:Periodic Company Report #A0105912A  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Diarrhoea Herpes Simplex	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Lip Ulceration					
PER DAY/ORAL		Swelling					
		Urticaria					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/00ISR Number: 3468081-XReport Type:Expedited (15-DaCompany Report #A0112057A  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Joint Swelling Pruritus Urticaria	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:03/02/00ISR Number: 3468093-6Report Type:Expedited (15-DaCompany Report #A0112807A  
 Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction Sudden Death	Health Professional Company	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / ORAL	4 DAY		Representative	Fentanyl Celecoxib	C C		

Date:03/02/00ISR Number: 3468094-8Report Type:Expedited (15-DaCompany Report #A0112851A  
 Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Burning Sensation Dermatitis Pain	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /		Rash Erythematous					
TWICE DAY /							
ORAL							

Date:03/06/00ISR Number: 3470357-7Report Type:Expedited (15-DaCompany Report #A0113407A  
Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Ecchymosis Epistaxis Skin Disorder Thrombocytopenia	Health Professional Company Representative	Wellbutrin Tablet -Controlled Release (Bupropion Hydrochloride)	PS		ORAL
UNK / SEE							
TEXT / ORAL							

Date:03/07/00ISR Number: 3471269-5Report Type:Expedited (15-DaCompany Report #A0113660A  
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dermatitis Drug Interaction Feeling Hot	Foreign Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL							
		Joint Swelling		Pseudoephedrine Hcl (Formulation Unknown) (Non-Us Product)	SS		
		Rash Erythematous		Chlorpheniramine Maleate (Formulation Unknown) (Chlorpheniramine Maleate)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/07/00ISR Number: 3471367-6Report Type:Expedited (15-DaCompany Report #A0112576A

Age:39 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	PT Hepatic Failure Rash Vesicular Renal Failure	Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		ORAL

ORAL

Date:03/07/00ISR Number: 3471517-1Report Type:Expedited (15-DaCompany Report #B0075853A

Age:36 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG / SEE TEXT / UNKNOWN	PT Anxiety Chest Discomfort Dizziness Dry Mouth Herpes Simplex Hypertension Hyperventilation Insomnia Nausea Oral Intake Reduced Palpitations Tachycardia Tremor	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		

Date:03/08/00ISR Number: 3471158-6Report Type:Direct

Company Report #

Age:58 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 100MG PO QD Initial or Prolonged	PT Dizziness Dysphagia Dyspnoea Finger Deformity	Health Professional	Wellbutrin Sr 100 Mg	PS		ORAL

Influenza Like Illness  
Motor Dysfunction  
Nausea  
Paraesthesia

Date:03/08/00ISR Number: 3472130-2Report Type:Expedited (15-DaCompany Report #A0110370A  
Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cerebrovascular Accident Speech Disorder	Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Date:03/08/00ISR Number: 3472139-9Report Type:Expedited (15-DaCompany Report #A0107781A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion Drug Interaction	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL				Citalopram Hydrobromide			

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Freedom Of Information (FOI) Report

(Formulation  
Unknown) (Citalopram  
Hydrobromide) SS

Date:03/08/00ISR Number: 3472140-5Report Type:Expedited (15-DaCompany Report #A0107071A  
Age:78 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG/ PER DAY/ ORAL	Atrophy Confusional State Convulsion Delusion	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
	20 MG/ UNKNOWN	Disorientation Drug Interaction Hypersensitivity Joint Stiffness Miosis Nervous System Disorder		Citalopram Hydrobromide (Formulation Unknown) (Citalopram Hydrobromide)	SS		
		Pupil Fixed Pupillary Reflex Impaired Sedation Sleep Disorder Urinary Tract Infection Visual Disturbance		Hormones Multivitamin	C C		

Date:03/09/00ISR Number: 3472617-2Report Type:Expedited (15-DaCompany Report #B0078241A  
Age:78 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG / SINGLE DOSE / ORAL	Hypotension	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
				Captopril	C		

Date:03/09/00ISR Number: 3472619-6Report Type:Expedited (15-DaCompany Report #A0114129A  
Age:49 YR Gender:Female I/FU:I

Outcome	PT
Other	Arthralgia
	Asthenia
	Chest Pain
	Coordination Abnormal
	Cough
	Cyanosis
	Disorientation
	Dizziness
	Dyspepsia
	Fatigue
	Feeling Abnormal
	Feeling Cold
	Feeling Hot
	Feeling Jittery
	Headache
	Hearing Impaired
	Insomnia
	Neck Pain

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE	PER DAY /	Oedema Pallor Paraesthesia Parkinsonian Gait Pollakiuria Sneezing	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL		Tinnitus Tremor Vertigo Visual Acuity Reduced		Diuretic	C		

Date:03/09/00ISR Number: 3473462-4Report Type:Expedited (15-DaCompany Report #A0110477A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL		Abdominal Pain Faecaloma	Consumer	Zyban	PS		ORAL

Date:03/09/00ISR Number: 3473464-8Report Type:Expedited (15-DaCompany Report #A0055937A  
Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT
Hospitalization - Initial or Prolonged Disability		Alopecia Anuria Anxiety Arrhythmia Supraventricular Asthenia Autonomic Nervous System Imbalance Bipolar Ii Disorder Blood Triglycerides Increased Bradycardia Chills Condition Aggravated

Convulsion  
Depression  
Disorientation  
Dizziness  
Drug Hypersensitivity  
Dysarthria  
Dyspnoea  
Fatigue  
Haematuria  
Headache  
Hyperglycaemia  
Hyperhidrosis  
Hypertension  
Hyperventilation  
Hypoaesthesia  
Hypomania  
Labile Blood Pressure  
Localised Oedema  
Memory Impairment  
Mental Impairment  
Micturition Urgency  
Mitral Valve Incompetence  
Muscle Rigidity



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL	Muscle Twitching Musculoskeletal Stiffness Nervousness Obsessive-Compulsive Disorder	Health Professional	Zyban	PS		ORAL
	Oedema		Conjugated Estrogens	C		
	Palpitations		Vitamin	C		
	Panic Disorder Without Agoraphobia		Aspirin	C		
	Paraesthesia		Melatonin	C		
	Renal Disorder					
	Renal Impairment					
	Sensation Of Pressure					
	Tachycardia					
	Tinnitus					
	Tongue Oedema					
	Transient Ischaemic Attack					
	Tremor					
	Urethral Stricture					
	Urinary Incontinence					
	Ventricular Extrasystoles					
	Weight Increased					

Date:03/10/00ISR Number: 3474118-4Report Type:Periodic Company Report #8-99137-155A  
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1 TABLET 1X PER DAY, ORAL		Alopecia	Consumer	Lo/Ovral Tablet	PS		ORAL
ORAL				Lithium	SS		ORAL
ORAL				Wellbutrin	SS		ORAL
				Lithium	C		
				Ritalin (Methylphenidate)	C		
				Wellbutrin (Bupropion)	C		

Date:03/10/00ISR Number: 3474219-0Report Type:Expedited (15-DaCompany Report #A007269

Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Angioneurotic Oedema	Consumer	Zoloft Tablets	PS	50.00 Mg	
Intervention to Prevent Permanent Impairment/Damage		Anorgasmia				Total:Daily:Oral	ORAL
		Face Oedema		Wellbutrin	SS		
		Libido Decreased		Ortho Tri-Cyclen	C		
		Oedema Peripheral					
		Pruritus					
		Tongue Oedema					
		Urticaria					

Date:03/10/00ISR Number: 3474517-0Report Type:Expedited (15-DaCompany Report #A007267

Age:18 YR Gender:Female I/FU:I

Outcome  
Hospitalization -  
Initial or Prolonged  
Required

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Intervention to Prevent Permanent Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
5.00 MG		Bone Marrow Depression Drug Interaction Infection In An Immunocompromised Host	Health Professional Company Representative	Glucotrol Xl Extended Release Tablets	PS		ORAL
TOTAL; DAILY;		Leukopenia					
ORAL		Varicella Vomiting		Wellbutin Ritalin	SS SS		

Date:03/13/00ISR Number: 3474992-1Report Type:Expedited (15-DaCompany Report #A0107071A  
Age:78 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cerebral Atrophy Coma Confusional State Convulsion	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ PER DAY/ ORAL		Delusion Disorientation Drug Interaction Hypersensitivity Joint Stiffness		Citalopram Hydrobromide (Formulation Unknown) (Citalopram Hydrobromide)	SS		
20 MG		Miosis Nervous System Disorder Pupil Fixed Sedation Urinary Tract Infection Visual Disturbance		Hormones Multivitamin	C C		

Date:03/13/00ISR Number: 3474994-5Report Type:Expedited (15-DaCompany Report #A0068941A  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation Conversion Disorder Dizziness	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG ORAL		Feeling Abnormal Hallucination, Auditory Nephrolithiasis Renal Colic Tremor Urinary Tract Infection Vomiting					

Date:03/13/00ISR Number: 3475026-5Report Type:Expedited (15-DaCompany Report #A0109223A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia Inflammation Pain In Extremity	Health Professional	Zyban Tablet- Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE		Pruritus					
PER DAY/ ORAL		Urticaria Vasculitis					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/14/00ISR Number: 3475615-8Report Type:Expedited (15-DaCompany Report #B0078436A  
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Atrial Tachycardia Stress	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL				Nicotine	C		

Date:03/14/00ISR Number: 3475623-7Report Type:Expedited (15-DaCompany Report #D0006860A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coronary Artery Surgery Influenza Respiratory Disorder	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Date:03/14/00ISR Number: 3475624-9Report Type:Expedited (15-DaCompany Report #A0114289A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Glaucoma	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Date:03/14/00ISR Number: 3475628-6Report Type:Expedited (15-DaCompany Report #A0110705A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged Other		Grand Mal Convulsion Sleep Disorder	Foreign Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

150 MG /

TWICE PER DAY

/ ORAL

Doxepin  
Hydrochloride C  
Herbal Medication C  
Thanol C

Date:03/14/00ISR Number: 3475629-8Report Type:Expedited (15-DaCompany Report #A0107356A

Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chromaturia Drug Interaction Haematuria	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ ORAL		Haemorrhage International Normalised Ratio Increased Prothrombin Time Prolonged Pyrexia Renal Colic Tremor		Warfarin Sodium (Formulation Unknoen) (Warfarin Sodium)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/15/00ISR Number: 3476290-9Report Type:Direct  
Age:31 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1T QD		Dermatitis		Zyban 150mg	PS		

Date:03/15/00ISR Number: 3476417-9Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to Prevent Permanent Impairment/Damage		Chills Erythema Joint Stiffness Joint Swelling Skin Disorder	Health Professional	Zyban Sr 150 Mg Tablet	PS		

Date:03/15/00ISR Number: 3476990-0Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #19990600691

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100 MG QD PO		Fatigue	Consumer	Toprol-Xl	PS		ORAL
10 MG QD PO		Nausea		Meridia	SS		ORAL
100 MG BID PO		Vision Blurred		Wellbutrin	SS		ORAL
				Ogen	C		

Date:03/16/00ISR Number: 3476537-9Report Type:Expedited (15-DaCompany Report #A0112851A  
Age:35 YR Gender:Male I/FU:F

Company Report #A0112851A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required Intervention to 150 MG/TWICE		Arthralgia Burning Sensation Dyspnoea	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Prevent Permanent Face Oedema  
 PER DAY/ORAL  
 Impairment/Damage Pain  
 Rash Erythematous  
 Rash Macular

Date:03/16/00ISR Number: 3477052-9Report Type:Expedited (15-DaCompany Report #A0111797A  
 Age:67 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG/ PER DAY/ ORAL	Arrhythmia Confusional State Dehydration Depressed Level Of Consciousness Dizziness	Study Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
1 TABLET/ PER DAY/ ORAL	Dysarthria Hypotension Myocardial Ischaemia Ventricular Tachycardia		Placebo Tablet (Placebo)	SS		ORAL
			Clopidogrel Bisulphate Atenolol Lisinopril Omeprazole Aspirin Atorvastatin Calcium Nitroglycerin	C C C C C C C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/16/00ISR Number: 3477054-2Report Type:Expedited (15-DaCompany Report #A0111148A  
Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required		Amnesia	Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		ORAL
Intervention to Prevent Permanent 100 MG/ ORAL Impairment/Damage		Gait Disturbance Muscle Rigidity Parkinson'S Disease Skin Disorder Speech Disorder		Trazodone Estratest Thyroxine Sodium Lisinopril Flunisolide Salbutamol Sulphate Clorazepate Dipotassium Salmeterol Xinafoate Nabumetone Ibuprofen	C C C C C C C C C C C		

Date:03/17/00ISR Number: 3477421-7Report Type:Expedited (15-DaCompany Report #A0114528A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Asthenia Chest Discomfort Communication Disorder	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER DAY / ORAL		Jaw Disorder Nausea Respiratory Distress Tongue Disorder Tongue Oedema Tremor		Ethanol (Formulation Unknown) (Alcohol)	SS		

Date:03/20/00ISR Number: 3477570-3Report Type:Expedited (15-DaCompany Report #A0107356A  
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chills	Health	Zyban Tablet - Zyban			
Other		Chromaturia	Professional	(Bupropion			
		Drug Interaction		Hydrochloride)	PS		ORAL
150 MG / UNK		Ear Haemorrhage					
/ ORAL		Haematuria		Warfarin Sodium			
		Haemorrhage		(Formulation			
		International Normalised		Unknown)(Warfarin			
		Ratio Increased		Sodium)	SS		ORAL
10 MG / UNK /		Prothrombin Time					
ORAL		Prolonged					
		Pyrexia					
		Renal Colic					

Date:03/21/00ISR Number: 3477185-7Report Type:Direct Company Report #  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Balance Disorder		Zyban Sr 150mg Glaxo			
Disability		Difficulty In Walking		Wellcome	PS	Glaxowellcome	ORAL
150MG ONE A		Diplopia					
DAY ORAL		Sleep Disorder					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/21/00ISR Number: 3477736-2Report Type:Direct  
Age:28 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS		
150MG/D				Zyban	SS		
150 MG/D.							

Date:03/22/00ISR Number: 3478723-0Report Type:Expedited (15-DaCompany Report #A0114289A  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Foreign	Zyban (Bupropion			
ORAL		Glaucoma	Consumer	Hydrochloride)	PS		ORAL

Date:03/22/00ISR Number: 3478777-1Report Type:Expedited (15-DaCompany Report #A0115400A  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Ecchymosis	Foreign	Zyban Tablet - Zyban			
Initial or Prolonged		Eyelid Oedema	Consumer	(Bupropion			
150 MG /		Joint Swelling		Hydrochloride)	PS		ORAL
TWICE PER DAY		Pharyngeal Oedema					
/ ORAL		Rash Generalised					
		Swelling					
		Urticaria					
		Vaginal Haemorrhage					

Date:03/23/00ISR Number: 3479269-6Report Type:Expedited (15-DaCompany Report #A0115630A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged	Anxiety Dermatitis Bullous Eczema Nervousness	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS	ORAL
150 MG/ TWICE  PER DAY/ ORAL	Rash Erythematous  Rash Generalised Rash Macular Rash Papular Toxic Epidermal Necrolysis		Lisinopril Hydrochlorothiazide	C C	

Date:03/24/00ISR Number: 3479745-6Report Type:Expedited (15-DaCompany Report #044-0991-980010  
Age:78 YR Gender:Female I/FU:I

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - PER ORAL	Angioneurotic Oedema Dyspnoea	Foreign Health	Rezulin (Troglitazone)	PS		ORAL
Initial or Prolonged  300 MG,  DAILY, PER  ORAL	Face Oedema Pharyngeal Oedema  Speech Disorder  Tachycardia	Professional	Zyban (Bupropion Hydrochloride)	SS		ORAL
			Amaryl (Glimepiride) Cardizem (Diltiazem Hydrochloride) Compazine (Prochlorperazine Edisylate)	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/24/00ISR Number: 3480099-XReport Type:Expedited (15-DaCompany Report #A0109179A

Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Chest Pain Drug Interaction	Health Professional Company Representative	Bupropion Hydrochloride Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
100							
MG/VARIABLE							
DOSE/ORAL							
ORAL				Pemoline Unspecified Tablet (Pemoline)	SS		ORAL
				Methylphenidate Hcl Tablet	SS		ORAL
5 MG TWICE							
PER DAY ORAL							

Date:03/24/00ISR Number: 3480102-7Report Type:Expedited (15-DaCompany Report #A0114679A

Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Convulsion Drug Interaction Feeling Jittery Fungal Infection	Foreign	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG TWICE							
PER DAY ORAL		Pain					
		Pneumonia Tremor		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	SS		ORAL
300 MG/FOUR							
TIMES PER DAY							
ORAL							

300 MG/PER

Fluconazole  
(Fluconazole)

SS

ORAL

DAY/ORAL

Cyclophosphamide C  
Fluorouracil C  
Methotrexate C

Date:03/27/00ISR Number: 3480380-4Report Type:Expedited (15-DaCompany Report #A0114129A

Age:46 YR Gender:Female I/FU:F

Outcome	PT
Disability	Arthralgia
Other	Asthenia
	Chest Pain
	Coordination Abnormal
	Cough
	Disorientation
	Dizziness
	Dyspepsia
	Fatigue
	Feeling Abnormal
	Feeling Cold
	Feeling Hot
	Feeling Jittery
	Headache
	Hearing Impaired
	Insomnia
	Oedema

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Pallor Palpitations Paraesthesia	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
PER DAY/ ORAL		Sneezing		Diuretic	C		
		Tinnitus Visual Acuity Reduced					

Date:03/28/00ISR Number: 3480292-6Report Type:Direct  
Age:56 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required ONE TWICE		Hepatitis	Health Professional	Wellbutrin 75mg Bid	PS		
Intervention to DAILY		Hepatitis Cholestatic					
Prevent Permanent Impairment/Damage		Hepatocellular Damage Liver Function Test Abnormal		Ritalin 20mg Allergy Injection Alburteral Vancenase Ibuprofen 600mg	C C C C C		

Date:03/28/00ISR Number: 3481187-4Report Type:Expedited (15-DaCompany Report #A0115676A  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dizziness Dysphagia Dyspnoea Finger Deformity	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
100 MG/ PER DAY/ ORAL		Influenza Like Illness					
		Nausea Paraesthesia Retching		Nefazodone Hydrochloride	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Akinesia	Health	Zyban Tablet-Zyban			
		Angiopathy	Professional	(Bupropion			
		Arthralgia		Hydrochloride)	PS		ORAL
150 MG/ ORAL		Inflammation					
		Leukocytoclastic					
		Vasculitis					
		Pharyngolaryngeal Pain					
		Pruritus					
		Purpura					
		Rheumatoid Arthritis					
		Sneezing					
		Urticaria					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/00ISR Number: 3481311-3Report Type:Direct  
 Age:36 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Oedema Peripheral		Zyban 150 Mg	PS		
150 BID		Urticaria		Caladryl	C		

Date:03/30/00ISR Number: 3482141-9Report Type:Expedited (15-DaCompany Report #A0112392A  
 Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Foreign	Zyban Tablet - Zyban			
Required		Burning Sensation	Health	(Bupropion			
Intervention to		Complement Factor	Professional	Hydrochloride)	PS		ORAL
150 MG/ TWICE							
Prevent Permanent		Increased					
PER DAY/ ORAL							
Impairment/Damage		Decreased Activity		Mometasone Furoate	C		
		Dermatitis					
		Difficulty In Walking					
		Discomfort					
		Dysphagia					
		Erythema					
		Eyelid Oedema					
		Face Oedema					
		Fatigue					
		Feeling Cold					
		Insomnia					
		Obstruction					
		Oedema					
		Pain					
		Peripheral Coldness					
		Pharyngolaryngeal Pain					
		Rash Pruritic					
		Skin Discolouration					
		Type Iv Hypersensitivity					
		Reaction					
		Urticaria					
		Vasculitis					

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG/TWICE PER DAY/ORAL	Ecchymosis Eyelid Oedema Joint Swelling Pharyngeal Oedema Rash Generalised Urticaria Vaginal Haemorrhage	Foreign Health Professional	Zyban Tablet -Zyban (Bupropion Hydrochloride)	PS		ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other ONE TABLET 2 TIMES ORAL	Convulsion		Welbutrin Sr 150mg  Dilantin	PS  C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/31/00ISR Number: 3482801-XReport Type:Expedited (15-DaCompany Report #A0115944A

Age:8 MON Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Medication Error	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/	ORAL	Toxicologic Test Abnormal					

Date:03/31/00ISR Number: 3483217-2Report Type:Periodic Company Report #A002002

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Diabetes Mellitus	Consumer	Cardura	PS		ORAL
ORAL				Pravachol	SS		ORAL
				Lamisil	SS		ORAL
				Wellbutrin	SS		

Date:04/03/00ISR Number: 3483994-0Report Type:Expedited (15-DaCompany Report #B0079458A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG/ORAL	Angina Pectoris Drug Interaction	Foreign Health Professional Other	Zyban (Bupropion Hydrochloride)	PS		
				Thyroxine (Levothyroxine Sodium)	SS		

Date:04/03/00ISR Number: 3483997-6Report Type:Expedited (15-DaCompany Report #A0115984A

Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Angiopathy Blood Potassium Decreased	Consumer	Zyban Tablet - Zyban (Bupropion			

150 MG TWICE	Dermatitis	Hydrochloride)	PS	ORAL
PER DAY ORAL	Diarrhoea			
	Drug Interaction	Cisapride		
	Dyspnoea	(Formulation		
	Eyelid Oedema	Unknown) (Cisapride)	SS	
	Headache			
	Oedema			
	Pharyngeal Oedema			
	Pyrexia			

Date:04/03/00ISR Number: 3484034-XReport Type:Expedited (15-DaCompany Report #A0116641A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Circulatory Collapse	Foreign	Zyban	PS		ORAL
150 MG /		Myocardial Infarction	Consumer				
TWICE PER DAY							
/ ORAL							

Date:04/05/00ISR Number: 3484446-4Report Type:Expedited (15-DaCompany Report #00P-163-0087711-00(0)  
Age:29 YR Gender:Female I/FU:I

Outcome	PT
Life-Threatening	Aggression
	Coma

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Confusional State Depression Drug Interaction					
1000 MG, 1 IN	1 D, PER ORAL	Drug Level Below Therapeutic	Health Professional	Depakote (Depakote) (Divalproex Sodium)	PS		ORAL
		Dyspnoea					
150 MG, 1 IN	1 D, PER ORAL	Ecchymosis Fatigue		Amfebutamone (Amfebutamone)	SS		ORAL
		Folliculitis					
PER ORAL		Headache Hypotension Influenza Like Illness		Amantadine (Amantadine) (Amantadine)	SS		ORAL
		Movement Disorder Muscle Rigidity Nausea Oxygen Saturation Decreased Pallor Pupillary Reflex Impaired Respiratory Rate Decreased Speech Disorder Tachycardia Tremor		Tetracycline Hydrochloride (Tetracycline Hydrochloride) Skin Discoloration Pupillary Reflex Impaired Ecchymosis Lab Test Abnormality Hostility Confusion Headache Personality Disorder	C C C C C C C C C C C		

Date:04/06/00ISR Number: 3485009-7Report Type:Expedited (15-DaCompany Report #A0111455A  
Age:19 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Creatine Phosphokinase Increased Chest Pain	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL		Myocarditis		Cannabis Mushrooms	C C		

Date:04/07/00ISR Number: 3485172-8Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Grand Mal Convulsion		Bupropion	PS		ORAL
ORAL							
Hospitalization -							
Initial or Prolonged							

Date:04/07/00ISR Number: 3485505-2Report Type:Expedited (15-DaCompany Report #A0116863A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional Company Representative	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		ORAL
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/07/00ISR Number: 3485508-8Report Type:Expedited (15-DaCompany Report #A0116939A  
 Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Aspartate Aminotransferase Increased	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY / ORAL		Dermatitis Urticaria					

Date:04/07/00ISR Number: 3485511-8Report Type:Expedited (15-DaCompany Report #A0116472A  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Condition Aggravated Dizziness Meniere'S Disease	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY / ORAL		Nausea Syncope Vomiting					

Date:04/07/00ISR Number: 3485815-9Report Type:Expedited (15-DaCompany Report #A0116999A  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Change Of Bowel Habit Condition Aggravated Muscle Spasms	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY / ORAL							

Date:04/10/00ISR Number: 3486408-XReport Type:Expedited (15-DaCompany Report #A0116641A  
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Circulatory Collapse Coronary Ostial Stenosis Myocardial Infarction	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL							

Date:04/11/00ISR Number: 3486449-2Report Type:Direct Company Report #  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 9 DAYS PRIOR Initial or Prolonged TO EVENT Required Intervention to Prevent Permanent Impairment/Damage		Dermatitis Hypersensitivity Pruritus Throat Tightness		Bupropion	PS		

Date:04/11/00ISR Number: 3487016-7Report Type:Expedited (15-DaCompany Report #A0055937A  
Age:55 YR Gender:Female I/FU:F

Outcome  
Hospitalization -  
Initial or Prolonged

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## Freedom Of Information (FOI) Report

## Disability

PT  
Agitation  
Anaphylactic Shock  
Anxiety  
Autonomic Nervous System  
Imbalance  
Back Pain  
Bipolar Disorder  
Blood Pressure  
Fluctuation  
Bradycardia  
Chills  
Condition Aggravated  
Convulsion  
Delusion  
Depersonalisation  
Disorientation  
Disturbance In Attention  
Dizziness  
Drug Hypersensitivity  
Dysarthria  
Dyspnoea  
Fatigue  
Haematuria  
Hallucination, Auditory  
Hallucination, Olfactory  
Hyperglycaemia  
Hyperhidrosis  
Hypertension  
Hyperventilation  
Hypoaesthesia  
Localised Oedema  
Logorrhoea  
Mania  
Mental Impairment  
Mitral Valve Incompetence  
Mood Swings  
Muscle Twitching  
Muscular Weakness  
Musculoskeletal Stiffness  
Nervous System Disorder  
Nervousness  
Palpitations  
Panic Disorder  
Paraesthesia

Paranoia  
Psychomotor Hyperactivity  
Renal Disorder  
Renal Impairment  
Schizophrenia  
Supraventricular  
Extrasystoles  
Tachycardia  
Throat Tightness  
Tinnitus  
Tongue Disorder  
Tongue Oedema  
Transient Ischaemic  
Attack

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER		Urethral Stricture Urinary Incontinence Ventricular Extrasystoles Weight Increased	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
DAY/ ORAL				Conjugated Estrogens	C		
				Vitamin	C		
				Aspirin	C		
				Melatonin	C		

Date:04/12/00ISR Number: 3487652-8Report Type:Expedited (15-DaCompany Report #A0102058A  
Age:65 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150 MG/ PER		Blood Pressure	Study	Zyban	PS		ORAL
Hospitalization - DAY/ ORAL		Fluctuation	Health Professional				
Initial or Prolonged Required		Blood Pressure Increased	Professional	Me-Prednisolone Na			
Intervention to Prevent Permanent Impairment/Damage		Chest Pain Condition Aggravated Confusional State Depressed Level Of Consciousness Hypertensive Encephalopathy Speech Disorder Vision Blurred		Succ. Theophylline Diltiazem Lisinopril Salbutamol Sulphate Ipratropium Bromide Procainamide Hcl Aspirin Isosorbide Dinitrate Metoprolol Tartrate Vitamin E Clopidogrel Bisulphate	C C C C C C C C C C C C		

Date:04/13/00ISR Number: 3488063-1Report Type:Expedited (15-DaCompany Report #A0073901A  
Age:29 YR Gender:Female I/FU:F

Outcome PT

Life-Threatening  
Hospitalization -  
Initial or Prolonged  
Disability  
Other  
Required  
Intervention to  
Prevent Permanent  
Impairment/Damage

Abdominal Distension  
Arthralgia  
Decreased Activity  
Decreased Appetite  
Dermatitis  
Difficulty In Walking  
Dyspnoea Exertional  
Erythema  
Erythema Multiforme  
Eyelid Oedema  
Face Oedema  
Fall  
Fatigue  
Feeling Cold  
Fibromyalgia  
Food Intolerance  
Insomnia  
Joint Stiffness  
Leukocytosis  
Migraine  
Mouth Ulceration  
Pain

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Periorbital Oedema Proteinuria Pruritus	Health Professional Company Representative	Wellbutrin Tablet - Controlled Release (Bupropion Hydrochloride)	PS		ORAL
TWICE PER DAY		Serum Sickness Stevens-Johnson Syndrome Throat Tightness Urticaria					
/ ORAL		Vomiting					

Date:04/14/00ISR Number: 3488692-5Report Type:Expedited (15-DaCompany Report #10349645  
Age:29 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300		Pancreatitis Sedation	Health Professional	Serzone Tabs (Nefazodone Hcl)	PS		ORAL
	MILLIGRAM,							
	ORAL				Wellbutrin (Bupropion Hcl)	SS		
	75 MILLIGRAM							

Date:04/17/00ISR Number: 3488881-XReport Type:Expedited (15-DaCompany Report #A0112807A  
Age:47 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG / UNK		Myocardial Infarction Sudden Death	Health Professional Company Representative	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
	/ ORAL	4 WK			Fentanyl Celecoxib	C C		

Date:04/17/00ISR Number: 3488942-5Report Type:Expedited (15-DaCompany Report #A0117477A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Macular Degeneration Retinal Haemorrhage	Health Professional	Bupropion Hydrochloride Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
SEE TEXT/ORAL		YR					

Date:04/17/00ISR Number: 3488945-0Report Type:Expedited (15-DaCompany Report #A0116737A

Age:29 YR Gender:Female I/FU:I

Outcome	PT
Life-Threatening	Aggression
Hospitalization -	Coma
Initial or Prolonged	Confusional State
	Depression
	Ecchymosis
	Fall
	Fatigue
	Folliculitis
	Grand Mal Convulsion
	Head Injury
	Headache

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /SEE		Heart Rate Increased Influenza Like Illness Movement Disorder	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
TEXT/ ORAL		Muscle Rigidity Nausea Oxygen Saturation Decreased  Pallor					
TWICE PER DAY		Pupillary Reflex Impaired Respiratory Rate Decreased Speech Disorder		Amantadine (Formulation Unknown) (Amantadine)	SS		
		Tremor		Semisodium Valproate	C		

Date:04/17/00ISR Number: 3489012-2Report Type:Expedited (15-DaCompany Report #A0115993A  
Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Amblyopia	Foreign	Zyban Tablet	PS		ORAL
		Brain Neoplasm	Health Professional				

Date:04/17/00ISR Number: 3489024-9Report Type:Expedited (15-DaCompany Report #A0112392A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 150 MG/ TWICE Intervention to PER DAY/ ORAL Prevent Permanent Impairment/Damage		Abdominal Pain	Foreign	Zyban	PS		ORAL
		Arthralgia	Health Professional				
		Burning Sensation Decreased Activity Dermatitis Difficulty In Walking Dysphagia Eyelid Oedema		Mometasone Furoate	C		

Face Oedema  
Fatigue  
Feeling Cold  
Insomnia  
Oedema Peripheral  
Pain  
Pharyngolaryngeal Pain  
Pruritus  
Rash Erythematous  
Rash Pruritic  
Raynaud'S Phenomenon  
Sensation Of Heaviness  
Skin Discolouration  
Type Iv Hypersensitivity  
Reaction  
Urticaria  
Vasculitic Rash  
Vasculitis



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/17/00ISR Number: 3489029-8Report Type:Expedited (15-DaCompany Report #A0110688A  
Age:44 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - Initial or Prolonged 150 MG/ THREE TIMES PER DAY/ ORAL	Pneumonia Renal Failure Wegener'S Granulomatosis	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
			Olanzapine Alprazolam	C C		

Date:04/18/00ISR Number: 3489624-6Report Type:Direct Company Report #  
Age:60 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 PO BID Initial or Prolonged	Delusion Dizziness Dyspnoea Hyperventilation Hypoaesthesia Nausea Paraesthesia	Health Professional	Wellbutrin Premarin	PS C		ORAL

Date:04/18/00ISR Number: 3489728-8Report Type:Expedited (15-DaCompany Report #A0113640A  
Age:57 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/ PER DAY/ ORAL	Collapse Of Lung Emphysema Lung Injury Tobacco Abuse Weight Increased	Consumer	Wellbutrin (Bupropion Hydrochloride) Sumatriptan	PS		ORAL

Succinate

C

Date:04/18/00ISR Number: 3489729-XReport Type:Expedited (15-DaCompany Report #A0110912A  
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Arthralgia	Consumer	Zyban	PS		ORAL
Initial or Prolonged		Hypersensitivity Hypotension Pruritus Respiratory Arrest Swelling Syncope Urticaria					

Date:04/18/00ISR Number: 3489734-3Report Type:Expedited (15-DaCompany Report #A0106978A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Deafness	Health	Wellbutrin	PS		ORAL
100 MG/ THREE TIMES PER DAY/ ORAL		Oedema	Professional				
200 MG/ THREE				Carbamazepine Tablet (Carbamazepine)	SS		ORAL

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Freedom Of Information (FOI) Report

TIMES PER

DAY/ ORAL

Oestradiol C

Date:04/18/00ISR Number: 3489897-XReport Type:Expedited (15-DaCompany Report #A0117782A

Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Creatinine Increased Creatinine Renal Clearance Decreased	Foreign Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
100 MG /		Multiple Sclerosis					
TWICE PER DAY		Renal Failure Acute					
/ ORAL				Olanzapine Semisodium Valproate	C C		

Date:04/19/00ISR Number: 3490241-2Report Type:Expedited (15-DaCompany Report #A0116939A

Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged		Dermatitis Dysphagia Dyspnoea	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /		Face Oedema					
Disability		Liver Function Test					
TWICE PER DAY		Abnormal Mean Cell Haemoglobin Concentration Increased Tongue Oedema Urticaria					
/ ORAL							

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG / TWICE PER DAY /ORAL	Agitation Asthenia Decreased Appetite Dizziness Postural Dysphemia Electroencephalogram Abnormal Feeling Jittery Grand Mal Convulsion Hypersomnia Irritability Loss Of Consciousness Memory Impairment Sedation Tremor Vomiting	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/19/00ISR Number: 3490257-6Report Type:Expedited (15-DaCompany Report #A0117821A  
 Age:31 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Adrenal Disorder Amnesia Anaphylactic Shock Gastrointestinal Disorder	Consumer	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER DAY / ORAL	Panic Attack Tremor					

Date:04/19/00ISR Number: 3490272-2Report Type:Direct Company Report #  
 Age:38 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG BID PO, APPROX. 2 WEEKS	Anxiety Balance Disorder Medication Error Vision Blurred		Wellbutrin Sr 150 Mg	PS		ORAL

Date:04/19/00ISR Number: 3490919-0Report Type:Expedited (15-DaCompany Report #JRFUSA2000000083  
 Age:30 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Hospitalization - TRANSDERMAL 50 MCH/H, 1 Initial or Prolonged IN 72 HOUR(S), TRANSD	Accidental Overdose Alcohol Poisoning Cardiomegaly Coma Drug Level Above Therapeutic	Consumer Health Professional	Duragesic (Patch) (Fentanyl)	PS		
			Nortriptyline			

ORAL	Medication Error	(Nortriptyline)	SS	ORAL
ORAL		Alprazolam (Alprazolam)	SS	ORAL
PRN, ORAL		Hydrocodone (Hydrocodone)	SS	ORAL
ORAL		Wellbutrin (Amfebutamone Hydrochloride)	SS	ORAL
ORAL		Alcohol (Ethanol)	SS	ORAL
		Stimulants (Psychostimulants And Nootropics)	SS	
		Diazepam (Diazepam)	SS	
		Chlordiazepoxide (Chlordiazepoxide)	SS	

Date:04/20/00ISR Number: 3490806-8Report Type:Expedited (15-DaCompany Report #A0112668A  
Age:26 YR Gender:Female I/FU:I

Outcome	PT
Required	Complex Partial Seizures
Intervention to	Confusional State
Prevent Permanent	Convulsion
Impairment/Damage	Disorientation
	Dizziness
	Grand Mal Convulsion
	Headache
	Hyperhidrosis
	Hypoglycaemia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypokalaemia Migraine Nausea					
150 MG / TWICE PER DAY / ORAL		Nervousness Panic Attack Visual Acuity Reduced  Transiently	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
				Tri-Levlen	C		

Date:04/21/00ISR Number: 3491038-XReport Type:Expedited (15-DaCompany Report #B0080118A  
Age:59 YR Gender:Male I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 TABLET / TWICE PER DAY/ ORAL		Hospitalization - Initial or Prolonged  Cardiovascular Disorder  Respiratory Failure	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:04/21/00ISR Number: 3491041-XReport Type:Expedited (15-DaCompany Report #A0117940A  
Age:47 YR Gender:Male I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL		Hospitalization - Initial or Prolonged  Chest Pain  Dizziness  Dyspnoea Eyelid Oedema	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
				Carisoprodol	C		
				Diazepam	C		

Hypersensitivity  
Irritability  
Palpitations  
Pruritus  
Swelling  
Toxic Shock Syndrome  
Urticaria

Date:04/21/00ISR Number: 3491920-3Report Type:Expedited (15-DaCompany Report #A0118416A

Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Ageusia Anosmia Face Oedema	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG TWICE PER DAY ORAL		Hypoaesthesia Pharyngeal Oedema Pyrexia Rash Erythematous Rash Pruritic					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/24/00ISR Number: 3491515-1Report Type:Expedited (15-DaCompany Report #JRFUSA2000000083

Age:30 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Death	Accidental Overdose	Consumer	Duragesic (Patch)			
Hospitalization -	Alcohol Poisoning	Health	(Fentanyl)	PS		
TRANSDERMAL	50 MCG/H, 1					
Initial or Prolonged	Coma	Professional				
IN 72						
	Drug Level Above					
HOUR(S),						
	Therapeutic					
TRANSD/200						
	Medication Error					
MCG/H						
			Nortriptyline			
			(Nortriptyline)	SS		ORAL
ORAL						
			Alprazolam			
			(Alprazolam)	SS		ORAL
ORAL						
			Hydrocodone			
			(Hydrocodone)	SS		ORAL
PRN, ORAL						
			Wellbutrin			
			(Amfebutamone			
			Hydrochloride)	SS		ORAL
ORAL						
			Alcohol (Alcohol)	SS		ORAL
ORAL						
			Diazepam (Diazepam)	SS		
			Chlordiazepoxide			
			(Chlordiazepoxide)	SS		
			Phenylpropanolamine			
			(Phenylpropanolamine			
			)	SS		
			Ephedrine			
			(Ephedrine)	SS		
			Pseudoephedrine			
			(Pseudoephedrine)	SS		

Date:04/24/00ISR Number: 3491575-8Report Type:Expedited (15-DaCompany Report #A0118209A

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Haemolysis Hyperkalaemia	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / ORAL				Nicotine	C		

Date:04/24/00ISR Number: 3491924-0Report Type:Expedited (15-DaCompany Report #B0079430A  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage 150 MG SEE		Drug Interaction International Normalised Ratio Decreased Prothrombin Time  Prolonged	Foreign Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
TEXT ORAL				Warfarin Sodium	C		
				Moduretic	C		
				Simvastatin	C		
				Paroxetine	C		
				Zolpidem	C		
				Alprazolam	C		



Gait Disturbance	Amantadine	
Neurotoxicity	(Formulation	
Restlessness	Unknown)	
Tremor	(Amantadine)	SS
Vertigo		

Date:04/28/00ISR Number: 3494456-9Report Type:Expedited (15-DaCompany Report #A0117770A  
Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Agitation Confusional State Coordination Abnormal Dizziness Drug Interaction	Literature Health Professional	Bupropion Hydrochloride Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL	Gait Disturbance Neurotoxicity Restlessness Tremor Vertigo		Amantadine (Formulation Unknown) (Amantadine)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/01/00ISR Number: 3494992-5Report Type:Expedited (15-DaCompany Report #B0079458A

Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Angina Pectoris Palpitations	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150							
MG/UNK/ORAL							

				Thyroxine Sodium Tablet (Levothyroxine Sodium)	SS		ORAL
.1							
MG/UNKNOWN/OR							

AL

Date:05/01/00ISR Number: 3495043-9Report Type:Expedited (15-DaCompany Report #A0110705A

Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Grand Mal Convulsion Insomnia	Foreign Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
1509 MG/							

TWICE PER DAY

/ ORAL

Doxepin Hydrochloride	C
Herbal Medication	C
Ethanol	C

Date:05/01/00ISR Number: 3495117-2Report Type:Expedited (15-DaCompany Report #A0110103A

Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to 150 MG /TWICE Prevent Permanent PER DAY/ ORAL Impairment/Damage		Colitis Ischaemic Rectal Haemorrhage	Study Health	Zyban (Bupropion Hydrochloride)	PS		ORAL
		Syncope	Professional				
				Oral Contraceptive Tablet (Oral Contraceptive)	SS		ORAL
150 MF /TWICE PER DAY/ ORAL ORAL				Quinapril Hydrochloride	SS		ORAL

Date:05/01/00ISR Number: 3495119-6Report Type:Expedited (15-DaCompany Report #A0104240A  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 400 MG /PER Hospitalization - DAY/ ORAL Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage		Aggression Child Abuse Mania Suicidal Ideation	Study Health Professional	Wellbutrin Paracetamol	PS C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/01/00ISR Number: 3495122-6Report Type:Expedited (15-DaCompany Report #A0113640A

Age:57 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG /PER DAY/ ORAL	Collapse Of Lung Condition Aggravated Emphysema Lung Injury Smoker Weight Increased	Consumer	Wellbutrin Tablet-Controlled Release  Sumatriptan Succinate Vicodin Cortisone Procaine Hydrochloride	PS    C C C C		ORAL

Date:05/01/00ISR Number: 3495142-1Report Type:Expedited (15-DaCompany Report #A0117127A

Age:22 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG SEE TEXT ORAL 13 DAY	Angioneurotic Oedema Arthralgia Escherichia Infection Joint Swelling Livedo Reticularis Myalgia Pruritus Pyrexia Serum Sickness Urinary Tract Infection Urticaria White Blood Cell Count Increased	Foreign Literature Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)  Oral Contraceptive	PS   C		ORAL

Date:05/01/00ISR Number: 3495164-0Report Type:Expedited (15-DaCompany Report #B0079430A

Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to Prevent Permanent Impairment/Damage		Drug Interaction International Normalised Ratio Decreased Prothrombin Time Prolonged	Foreign Health Professional	Bupropion Hydrochloride Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG	SEE						
TEXT	ORAL			Warfarin Sodium (Formulation Unknown) (Warfarin Sodium)	SS		ORAL
6.25 MG	PER						
DAY	ORAL			Moduretic Simvastatin Paroxetine Zolpidem Alprazolam	C C C C C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/01/00ISR Number: 3496404-4Report Type:Periodic  
 Age:44 YR Gender:Male I/FU:I

Company Report #PRIUSA1999006408

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - Initial or Prolonged ORAL Required Intervention to Prevent Permanent 150 MG, 2 IN Impairment/Damage 1 DAILY, ORAL	Grand Mal Convulsion	Health Professional	Ultram (50 Mg Tablet) (Tramadol Hydrochloride)	PS		ORAL
			Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL
			Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C		

Date:05/02/00ISR Number: 3495398-5Report Type:Direct  
 Age:8 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required Intervention to 100MG BID Prevent Permanent ORAL Impairment/Damage	Arthralgia Arthropathy Conjunctivitis Dermatitis Erythema Multiforme Myalgia Serum Sickness Urticaria		Bupropion-Wellbutrin 100mg	PS		ORAL
			Adderall	C		
			Dexedrine	C		

Date:05/03/00ISR Number: 3495646-1Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Arthralgia Arthritis		Wellbutrin Adderall	PS C		

Dyspnoea  
 Erythema Multiforme  
 Laryngeal Oedema  
 Swelling  
 Urticaria

Ritalin

C

Date:05/03/00ISR Number: 3495834-4Report Type:Direct  
 Age:38 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG BID PO	Convulsion Dyskinesia	Health Professional	Bupropion (Wellbutrin Sr)	PS		ORAL
		Hypoglycaemia		Human Insulin (Humalog)	SS		
	SUBCUTANEOUS	HUMALOG 75U					
	SQ			Human Insulin (Humulin U)	SS		
	SUBCUTANEOUS	HUMALIN U 40					
	U SQ			Prevacid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/03/00ISR Number: 3496212-4Report Type:Direct  
Age:41 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG BID 1 WK Initial or Prolonged	Confusional State	Health	Wellbutrin	PS		
	Eye Rolling	Professional	Levaquin	C		
	Grand Mal Convulsion		Colace	C		
	Lethargy		Xanax	C		
			Toradol	C		
			Demerol	C		
			Zoloft	C		
			Bellergal	C		
			Depakote	C		
			Serzone	C		

Date:05/03/00ISR Number: 3496527-XReport Type:Expedited (15-DaCompany Report #A0116863A  
Age:50 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Required Intervention to Prevent Permanent 150 MG / Impairment/Damage TWICE PER DAY/ ORAL	Completed Suicide	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
			Ethanol (Formulation Unknown) (Alcohol)	SS		
			Aspirin (Formulation Unknown) (Aspirin)	SS		

Date:05/05/00ISR Number: 3497095-9Report Type:Expedited (15-DaCompany Report #A0114122A  
Age:16 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Aphasia Blepharospasm	Health Professional	Wellbutrin Tablet-Controlled			

Disability Required 150 MG / Intervention to TWICE PER DAY Prevent Permanent / ORAL Impairment/Damage	Dystonia Oculogyration  Trismus	Release (Bupropion Hydrochloride)	PS	ORAL
		Semisodium Valproate	C	

Date:05/05/00ISR Number: 3497096-0Report Type:Expedited (15-DaCompany Report #A0119185A  
Age:58 YR Gender:Female I/FU:I

Outcome Dose Hospitalization - Initial or Prolonged	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / UNK  / ORAL		Decreased Activity Multiple Sclerosis	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
				Prednisone	C		

Date:05/05/00ISR Number: 3497302-2Report Type:Expedited (15-DaCompany Report #A0110940A  
Age:44 YR Gender:Male I/FU:F

Outcome Death	PT Alcohol Poisoning Completed Suicide
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Injury Toxicologic Test Abnormal				
Dose	Duration		Report Source	Product	Role	Manufacturer
			Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS	
150 MG/ORAL						ORAL

Date:05/05/00ISR Number: 3497883-9Report Type:Direct Company Report #  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health Professional	Zyban	PS		ORAL
Other		Rash Papular					
PO 1 QD X 3							
		Rash Pruritic	Professional				
THEN 2 QD							

Date:05/08/00ISR Number: 3497771-8Report Type:Expedited (15-DaCompany Report #WAES 00032541  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Tab Vioxx 25 Mg	PS		ORAL
Other		Dermatitis	Health Professional				
25							
		Dizziness					
MG/DAILY/PO							
		Dysphagia	Professional	Wellbutrin Sr 150 Mg	SS		ORAL
150 MG/BID/PO							
		Pharyngeal Oedema					
		Pharyngitis					
		Pyrexia					
		Urticaria					

Date:05/08/00ISR Number: 3498215-2Report Type:Expedited (15-DaCompany Report #9915007  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Required	Anxiety	Consumer	Marax Tablets	PS	ORAL
ORAL					
Intervention to	Depression	Health	Wellbutin	SS	
Prevent Permanent	Depression Suicidal	Professional	Azmacort	C	
Impairment/Damage	Drug Ineffective		Ventolin	C	
	Gastric Disorder		Vancenase	C	
	Hypertension		Potassium	C	
	Irritable Bowel Syndrome		Prilosec	C	
	Suicidal Ideation		Tagamet Hb	C	
	Weight Decreased				

Date:05/08/00ISR Number: 3498277-2Report Type:Expedited (15-DaCompany Report #A0117940A  
Age:47 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Asthenia
Initial or Prolonged	Blister
	Bronchospasm
	Burning Sensation
	Chest Pain
	Dizziness
	Drug Hypersensitivity
	Dyspnoea
	Eye Irritation
	Eyelid Oedema
	Haemorrhage
	Irritability
	Palpitations

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Pruritus Sunburn Swelling	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
TWICE PER DAY		Toxic Shock Syndrome Urticaria					
/ ORAL				Carisoprodol Diazepam	C C		

Date:05/08/00ISR Number: 3498279-6Report Type:Expedited (15-DaCompany Report #A0119044A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abortion Induced	Study	Bupropion			
Congenital Anomaly		Anal Atresia	Health	Hydrochloride Tablet			
		Calcinosis	Professional	(Bupropion Hydrochloride)	PS		ORAL
ORAL		Cleft Palate					
		Complications Of Maternal Exposure To Therapeutic Drugs		Influenza Vaccine	C		
		Mesomelia					
		Multiple Congenital Abnormalities					
		Pierre Robin Syndrome					

Date:05/09/00ISR Number: 3497779-2Report Type:Direct Company Report #  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG TWICE		Stevens-Johnson Syndrome		Wellbuterin 300mg	PS		ORAL
Initial or Prolonged DAY ORAL				Vitamin B			

Date:05/10/00ISR Number: 3498196-1Report Type:Direct  
 Age:18 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG PO BID		Arthralgia Pharyngeal Oedema Urticaria	Health Professional	Bupropion Sr (Zyban)	PS		ORAL

Date:05/10/00ISR Number: 3504424-6Report Type:Periodic  
 Age:51 YR Gender:Male I/FU:I

Company Report #990926.01

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2.5MG QD, ORAL		Cough	Health Professional	Indapamide	PS	Mylan Pharmaceuticals Inc	ORAL
2 PUFFS UD, ORAL				Allopurinol 300mg Beconase Glaxo Wellcome	SS SS		ORAL
50MG QD, ORAL				Cozaar 50mg Merck Glucophage Bristol	SS	Merck	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

1 GRAM TID, ORAL	Myers Squibb	SS	Bristol Myers Squibb	ORAL
10MG QD, ORAL	Lipitor 10mg Parke-Davis	SS	Parke-Davis	ORAL
200MG BID, ORAL	Rezulin Parke-Davis	SS	Parke-Davis	ORAL
400MG TID, ORAL	Trental 400mg Hoechst Marion Roussel	SS	Hoechst Marion Roussel	ORAL
150MG QD, ORAL	Wellbutrin 150mg Glaxo Wellcome	SS	Glaxo Wellcome	ORAL
10MG BID, ORAL	Glipizide 10mg	SS		ORAL
325MG TID, ORAL	Quinine 325mg	SS		ORAL

Date:05/12/00ISR Number: 3499875-2Report Type:Expedited (15-DaCompany Report #A0117940A

Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Asthenia Bronchospasm Burning Sensation	Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE PER DAY/ORAL		Chest Pain					
		Dermatitis		Carisoprodol	C		
		Dizziness		Diazepam	C		
		Dyspnoea					

Dyspnoea Exertional  
 Erythema  
 Eye Irritation  
 Eyelid Oedema  
 Hypersensitivity  
 Irritability  
 Palpitations  
 Periorbital Oedema  
 Photosensitivity Reaction  
 Pruritus  
 Swelling  
 Toxic Shock Syndrome  
 Urticaria

Date:05/12/00ISR Number: 3499878-8Report Type:Expedited (15-DaCompany Report #A0112039A  
 Age:43 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Convulsion Loss Of Consciousness Memory Impairment	Health Professional Company	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE  PER DAY/ORAL	Pain  Syncope Tongue Disorder	Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/12/00ISR Number: 3499897-1Report Type:Expedited (15-DaCompany Report #D0006860A  
Age:36 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - ORAL	Angina Pectoris Cardiac Disorder	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged Disability	Coronary Artery Surgery Influenza	Professional				

Date:05/12/00ISR Number: 3499898-3Report Type:Expedited (15-DaCompany Report #A0119608A  
Age:16 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL	Bronchospasm Dehydration Difficulty In Walking Dyspnoea Oedema Pharyngolaryngeal Pain Pruritus Urticaria	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:05/15/00ISR Number: 3500475-6Report Type:Expedited (15-DaCompany Report #A0115993A  
Age:60 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other ORAL	Amblyopia Brain Neoplasm	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/15/00ISR Number: 3500484-7Report Type:Expedited (15-DaCompany Report #A0069088A  
Age:65 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Choking	Professional				
TWICE PER DAY		Convulsion	Other				
/ ORAL		Feeling Jittery		Simvastatin	C		
		Thirst		Aspirin	C		
		Tremor					

Date:05/15/00ISR Number: 3501204-2Report Type:Expedited (15-DaCompany Report #00P-163-0089665-00(0)  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Chest Pain	Health	Cylert	PS	Abbott Laboratories	
Intervention to		Drug Interaction	Professional			Pharmaceutical	
Prevent Permanent						Products Div	ORAL
PER ORAL							
Impairment/Damage				Amfebutamone			
				Hydrochloride			
				(Amfebutamone			
				Hydrochloride)	SS		ORAL
400 MG, 1 IN							
1 D, PER ORAL							
				Methylphenidate			
				Hydrochloride			
				(Methylphenidate			
				Hydrochloride)	SS		ORAL
5 MG, 2 IN 1							
D, PER ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/15/00ISR Number: 3501216-9Report Type:Expedited (15-DaCompany Report #A0120078A  
 Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / PER Initial or Prolonged DAY / ORAL		Bipolar Disorder	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Other		Bipolar I Disorder	Consumer				
		Convulsion		Clonazepam	C		
		Disorientation		Paroxetine			
		Disturbance In Attention		Hydrochloride	C		
		Mood Swings		Perphenazine	C		
		Visual Disturbance					

Date:05/16/00ISR Number: 3501192-9Report Type:Expedited (15-DaCompany Report #200011497HMRI  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Health Professional	Lasix	PS	Aventis Pharmaceuticals Inc	
				Amfebutamone Hydrochloride (Wellbutrin - Slow Release)	SS		

Date:05/16/00ISR Number: 3501780-XReport Type:Periodic Company Report #HQ0846901FEB2000  
 Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional	Effexor Xr	PS	Wyeth Ayerst Laboratories Inc	ORAL
37.5 MG 1X PER 1 DAY, ORAL				Prozac	SS		ORAL
20 MG 2X PER 1 DAY ORAL							

14 CAPSULES	Effexor Xr	SS	ORAL
OVERDOSE			
AMOUNT, ORAL			
OVERDOSE	Prozac	SS	ORAL
AMOUNT			
UNKNOWN, ORAL			
150 MG 1X PER	Wellbutrin	SS	ORAL
1 DAY, ORAL			
OVERDOSE	Wellbutrin	SS	ORAL
AMOUNT			
UNKNOWN, ONE			
TIME, ORAL			

Date:05/17/00ISR Number: 3561205-5Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #S00-USA-00214-01

Outcome PT  
Anxiety  
Ejaculation Disorder  
Emotional Disorder  
Headache  
Hyperhidrosis  
Libido Decreased  
Nervousness  
Paraesthesia  
Sedation

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tooth Disorder  
Vasodilatation

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Consumer	Celexa	PS	Forest Laboratories Inc	ORAL
15 MG QD PO			Celexa	SS		ORAL
10 MB BID PO			Celexa	SS		ORAL
10 MG QD PO			Celexa	SS		ORAL
5 MG QD PO			Wellbutrin	SS		ORAL
75 MG TID PO			Wellbutrin	SS		ORAL
75 MG QD PO						

Date:05/17/00ISR Number: 3561232-8Report Type:Periodic Company Report #S00-USA-00234-01  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accident	Health Professional	Celexa	PS	Forest Laboratories Inc	
				Wellbutrin	SS		

Date:05/18/00ISR Number: 3501390-4Report Type:Direct Company Report #USP 53031  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxo Wellcome	
				Wellbutrin (Bupropion Hydrochloride)	SS	Glaxo Wellcome	

Date:05/18/00ISR Number: 3501462-4Report Type:Direct Company Report #  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300MG DAILY Disability ORAL Required Intervention to Prevent Permanent Impairment/Damage		Cerebrovascular Accident Convulsion	Consumer	Zyban / 150mg / Glaxo	PS	Glaxo	ORAL

Date:05/18/00ISR Number: 3501483-1Report Type:Direct Company Report #  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG BID PO		Swelling Urticaria	Health Professional	Zyban 150mg Bid	PS		ORAL

Date:05/18/00ISR Number: 3502211-6Report Type:Expedited (15-DaCompany Report #M0316-2000  
Age:35 YR Gender:Female I/FU:I

Outcome  
Life-Threatening  
Hospitalization -  
Initial or Prolonged  
Required  
Intervention to

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Prevent Permanent Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
560 MG PO		Abnormal Behaviour Angina Pectoris	Health Professional	Remeron	PS	Organon Inc Sub Akzona Inc	ORAL
900 MG PO		Aspartate	Other	Wellbutrin	SS		ORAL
		Aminotransferase Increased		Librium	SS		
		Blood Creatine		Benzonatate	C		
		Phosphokinase Increased		Cefuroxime	C		
		Cardiac Disorder		Docusate	C		
		Cardiomyopathy		Doxycycline	C		
		Chest Pain		Kcl	C		
		Dyspnoea		Levaquin	C		
		Electrocardiogram Qt Corrected Interval Prolonged		Methadone	C		
		Electrocardiogram Qt Prolonged		Robitussin	C		
		Electrocardiogram T Wave Inversion		Tylenol	C		
		Hypokalaemia					
		Mitral Valve Incompetence					
		Myocardial Ischaemia					
		Overdose					
		Pneumonia					
		Toxicologic Test Abnormal					
		Tricuspid Valve Incompetence					

Date:05/19/00ISR Number: 3501934-2Report Type:Expedited (15-DaCompany Report #A0110912A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening ORAL		Anaphylactic Reaction	Health Professional	Zyban Tablet	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent		Arthralgia Dyspnoea Hypersensitivity Hypotension Pruritus					

Impairment/Damage      Respiratory Arrest  
Swelling  
Syncope  
Urticaria

Date:05/19/00ISR Number: 3501935-4Report Type:Expedited (15-DaCompany Report #A0116737A  
Age:29 YR    Gender:Female      I/FU:F

Outcome	PT
Life-Threatening	Aggression
Hospitalization -	Amnesia
Initial or Prolonged	Coma
	Confusional State
	Depression
	Drug Level Below
	Therapeutic
	Ecchymosis
	Fatigue
	Folliculitis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 150 MG / SEE TEXT / ORAL		Grand Mal Convulsion Head Injury Headache	Health				
		Heart Rate Increased	Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
		Influenza Like Illness					
TWICE PER DAY		Movement Disorder Muscle Rigidity Nausea Oxygen Saturation Decreased Pallor Pupillary Reflex Impaired Respiratory Rate Decreased Speech Disorder		Amantadine (Formulation Unknown) (Amantadine)	SS		
				Semisodium Valproate	C		

Date:05/19/00ISR Number: 3502241-4Report Type:Expedited (15-DaCompany Report #A0113561A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY / ORAL		Ageusia Angioneurotic Oedema Anosmia Face Oedema Paraesthesia Pharyngeal Oedema Pyrexia Rash Pruritic Tongue Oedema	Foreign Health Professional Company Representative	Zyban Salbutamol Sulphate	PS C	Glaxo Wellcome Inc	ORAL

Date:05/19/00ISR Number: 3502244-XReport Type:Expedited (15-DaCompany Report #A0118416A  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY  / ORAL	Ageusia Anosmia  Dermatitis  Face Oedema Paraesthesia Pharyngeal Oedema Pruritus Pyrexia	Foreign  Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
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Date:05/23/00		ISR Number: 3503758-9		Report Type:Direct		Company Report #	
Age:42 YR	Gender:Male	I/FU:I					
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash Pruritic Urticaria	Health Professional	Zyban 150 Mg (Glaxo-Wellcome)	PS	Glaxo-Wellcome	
150MG QD X 3							
THEN BID				Biaxin Ibuprofen Robitussin	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/23/00ISR Number: 3503844-3Report Type:Expedited (15-DaCompany Report #A0119823A  
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/PER Initial or Prolonged DAY/ORAL		Chest Discomfort	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Dyspnoea					
		Hypoxia		Metoprolol Succinate	C		
		Palpitations		Amlodipine	C		
		Ventricular Extrasystoles		Isosorbide Mononitrate	C		

Date:05/24/00ISR Number: 3504498-2Report Type:Direct Company Report #  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG BID		Partial Seizures		Bupropion	PS		
				Prednisone	C		
				Flovent	C		
				Lanoxin	C		
				Lasix	C		
				Kcl	C		
				Singulair	C		
				Atrovent	C		
				Albuterol	C		

Date:05/24/00ISR Number: 3504499-4Report Type:Direct Company Report #  
 Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG BID		Erythema Rash Papular		Wellbutrin-Sr (Glaxo-Wellcome)	PS	Glaxo -Wellcome	
				Deconamine	C		
				Allegra	C		

Date:05/24/00ISR Number: 3504501-XReport Type:Direct  
Age:61 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dermatitis Urticaria		Zyban 150mg (Glaxo-Wellcome)	PS	Glaxo-Wellcome	ORAL
150MG PO QD X							
3 THEN BID				Ambien Aspirin Nicoderm Patch	C C C		

Date:05/24/00ISR Number: 3504531-8Report Type:Expedited (15-DaCompany Report #A0117763A  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Ageusia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Decreased Appetite	Professional				
PER DAY/ ORAL		Headache	Company Representative	Aspirin Armour Thyroid Terbinafine	C C C		



ORAL	Infection	Flagyl	SS	ORAL
300.00 MG	Lymphocyte Count	Wellbutrin	SS	ORAL
TOTAL DIALY	Increased			
ORAL	Malaise			
	Oral Candidiasis	Azidothymidine	C	
	Tongue Discolouration	Viracept	C	
		Combivir	C	
		Methamphetamine	C	
		Prevacid	C	
		Zyrtec	C	

Date:05/26/00ISR Number: 3566696-1Report Type:Periodic Company Report #10868-AR  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia	Consumer	Bupropion Hcl	PS	Teva Pharmaceuticals	
280 MG		Hostility				Usa Inc	
DECREASED TO		Hyperhidrosis					
75 MG BID		Vasodilatation					
				Acyclovir	C		
				Allopurinol	C		
				Celexa	C		
				Remeron	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/26/00ISR Number: 3566702-4Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #10872-AR

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urinary Retention	Health Professional	Bupropion Hcl	PS	Teva Pharmaceuticals Usa Inc	
75 MG BID							

Date:05/26/00ISR Number: 3566704-8Report Type:Periodic  
 Age:13 YR Gender:Male I/FU:I

Company Report #10878-AR

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Headache	Consumer	Bupropion Hcl	PS	Teva Pharmaceuticals Usa Inc	
150 MG AM, 75 MG NOON AND HS		Hostility					
				Bupropion Hydrochloride Tablets, 75 Mg	SS		
150 MG AM, NOON AND HS							

Date:05/30/00ISR Number: 3505698-8Report Type:Direct  
 Age:73 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 150MG BID		Eye Rolling		Zyban 150mg	PS		ORAL
Intervention to ORAL		Hallucination					
Prevent Permanent Impairment/Damage		Muscle Rigidity Tremor		Lipitor	C		
				Prevacid	C		
				Hytrin	C		
				Zoloft	C		
				Asa	C		
				Clonidine	C		

Date:05/30/00ISR Number: 3506782-5Report Type:Expedited (15-DaCompany Report #A0120940A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache Neck Pain	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Night Sweats					
TWICE PER DAY		Pain					
/ ORAL		Paraesthesia Visual Acuity Reduced					

Date:05/30/00ISR Number: 3506783-7Report Type:Expedited (15-DaCompany Report #A0120791A  
Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Agitation	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Delusion	Professional				
Initial or Prolonged		Psychotic Disorder					
THREE TIMES		Self Mutilation		Baclofen Oxycodone Hydrochloride	C C		
PER DAY ORAL	6	MON					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/30/00ISR Number: 3506784-9Report Type:Expedited (15-DaCompany Report #A0120487A  
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eye Disorder Skin Degenerative Disorder	Literature Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / UNKNOWN / ORAL		Stevens-Johnson Syndrome Toxic Epidermal Necrolysis					

Date:05/30/00ISR Number: 3506834-XReport Type:Periodic Company Report #3654/11757  
 Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anorexia Dizziness Dyspnoea	Consumer Company Representative	Depo-Medrol	PS	Pharmacia And Upjohn Co	
INTRAMUSCULAR DOSE;IM 300 MG/DAY	20 MG-1	Feeling Jittery Headache Mood Swings Nervousness Pain In Extremity Photophobia Restlessness Sedation Vomiting		Wellbutrin Celexa Risperdol	SS SS C		

Date:05/31/00ISR Number: 3507206-4Report Type:Periodic Company Report #A0109497A  
 Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspnoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							

PER DAY/ ORAL

Hypersensitivity

Joint Swelling  
Oedema Peripheral  
Pharyngeal Oedema  
Rash Generalised  
Throat Tightness  
Urticaria

Date:05/31/00ISR Number: 3507207-6Report Type:Periodic  
Age:46 YR Gender:Female I/FU:F

Company Report #A0107904A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150 MG/ TWICE Hospitalization - PER DAY/ ORAL Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage		Grand Mal Convulsion  Pyrexia	Health  Professional	Zyban   Conjugated Estrogens Medroxyprogesterone Ace.	   C  C	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3507208-8Report Type:Periodic  
Age:49 YR Gender:Male I/FU:F

Company Report #A0060628A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG/TWICE	Back Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged PER DAY/ORAL	Chest Pain					
	Decreased Appetite		Isosorbide			
	Dizziness		Mononitrate	C		
	Drug Ineffective		Metoprolol Succinate	C		
	Dry Mouth		Lansoprazole	C		
	Dysgeusia		Aspirin	C		
	Dyspepsia					
	Dyspnoea					
	Headache					
	Insomnia					

Date:05/31/00ISR Number: 3507209-XReport Type:Periodic  
Age:75 YR Gender:Female I/FU:I

Company Report #A0118622A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG/ ORAL	Convulsion	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Triamcinolone Acetonide	C		

Date:05/31/00ISR Number: 3507210-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0116791A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG/TWICE	Dizziness	Health	Zyban	PS	Glaxo Wellcome Inc	
Initial or Prolonged PER DAY/		Professional				
			Atorvastatin Calcium	C		

Date:05/31/00ISR Number: 3507211-8Report Type:Periodic Company Report #A0116361A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ TWICE		Anaphylactic Reaction	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged PER DAY/ ORAL		Dermatitis					
		Hypersensitivity					

Date:05/31/00ISR Number: 3507212-XReport Type:Periodic Company Report #A0115677A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dermatitis Urticaria	Health Professional Company Representative	Zyban	PS	Glaxo Wellcome Inc	

Date:05/31/00ISR Number: 3507213-1Report Type:Periodic Company Report #A0114547A  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG/ PER DAY/ ORAL		Convulsion	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3507214-3Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0114398A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
Other		Convulsion					
150 MG/		Tremor	Professional				
SINGLE DOSE/			Company				
ORAL			Representative				

Date:05/31/00ISR Number: 3507215-5Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #A0114203A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Other		Convulsion					
150 MG/ TWICE		Dry Mouth					
PER DAY/ ORAL		Insomnia					

Date:05/31/00ISR Number: 3507216-7Report Type:Periodic  
Age:65 YR Gender:Female I/FU:I

Company Report #A0113908A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	
Other		Convulsion					

Date:05/31/00ISR Number: 3507217-9Report Type:Periodic  
Age:25 YR Gender:Female I/FU:I

Company Report #A0111695A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Other		Dermatitis					
150 MG/TWICE		Eyelid Oedema					
PER DAY/ ORAL		Face Oedema		Oral Contraceptive	C		

Hypersensitivity  
Oedema Peripheral  
Pain  
Pharyngeal Oedema  
Urticaria

Date:05/31/00ISR Number: 3507218-0Report Type:Periodic Company Report #A0113182A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
Other		Convulsion					
150 MG/ ORAL		Drug Toxicity	Professional				

Date:05/31/00ISR Number: 3563220-4Report Type:Periodic Company Report #A0115654A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Diarrhoea					
150 MG / ORAL				Nicotine (Nicotine)	C		

Date:05/31/00ISR Number: 3563225-3Report Type:Periodic Company Report #A0115703A  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Headache Suicidal Ideation					
150 MG /							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

TWICE PER

DAY/ ORAL

Hypericum

C

Date:05/31/00ISR Number: 3563227-7Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0115704A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /TWICE		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL		Irritability					

Date:05/31/00ISR Number: 3563229-0Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0115772A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Euphoric Mood					
/ ORAL		Feeling Abnormal					
		Feeling Drunk					
		Nervousness					

Date:05/31/00ISR Number: 3563233-2Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #A0115784A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY /		Joint Swelling					
ORAL		Pruritus					

Date:05/31/00ISR Number: 3563235-6Report Type:Periodic Company Report #A0115837A  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ ORAL	3 WK	Hypersensitivity	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Rash Generalised					

Date:05/31/00ISR Number: 3563236-8Report Type:Periodic Company Report #A0115873A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG / ORAL		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563237-XReport Type:Periodic Company Report #A0115879A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ ORAL	1 DAY	Vision Blurred	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563240-XReport Type:Periodic Company Report #A0115882A  
Age:73 YR Gender:Female I/FU:I

Outcome	PT
	Dissociation
	Dizziness
	Headache

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Nausea Vertigo				
Dose	Duration		Report Source	Product	Role	Manufacturer
150 MG /			Health	Zyban	PS	Glaxo Wellcome Inc
SINGLE DOSE/			Professional			
ORAL						Route
						ORAL

Date:05/31/00ISR Number: 3563242-3Report Type:Periodic Company Report #A0115926A  
 Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Rash Generalised	Consumer	Zyban	PS	Glaxo Wellcome Inc
150 MG /						Route
TWICE PER						ORAL
DAY/ ORAL						

Date:05/31/00ISR Number: 3563244-7Report Type:Periodic Company Report #A0115927A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Arthralgia	Consumer	Zyban	PS	Glaxo Wellcome Inc
ORAL						Route
		Difficulty In Walking				ORAL
		Dyspnoea				
		Dyspraxia				
		Eyelid Oedema				
		Joint Swelling				
		Pain				
		Pruritus				
		Rheumatoid Arthritis				
		Serum Sickness				
		Swelling				
		Urticaria				

Date:05/31/00ISR Number: 3563245-9Report Type:Periodic Company Report #A0115928A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphagia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Insomnia					
TWICE PER		Lymphadenopathy					
DAY/ ORAL		Tinnitus					
		Urticaria					

Date:05/31/00ISR Number: 3563248-4Report Type:Periodic Company Report #A0115929A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL				Nicotine Patch (Nicotine)	SS		
TRANSDERMAL	21 MG /						
TRANSDERMAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3563250-2Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0115932A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL		Agitation Nervousness	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563251-4Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #A0115949A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Feeling Abnormal	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563252-6Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #A0115953A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Agitation Dizziness Eating Disorder Feeling Abnormal Headache Insomnia Malaise Nausea Psychomotor Hyperactivity Tremor	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563253-8Report Type:Periodic  
Age:52 YR Gender:Male I/FU:I

Company Report #A0115955A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Rash Generalised	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER		Rash Pruritic					
DAY/ ORAL							

Date:05/31/00ISR Number: 3563254-XReport Type:Periodic Company Report #A0115958A  
 Age:55 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /ORAL		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Dry Skin		Simvastatin	C		
		Dysphagia		Nadolol	C		
		Mucosal Dryness		Aspirin	C		

Date:05/31/00ISR Number: 3563255-1Report Type:Periodic Company Report #A0115960A  
 Age:32 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Galactorrhoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3563256-3Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0116037A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL		Amnesia Chest Pain Conjunctival Irritation Hallucination, Visual Mental Impairment Paraesthesia Sensation Of Heaviness	Consumer	Zyban   Conjugated Estrogens	PS   C	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563257-5Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #A0116067A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG (PER DAY) ORAL		Decreased Appetite Disturbance In Attention Malaise Tobacco Abuse	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563258-7Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #A0116070A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG (TWICE PER DAY) ORAL		Dermatitis Drug Ineffective Erythema Irritability Scar	Consumer	Bupropion Hydrochloride   Glucophage	PS   C	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563260-5Report Type:Periodic Company Report #A0116090A  
Age:30 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG (TWICE PER DAY) ORAL		Hyperaesthesia Oedema Peripheral Pain In Extremity Pruritus Rash Erythematous Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563265-4Report Type:Periodic Company Report #A0116091A  
Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL	1 WK	Erythema Neoplasm Skin Rash Pruritic	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563268-XReport Type:Periodic Company Report #A0116092A  
Age:45 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG (TWICE PER		Dysgeusia Parosmia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

DAY) ORAL

Date:05/31/00ISR Number: 3563272-1Report Type:Periodic Company Report #A0116094A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oliguria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL							

Date:05/31/00ISR Number: 3563274-5Report Type:Periodic Company Report #A0116133A  
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decreased Appetite Dysgeusia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL							

Date:05/31/00ISR Number: 3563279-4Report Type:Periodic Company Report #A0116136A  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL							

Date:05/31/00ISR Number: 3563282-4Report Type:Periodic Company Report #A0116148A  
 Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain Neck Pain	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL							

Date:05/31/00ISR Number: 3563284-8Report Type:Periodic Company Report #A0116154A  
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Twitching Sleep Disorder	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL 8 MON							
				Multivitamin	C		
				Ibuprofen	C		
				Nicotine	C		

Date:05/31/00ISR Number: 3563285-XReport Type:Periodic Company Report #A0116180A  
 Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Faeces Flatulence	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3563286-1Report Type:Periodic  
 Age:68 YR Gender:Male I/FU:I

Company Report #A0116182A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tremor	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL							

Date:05/31/00ISR Number: 3563287-3Report Type:Periodic  
 Age:30 YR Gender:Female I/FU:I

Company Report #A0116185A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Burning Sensation Dermatitis	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL							
		Disorientation					
		Hypoaesthesia Nervousness Pruritus Tachycardia					

Date:05/31/00ISR Number: 3563288-5Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0116197A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Amnesia Confusional State	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
15-0 MG (TWICE PER DAY) ORAL							
		Disturbance In Attention					
		Dizziness					
		Learning Disorder Thinking Abnormal Vision Blurred					

Date:05/31/00ISR Number: 3563289-7Report Type:Periodic Company Report #A0116198A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							

Date:05/31/00ISR Number: 3563290-3Report Type:Periodic Company Report #A0116200A  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Disturbance In Attention	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL		Headache Major Depression Tremor Visual Disturbance					

Date:05/31/00ISR Number: 3563291-5Report Type:Periodic Company Report #A0116201A  
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3563292-7Report Type:Periodic  
Age:65 YR Gender:Male I/FU:I

Company Report #A0116202A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG (PER DAY) ORAL		Abdominal Pain Nausea Oral Intake Reduced Vomiting	Consumer	Bupropion Hydrochloride  Glucophage	PS  C	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563296-4Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #A0116204A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG (TWICE PER DAY) ORAL		Dry Mouth Insomnia Sedation	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563301-5Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #A0116205A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG (TWICE PER DAY) ORAL	1 WK	Headache	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563806-7Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0115305A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Ear Discomfort	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Tinnitus

TWICE PER

DAY/ ORAL 4 WK

Date:05/31/00ISR Number: 3563808-0Report Type:Periodic Company Report #A0115306A  
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /TWICE		Diplopia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL		Hypertension					
		Muscle Spasms		Amlodipine	C		
		Muscle Twitching		Lisinopril	C		
		Tremor		Latanoprost	C		

Date:05/31/00ISR Number: 3563810-9Report Type:Periodic Company Report #A0115317A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /TWICE		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY /ORAL							

Date:05/31/00ISR Number: 3563812-2Report Type:Periodic Company Report #A0115319A  
Age:34 YR Gender:Female I/FU:I

Outcome	PT
	Chest Discomfort
	Confusional State

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Depersonalisation Dyspnoea Fatigue	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL		Tremor	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563813-4Report Type:Periodic Company Report #A0115339A  
Age:59 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT Gastrooesophageal Reflux Disease Nausea	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563815-8Report Type:Periodic Company Report #A0115352A  
Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT Insomnia	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL	2 DAY		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563816-XReport Type:Periodic Company Report #A0115367A  
Age:23 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT Nausea	Report Source	Product	Role	Manufacturer	Route
250 MG / PER DAY / ORAL			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3563817-1Report Type:Periodic Company Report #A0115375A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Feeling Abnormal					
DAY / ORAL		Hyperhidrosis					
		Syncope					
		Visual Disturbance					

Date:05/31/00ISR Number: 3563818-3Report Type:Periodic Company Report #A0115413A  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Depression					
PER DAY/ ORAL		Dizziness		Wellbutrin			
		Drug Ineffective		Tablet-Controlled			
		Feeling Abnormal		Release (Bupropion			
		Nausea		Hydrochloride)	SS		ORAL
ORAL	2 WK	Syncope		Multiple Medication	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3563819-5Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #A0115415A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Throat Irritation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							

Date:05/31/00ISR Number: 3563820-1Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #A0115419A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Constipation					
TWICE PER		Dysgeusia					
DAY/ ORAL	3 WK	Nausea					

Date:05/31/00ISR Number: 3563821-3Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0115420A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anger	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Arthralgia					
TWICE PER		Eyelid Oedema					
DAY/ ORAL		Face Oedema					
		Insomnia					
		Joint Swelling					
		Oedema Peripheral					
		Pruritus					
		Urticaria					

Date:05/31/00ISR Number: 3563822-5Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #A0115492A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG / PER		Muscle Spasms	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY /ORAL		Sensation Of Heaviness					

Date:05/31/00ISR Number: 3563823-7Report Type:Periodic Company Report #A0115493A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG / ORAL 3 WK		Liver Function Test	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Abnormal					

Date:05/31/00ISR Number: 3563824-9Report Type:Periodic Company Report #A0115509A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Muscle Twitching	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER							
DAY/ ORAL 2 WK							



150 MG /  
TWICE PER  
DAY/ ORAL  
Drug Ineffective  
Headache  
Consumer  
Zyban  
PS  
Glaxo Wellcome Inc  
ORAL

Date:05/31/00ISR Number: 3563830-4Report Type:Periodic Company Report #A0115642A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

150 MG /TWICE  
PER DAY/ ORAL

Date:05/31/00ISR Number: 3564384-9Report Type:Periodic Company Report #A0118239A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Hypersensitivity Rash Generalised	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

150 MG/ TWICE  
PER DAY/ ORAL

Date:05/31/00ISR Number: 3564397-7Report Type:Periodic Company Report #A0118665A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia Mouth Ulceration Pharyngolaryngeal Pain	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

150MG / TWICE  
PER DAY/ ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3564412-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0118666A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
OPHTHALMIC PER DAY / ORAL	150 MG/ TWICE 3 WK	Accommodation Disorder Burning Sensation Confusional State Elevated Mood Erythema Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:05/31/00ISR Number: 3564428-4Report Type:Periodic  
 Age:44 YR Gender:Female I/FU:I

Company Report #A0117914A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL UNK / UNK / ORAL		Dizziness Syncope Therapeutic Response Unexpected	Consumer	Zyban   Bupropion Hydrochloride Unspecified Tablet (Bupropion Hydrochloride)	PS   SS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3564438-7Report Type:Periodic  
 Age:45 YR Gender:Male I/FU:I

Company Report #A0117916A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Rash Pruritic	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

TWICE PER DAY

/ ORAL

Date:05/31/00ISR Number: 3564442-9Report Type:Periodic Company Report #A0118240A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150MG / PER							
DAY / ORAL							

Date:05/31/00ISR Number: 3564443-0Report Type:Periodic Company Report #A0117917A  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Dry Mouth					
DAY / ORAL		Eructation					
		Flatulence					
		Gastrointestinal Disorder					
		Hyperchlorhydria					
		Insomnia					
		Nausea					
		Throat Irritation					
		Vomiting					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3564448-XReport Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #A0117919A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3564452-1Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0117924A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER		Confusional State	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY / ORAL							
Disturbance In Attention							
Sedation							
Omeprazole							
C							

Date:05/31/00ISR Number: 3564455-7Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0117938A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Asthenia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							
Chest Pain							
Derealisation							
Dizziness							
Dyspnoea							
Fatigue							
Malaise							
Memory Impairment							
Palpitations							
Speech Disorder							
Syncope							
Tachycardia							

Yawning

Date:05/31/00ISR Number: 3564457-0Report Type:Periodic Company Report #A0118283A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / UNK/ ORAL				Nicotine Nicotine	C C		

Date:05/31/00ISR Number: 3564463-6Report Type:Periodic Company Report #A0117945A  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fixed Eruption	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY / ORAL		Pruritus	Professional				
		Rash Erythematous					
		Rash Generalised Rash Macular		Prempro	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3564464-8Report Type:Periodic  
 Age:28 YR Gender:Male I/FU:I

Company Report #A0118310A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus Rash Papular	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Urticaria					
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3564466-1Report Type:Periodic  
 Age:53 YR Gender:Female I/FU:I

Company Report #A0118051A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphagia Hypersensitivity Hypoaesthesia Oral	Health Professional Company	Bupropion Hydrochloride Unspecified Tablet	PS	Glaxo Wellcome Inc	ORAL
75 MG / UNK /			Representative				
ORAL							

Date:05/31/00ISR Number: 3564467-3Report Type:Periodic  
 Age:30 YR Gender:Male I/FU:I

Company Report #A0118371A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Drug Ineffective	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Irritability					
DAY / ORAL		Laryngitis Lymphadenopathy Pain Pharyngolaryngeal Pain					

Date:05/31/00ISR Number: 3564468-5Report Type:Periodic Company Report #A0118052A  
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3564469-7Report Type:Periodic Company Report #A0118056A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
	1	WK					
Anxiety							
Feeling Abnormal							
Hyperhidrosis							

Date:05/31/00ISR Number: 3564470-3Report Type:Periodic Company Report #A0118393A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							
DAY / ORAL							



Drug Dependence Consumer Zyban PS Glaxo Wellcome Inc ORAL  
150 MG /  
TWICE PER DAY  
/ ORAL

Date:05/31/00ISR Number: 3564477-6Report Type:Periodic Company Report #A0118438A  
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Dyspnoea	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Oedema					
TWICE PER DAY		Vomiting					
/ ORAL							

Date:05/31/00ISR Number: 3564479-XReport Type:Periodic Company Report #A0118128A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL	14 DAY						

Date:05/31/00ISR Number: 3564482-XReport Type:Periodic Company Report #A0118133A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

/ ORAL 3 WK

Date:05/31/00ISR Number: 3564484-3Report Type:Periodic Company Report #A0118439A  
 Age:72 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY		Drug Ineffective Dysgeusia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

/ ORAL

Date:05/31/00ISR Number: 3564486-7Report Type:Periodic Company Report #A0118135A  
 Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / UNK		Anxiety Dyspepsia Pollakiuria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

/ ORAL 2 WK

Date:05/31/00ISR Number: 3564488-0Report Type:Periodic Company Report #A0118457A  
 Age:50 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Aggression Insomnia Irritability	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3564490-9Report Type:Periodic Company Report #A0118140A  
 Age:46 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Dysgeusia					
/ ORAL		Headache					
		Insomnia					
		Parosmia					
		Tinnitus					

Date:05/31/00ISR Number: 3564493-4Report Type:Periodic Company Report #A0118152A  
 Age:39 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Erythema	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Pruritus	Professional				
/ ORAL		Rash Erythematous					

Date:05/31/00ISR Number: 3564500-9Report Type:Periodic Company Report #A0118463A  
 Age:72 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Urticaria					
/ ORAL							



	Rash Erythematous	Consumer	Bupropion	PS	Glaxo Wellcome Inc	ORAL
	Rash Papular		Hydrochloride			
150 MG /						
TWICE PER DAY						
/ ORAL						

Date:05/31/00ISR Number: 3564510-1Report Type:Periodic Company Report #A0118488A  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abnormal Behaviour	Health	Bupropion	PS	Glaxo Wellcome Inc	ORAL
		Crying	Professional	Hydrochloride			
150 MG / PER		Depression					
DAY / ORAL							

Date:05/31/00ISR Number: 3564512-5Report Type:Periodic Company Report #A0118489A  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis	Consumer	Bupropion	PS	Glaxo Wellcome Inc	ORAL
		Eyelid Oedema		Hydrochloride			
150 MG /		Oedema					
TWICE PER DAY		Pruritus					
/ ORAL		Pyrexia					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3564515-0Report Type:Periodic  
 Age:48 YR Gender:Female I/FU:I

Company Report #A0118579A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Back Pain Renal Colic	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3564518-6Report Type:Periodic  
 Age:40 YR Gender:I/FU:I

Company Report #A0118596A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Condition Aggravated Constipation	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							
Headache							
Sinus Congestion							
Sinusitis							

Date:05/31/00ISR Number: 3564519-8Report Type:Periodic  
 Age:43 YR Gender:Male I/FU:I

Company Report #A0117473A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Central Nervous System	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL							
Function Test Abnormal							
Condition Aggravated Disturbance In Attention Irritability Restlessness							

Date:05/31/00ISR Number: 3564521-6Report Type:Periodic Company Report #A0118612A  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / UNK							
/ ORAL	4	DAY					

Date:05/31/00ISR Number: 3564524-1Report Type:Periodic Company Report #A0117475A  
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Nightmare					
PER DAY ORAL							

Date:05/31/00ISR Number: 3564526-5Report Type:Periodic Company Report #A0117485A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Cough					
PER DAY ORAL		Dry Mouth					
		Ejaculation Disorder					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3564530-7Report Type:Periodic  
 Age:61 YR Gender:Male I/FU:I

Company Report #A0117603A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL	3 DAY	Blister	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Dermatitis Excoriation Pruritus		Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	SS		ORAL
ORAL	1 WK						

Date:05/31/00ISR Number: 3564533-2Report Type:Periodic  
 Age:53 YR Gender:Female I/FU:I

Company Report #A0117654A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Insomnia Night Sweats	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Conjugated Estrogens	C		

Date:05/31/00ISR Number: 3564536-8Report Type:Periodic  
 Age:37 YR Gender:Female I/FU:I

Company Report #A0117659A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL	3 WK	Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TRANSDERMAL	21 MG			Nicotine Patch (Nicotine)	SS		
TRANSDERMAL							

Date:05/31/00ISR Number: 3564545-9Report Type:Periodic  
 Age:38 YR Gender:Male I/FU:I

Company Report #A0117660A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL		Paraesthesia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TRANSDERMAL	21 MG			Nicotine Patch (Nicotine)	SS		
TRANSDERMAL							

Date:05/31/00ISR Number: 3564548-4Report Type:Periodic Company Report #A0117686A  
 Age:42 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Migraine	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3564552-6Report Type:Periodic Company Report #A0117687A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL		Abnormal Faeces	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Dysgeusia					

Date:05/31/00ISR Number: 3564553-8Report Type:Periodic Company Report #A0117688A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Rash Erythematous	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Skin Discolouration					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3564556-3Report Type:Periodic Company Report #A0117694A  
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:05/31/00ISR Number: 3564563-0Report Type:Periodic Company Report #A0117719A  
 Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharyngolaryngeal Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							

Date:05/31/00ISR Number: 3564570-8Report Type:Periodic Company Report #A0117725A  
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Epistaxis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:05/31/00ISR Number: 3564573-3Report Type:Periodic Company Report #A0117779A  
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
		Pruritus					
		Rash Erythematous		Nicotine Patch			
		Therapeutic Response		(Nicotine)	SS		
TRANSDERMAL	TRANSDERMAL	Unexpected		Fluticasone			

Date:05/31/00ISR Number: 3564577-0Report Type:Periodic Company Report #A0117785A  
 Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Back Pain					
DAY ORAL		Dermatitis					
		Disturbance In Attention					
		Neck Pain					
		Pain In Jaw					

Date:05/31/00ISR Number: 3564580-0Report Type:Periodic Company Report #A0117793A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Dysgeusia					
PER DAY ORAL		Tobacco Abuse					

Date:05/31/00ISR Number: 3564583-6Report Type:Periodic Company Report #A0117879A  
 Age:42 YR Gender:Female I/FU:I

Outcome	PT
	Apathy
	Balance Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Dermatitis Difficulty In Walking Dizziness Dysarthria Excitability	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Fatigue Headache Hypoaesthesia Oral Sensation Of Foreign Body Speech Disorder Tremor		Atenolol Isosorbide Dinitrate Atorvastatin Calcium Clopidogrel Bisulphate Aspirin Vitamin B Complex Vitamin E Estrogen	C C C C C C C C C		

Date:05/31/00ISR Number: 3564592-7Report Type:Periodic Company Report #A0117903A  
Age: Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL			Dizziness	Consumer	Zyban Caffeine	PS C	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3564594-0Report Type:Periodic Company Report #A0117907A  
Age:73 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL			Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3564597-6Report Type:Periodic Company Report #A0117913A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Urticaria					
DAY ORAL							

Date:05/31/00ISR Number: 3564715-XReport Type:Periodic Company Report #A0118667A  
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Bupropion	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Feeling Abnormal		Hydrochloride			
TWICE PER DAY		Pain					
/ ORAL	2 WK	Pruritus					
		Rash Papular					
		Speech Disorder					
		Tremor					

Date:05/31/00ISR Number: 3564716-1Report Type:Periodic Company Report #A0118668A  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Bupropion	PS	Glaxo Wellcome Inc	ORAL
150 MG /				Hydrochloride			
TWICE PER DAY							
/ ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3564717-3Report Type:Periodic  
Age:58 YR Gender:Male I/FU:I

Company Report #A0118669A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL				Lisinopril	C		

Date:05/31/00ISR Number: 3564718-5Report Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #A0118675A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Decreased Lethargy	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Sedation					
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3564719-7Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0118676A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rectal Haemorrhage	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY /							
ORAL							

Date:05/31/00ISR Number: 3564720-3Report Type:Periodic Company Report #A0118682A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER DAY / ORAL 3 DAY							

Date:05/31/00ISR Number: 3564728-8Report Type:Periodic Company Report #A0118684A  
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER DAY / ORAL							

Date:05/31/00ISR Number: 3564730-6Report Type:Periodic Company Report #A0118686A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased Weight Increased	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER DAY / ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3564733-1Report Type:Periodic  
 Age:59 YR Gender:Male I/FU:I

Company Report #A0118696A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea Vomiting	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3564735-5Report Type:Periodic  
 Age:33 YR Gender:Female I/FU:I

Company Report #A0118729A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia Arthritis	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Dermatitis					
TWICE PER DAY		Muscle Disorder					
/ ORAL	2 WK	Muscle Spasms Muscular Weakness Pyrexia Tic Urticaria					

Date:05/31/00ISR Number: 3564737-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0118733A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Irritability	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Mood Swings					
TWICE PER DAY		Nervousness					
/ ORAL							

Date:05/31/00ISR Number: 3564739-2Report Type:Periodic Company Report #A0118734A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Erythema	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Flushing					
DAY / ORAL		Pain In Extremity					

Date:05/31/00ISR Number: 3564741-0Report Type:Periodic Company Report #A0118753A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia Bruxism	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Depression					
DAY / ORAL		Disorientation Disturbance In Attention Dysgeusia Feeling Abnormal Insomnia Irritability Malaise Movement Disorder Sensation Of Heaviness Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3564743-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0118754A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / UNK / ORAL	7 DAY	Headache	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3564744-6Report Type:Periodic  
 Age:44 YR Gender:Male I/FU:I

Company Report #A0118755A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG / PER DAY / ORAL		Abnormal Dreams Insomnia Therapeutic Response Unexpected Tremor	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3564747-1Report Type:Periodic  
 Age:42 YR Gender:Female I/FU:I

Company Report #A0118757A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Central Nervous System Stimulation Insomnia Mental Disorder Vision Blurred	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3564749-5Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0118758A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Irritability	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / UNK/							
ORAL							

Date:05/31/00ISR Number: 3564751-3Report Type:Periodic Company Report #A0118759A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG UNK /							
ORAL							
TRANSDERMAL UNK /UNK/				Nicotine Patch (Nicotine)	SS		
TRTRANSDERMAL							

Date:05/31/00ISR Number: 3564753-7Report Type:Periodic Company Report #A0118784A  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus Rash Erythematous	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3564754-9Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #A0111412A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3564755-0Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #A0118793A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Drug Ineffective	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3564756-2Report Type:Periodic  
Age:30 YR Gender:Male I/FU:I

Company Report #A0111413A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dermatitis Haemorrhage Subcutaneous Testicular Swelling	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3564757-4Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0111417A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Abdominal Pain Upper	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Chest Pain	Professional				
/ ORAL		Constipation					
		Dermatitis					
		Dyspnoea					
		Emotional Disorder					
		Hypersensitivity					
		Panic Attack					
		Pruritus					
		Swelling					
		Urticaria					

Date:05/31/00ISR Number: 3564758-6Report Type:Periodic Company Report #A0111419A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY / ORAL		Tremor					

Date:05/31/00ISR Number: 3564759-8Report Type:Periodic Company Report #A0111421A  
Age:57 YR Gender:Male I/FU:I

Outcome	PT
	Agitation
	Disturbance In Attention
	Dry Mouth
	Feeling Abnormal
	Gait Disturbance



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Irritability Sleep Disorder				
Dose	Duration		Report Source	Product	Role	Manufacturer
150 MG /			Health	Zyban	PS	Glaxo Wellcome Inc
TWICE PER DAY			Professional			
/ ORAL						Route
						ORAL

Date:05/31/00ISR Number: 3564760-4Report Type:Periodic Company Report #A0111424A  
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc
150 MG /						Route
TWICE PER DAY						ORAL
/ ORAL						

Date:05/31/00ISR Number: 3564761-6Report Type:Periodic Company Report #A0111541A  
 Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc
150 MG /		Cardiovascular Disorder				Route
TWICE PER DAY		Dry Mouth				ORAL
/ ORAL		Feeling Abnormal				
		Insomnia				

Date:05/31/00ISR Number: 3564762-8Report Type:Periodic Company Report #A0111542A  
 Age:47 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Arthralgia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Asthenia					
/ ORAL		Dizziness					
		Sensation Of Heaviness					

Date:05/31/00ISR Number: 3564763-XReport Type:Periodic Company Report #A0111595A  
Age:46 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3564764-1Report Type:Periodic Company Report #A0111604A  
Age:51 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / UNK		Cold Sweat	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
/ ORAL		Crying					
		Depression					
		Dry Mouth					
		Feeling Hot And Cold					
		Insomnia					
		Tachycardia					
		Tinnitus					
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3564765-3Report Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #A0111605A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Chromaturia					
TWICE PER DAY		Dermatitis					
/ ORAL		Dry Skin					
		Dysgeusia					
		Fatigue					
		Urine Odour Abnormal					

Date:05/31/00ISR Number: 3564766-5Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #A0111606A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Insomnia	Professional				
TWICE PER DAY		Tremor					
/ ORAL							

Date:05/31/00ISR Number: 3564767-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0111607A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / UNK							
/ ORAL	3 WK						

Date:05/31/00ISR Number: 3564768-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0111636A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / UNK							
/ ORAL							

Date:05/31/00ISR Number: 3564769-0Report Type:Periodic Company Report #A0111637A  
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Drug Ineffective					
TWICE PER DAY		Dry Mouth					
/ ORAL		Pruritus		Imipramine			
		Rash Erythematous		Hydrochloride	C		
		Tremor		Lansoprazole	C		
		Urinary Retention		Prempro	C		
		Visual Disturbance		Glucophage	C		
				Acarbose	C		
				Hyzaar	C		
				Nicotine	C		

Date:05/31/00ISR Number: 3564770-7Report Type:Periodic Company Report #A0111648A  
 Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG, ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3564771-9Report Type:Periodic Company Report #A0111743A  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG, TWICE		Urticaria					
PER DAY, ORAL	2 WK						

Date:05/31/00ISR Number: 3564772-0Report Type:Periodic Company Report #A0111744A  
 Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vomiting	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG, TWICE							
PER DAY, ORAL							

Date:05/31/00ISR Number: 3564774-4Report Type:Periodic Company Report #A0111745A  
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG, TWICE							
PER DAY, ORAL							

Date:05/31/00ISR Number: 3564775-6Report Type:Periodic Company Report #A0111746A  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG, TWICE		Eyelid Oedema	Professional				
PER DAY, ORAL		Joint Swelling					
		Oedema Peripheral					

Swelling

Date:05/31/00ISR Number: 3564862-2Report Type:Periodic Company Report #A0112744A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Bupropion			
150 MG /		Rash Generalised		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3564863-4Report Type:Periodic Company Report #A0112745A  
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decreased Appetite	Consumer	Bupropion			
150 MG /		Weight Decreased		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Weight Increased					
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3564864-6Report Type:Periodic Company Report #A0112765A  
Age:41 YR Gender:Female I/FU:I

Outcome	PT
	Dysgeusia
	Feeling Of Relaxation
	Insomnia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nausea

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG /		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY						
/ ORAL						

Date:05/31/00ISR Number: 3564866-XReport Type:Periodic Company Report #A0112766A  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Dizziness					
DAY / ORAL		Dysgeusia					
		Headache					
		Nausea					

Date:05/31/00ISR Number: 3564868-3Report Type:Periodic Company Report #A0112803A  
 Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Agitation					
TWICE PER DAY		Difficulty In Walking					
/ ORAL		Dry Mouth					
		Insomnia					
		Oedema Peripheral					
		Pain					
		Tinnitus					
		Tremor					
		Urticaria					

Date:05/31/00ISR Number: 3564870-1Report Type:Periodic Company Report #A0112922A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Lethargy					
TWICE PER DAY		Syncope					
/ORAL		Tremor					

Date:05/31/00ISR Number: 3564871-3Report Type:Periodic Company Report #A0112939A  
Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ORAL							

Date:05/31/00ISR Number: 3564874-9Report Type:Periodic Company Report #A0112970A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
UNKNOWN /							
ORAL							





150 MG /  
TWICE PER DAY  
/ ORAL

Date:05/31/00ISR Number: 3564885-3Report Type:Periodic Company Report #A0112996A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Pruritus					
UNKNOWN /							
ORAL							

Date:05/31/00ISR Number: 3564888-9Report Type:Periodic Company Report #A0113010A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Irritability	Professional				
TWICE PER DAY		Nervousness					
/ ORAL							

Date:05/31/00ISR Number: 3564890-7Report Type:Periodic Company Report #A0113012A  
Age:71 YR Gender:Female I/FU:I

Outcome	PT
	Dizziness
	Ill-Defined Disorder
	Pyrexia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vomiting

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG /		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY						
/ ORAL						

Date:05/31/00ISR Number: 3564891-9Report Type:Periodic Company Report #A0113017A  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Agraphia	Professional				
TWICE PER DAY		Anxiety					
/ ORAL		Tremor					

Date:05/31/00ISR Number: 3564892-0Report Type:Periodic Company Report #A0113019A  
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Urticaria					
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3564895-6Report Type:Periodic Company Report #A0113029A  
 Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG /	Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY	Hyperhidrosis					
/ ORAL	Insomnia					
	Tremor					

Date:05/31/00ISR Number: 3564896-8Report Type:Periodic Company Report #A0113074A  
 Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cough	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Drug Ineffective					
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3564897-XReport Type:Periodic Company Report #A0113081A  
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Disturbance In Attention					
TWICE PER DAY		Lethargy					
/ ORAL		Sedation					

Date:05/31/00ISR Number: 3564984-6Report Type:Periodic Company Report #A0111005A  
 Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

/ ORAL

Date:05/31/00ISR Number: 3564988-3Report Type:Periodic Company Report #A0111006A  
 Age:44 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dysgeusia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER		Nausea					
DAY/ ORAL				Protuss	C		
				Pilocarpine	C		
				Hydrochloride	C		
				Trazodone	C		

Date:05/31/00ISR Number: 3564993-7Report Type:Periodic Company Report #A0111007A  
 Age:47 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER							
DAY/ ORAL	3 WK						

Date:05/31/00ISR Number: 3564997-4Report Type:Periodic Company Report #A0111009A  
 Age:72 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Headache					
/ ORAL		Nausea					

Tremor

Date:05/31/00ISR Number: 3565003-8Report Type:Periodic Company Report #A0111010A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decreased Activity	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Pruritus	Professional				
DAY/ ORAL		Rash Papular					
		Urticaria					

Date:05/31/00ISR Number: 3565008-7Report Type:Periodic Company Report #A0111011A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Insomnia					
TWICE PER							
DAY/ ORAL							

Date:05/31/00ISR Number: 3565011-7Report Type:Periodic Company Report #A0111012A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE							
PER DAY /							
ORAL							



150 MG /  
TWICE PER  
DAY/ORAL

Date:05/31/00ISR Number: 3565025-7Report Type:Expedited (15-DaCompany Report #A0111095A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

150 MG/ PO

Date:05/31/00ISR Number: 3565028-2Report Type:Periodic Company Report #A0111152A  
Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decreased Appetite	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

150 MG /  
TWICE PER DAY  
/ ORAL

Dysgeusia  
Headache  
Nausea  
Psychomotor Hyperactivity  
Thirst



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565033-6Report Type:Periodic Company Report #A0111155A  
 Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Migraine	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:05/31/00ISR Number: 3565036-1Report Type:Periodic Company Report #A0111292A  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY /							
ORAL							

Date:05/31/00ISR Number: 3565039-7Report Type:Periodic Company Report #A0111293A  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Pruritus					
PER DAY/ ORAL							

Date:05/31/00ISR Number: 3565044-0Report Type:Periodic Company Report #A0111294A  
 Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:05/31/00ISR Number: 3565048-8Report Type:Periodic Company Report #A0111295A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Disturbance In Attention					
TWICE PER		Dizziness					
DAY/ ORAL		Headache					
		Tinnitus					

Date:05/31/00ISR Number: 3565051-8Report Type:Periodic Company Report #A0111302A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain Upper	Consumer	Bupropion			
150 MG /		Nervousness		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER		Obstructive Airways					
DAY/ ORAL		Disorder					
		Pruritus					
		Swelling					
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565055-5Report Type:Periodic  
 Age:34 YR Gender:Female I/FU:I

Company Report #A0111379A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Bronchospasm					
TWICE PER		Dyspnoea					
DAY/ ORAL		Fatigue					
		Pruritus					
		Tachycardia					
		Urticaria					

Date:05/31/00ISR Number: 3565111-1Report Type:Periodic  
 Age:65 YR Gender:Female I/FU:F

Company Report #A0104884A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Pruritus	Professional				
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3565112-3Report Type:Periodic  
 Age:39 YR Gender:Female I/FU:F

Company Report #A0106495A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL		Erythema	Professional				
		Pain					
		Pruritus					
		Urticaria					

Date:05/31/00ISR Number: 3565113-5Report Type:Periodic Company Report #A0118984A  
Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL	4 DAY	Insomnia Irritability Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565114-7Report Type:Periodic Company Report #A0118989A  
Age:61 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Arthralgia Subcutaneous Nodule Tendonitis	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565115-9Report Type:Periodic Company Report #A0119010A  
Age:49 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG./ TWICE PER DAY/ ORAL	12 WK	Drug Ineffective	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565116-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0119014A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL							

Date:05/31/00ISR Number: 3565117-2Report Type:Periodic  
Age:70 YR Gender:Female I/FU:I

Company Report #A0119015A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							
Pruritus							
Rash Erythematous							
Urticaria							

Date:05/31/00ISR Number: 3565118-4Report Type:Periodic  
Age:77 YR Gender:Female I/FU:F

Company Report #A0106504A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL 16 DAY							
Pain							
Pruritus							
Rash Erythematous							
Urticaria							
Professional							

Date:05/31/00ISR Number: 3565119-6Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #A0119016A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							

Disorientation

PER DAY/ ORAL

Emotional Disorder

Hypoaesthesia

Thinking Abnormal

Date:05/31/00ISR Number: 3565120-2Report Type:Periodic Company Report #A0119017A

Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							

DAY/ ORAL

Date:05/31/00ISR Number: 3565121-4Report Type:Periodic Company Report #A0119020A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Night Sweats	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							

PER DAY./

ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565122-6Report Type:Periodic  
Age:52 YR Gender:Female I/FU:F

Company Report #A0107720A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Conjunctival Hyperaemia Dermatitis Eyelid Oedema Pruritus Urticaria	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
1 SPRAY / PER DAY / INTRANASAL				Flonase Aqueous Spray (Fluticasone Propionate)	SS		NASAL

Date:05/31/00ISR Number: 3565123-8Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #A0119021A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Middle Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565124-XReport Type:Periodic  
Age:24 YR Gender:Female I/FU:F

Company Report #A0108626A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Insomnia	Health Professional	Zyban Desogen	PS C	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565125-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0119022A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/	ORAL	Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Headache		Nicotine Patch (Nicotine)	SS		
INTRADERMAL	21 MG	Ill-Defined Disorder					
INTRADERMAL	15 DAY	Nervousness					

Date:05/31/00ISR Number: 3565126-3Report Type:Periodic Company Report #A0119027A  
Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/	ORAL	Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Insomnia					

Date:05/31/00ISR Number: 3565127-5Report Type:Periodic Company Report #A0108628A  
Age:32 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Blood Pressure Increased	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Tachycardia	Professional				
/ ORAL				Oral Contraceptive	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565128-7Report Type:Periodic  
 Age:52 YR Gender:Female I/FU:I

Company Report #A0119050A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL	6 DAY	Insomnia Motion Sickness Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565129-9Report Type:Periodic  
 Age:45 YR Gender:Female I/FU:F

Company Report #A0108630A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Burning Sensation Dermatitis Dizziness Hypersensitivity Nausea Oedema Peripheral Pruritus Rash Generalised Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565130-5Report Type:Periodic  
 Age:35 YR Gender:Female I/FU:F

Company Report #A0109155A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abnormal Dreams	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL		Insomnia	Professional Other	Nicotine Patch (Nicotine)	SS		
TRANSDERMAL	21 MG/						
TRANSDERMAL				Nasonex	C		
				Cetirizine Hydrochloride	C		
				Medroxyprogesterone Ace	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Depression	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /			Professional				
TWICE PER DAY							
/ ORAL				Sertraline Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565133-0Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0119058A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ORAL		Hypoaesthesia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TRANSDERMAL	21 MG/ DAY/	Paraesthesia Pruritus PER Swelling		Nicotine Patch (Nicotine)	SS		
TRANSDERMAL		Weight Increased					

Date:05/31/00ISR Number: 3565134-2Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #A0119064A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Therapeutic Response	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
VARIABLE DOSE/ ORAL		Unexpected	Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	SS		ORAL

Date:05/31/00ISR Number: 3565135-4Report Type:Periodic  
Age:33 YR Gender:Female I/FU:F

Company Report #A0109322A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Rash Pruritic Vision Blurred	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565136-6Report Type:Periodic Company Report #A0119083A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE			Professional				
PER DAY/ ORAL	12	WK					

Date:05/31/00ISR Number: 3565137-8Report Type:Periodic Company Report #A0109324A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Dyspnoea	Professional				
TWICE PER DAY		Fatigue					
/ ORAL		Feeling Abnormal					
		Insomnia					
		Negativism					

Date:05/31/00ISR Number: 3565138-XReport Type:Periodic Company Report #A0119189A  
Age:49 YR Gender:Male I/FU:I

Outcome	PT
	Dermatitis
	Hyperaesthesia
	Myalgia
	Pyrexia
	Serum Sickness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Swelling

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL	10 DAY	Health Professional Company Representative	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565139-1Report Type:Periodic Company Report #A0109434A  
Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dermatitis Dizziness Dysphonia Erythema Rash Pruritic Throat Tightness	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565140-8Report Type:Periodic Company Report #A0109500A  
Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dermatitis Face Oedema Hypersensitivity Pruritus Urticaria	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565141-XReport Type:Periodic Company Report #A0119194A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL	4 MON						

Date:05/31/00ISR Number: 3565142-1Report Type:Periodic Company Report #A0109600A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Irritability	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Nausea	Professional				
TWICE PER DAY		Rash Erythematous					
/ ORAL		Skin Disorder		Isoniazid	C		
				Cyanocobalamin	C		
				Pyridoxine			
				Hydrochloride	C		
				Mylanta	C		
				Oral Contraceptive	C		

Date:05/31/00ISR Number: 3565143-3Report Type:Periodic Company Report #A0109723A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Pruritic	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565144-5Report Type:Periodic  
 Age:52 YR Gender:Female I/FU:F

Company Report #A0110048A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Dermatitis Pruritus Skin Lesion Speech Disorder Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565145-7Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:F

Company Report #A0110057A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dry Mouth Feeling Abnormal Insomnia Psychomotor Hyperactivity Tachycardia Tremor	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565146-9Report Type:Periodic  
 Age:59 YR Gender:Female I/FU:I

Company Report #A0119196A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Abdominal Pain Upper Depression Fatigue Feeling Abnormal Irritability Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Depressed Mood					
TWICE PER DAY		Dry Mouth					
/ ORAL		Dysgeusia					
		Fatigue					
		Insomnia					
		Malaise					
		Tremor					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphagia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Gastrointestinal Pain					
PER DAY/ ORAL		Pain					
		Pyrexia					
		Rash Pruritic					
		Urticaria					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565149-4Report Type:Periodic  
 Age:41 YR Gender:Female I/FU:F

Company Report #A0110114A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Leukocytoclastic	Professional				
DAY / ORAL		Vasculitis					
		Petechiae					
		Purpura					
		Rash Erythematous					
		Rash Generalised					
		Urticaria					

Date:05/31/00ISR Number: 3565150-0Report Type:Periodic  
 Age:55 YR Gender:Female I/FU:F

Company Report #A0110329A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Urticaria	Professional				
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3565153-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0118800A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Toxicologic Test Abnormal	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG UNK			Professional				
ORAL							

Date:05/31/00ISR Number: 3565155-XReport Type:Periodic  
 Age:50 YR Gender:Male I/FU:I

Company Report #A0111747A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Dizziness					
DAY/ ORAL		Headache					
		Nausea					
		Tremor					

Date:05/31/00ISR Number: 3565157-3Report Type:Periodic Company Report #A0118801A  
 Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Yawning					
PER DAY ORAL							

Date:05/31/00ISR Number: 3565161-5Report Type:Periodic Company Report #A0118844A  
 Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Alanine Aminotransferase					
PER DAY ORAL	1 MON	Increased		Simvastatin			
		Condition Aggravated		(Formulation			
		Hepatitis C		Unknown)			
				(Simvastatin)	SS		ORAL
UNK ORAL				Loratadine	C		
				Lansoprazole	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565162-7Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #A0111748A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:05/31/00ISR Number: 3565164-0Report Type:Periodic  
Age:20 YR Gender:Female I/FU:I

Company Report #A0118847A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
1 TABLET							
TWICE PER DAY							

ORAL

Date:05/31/00ISR Number: 3565165-2Report Type:Periodic  
Age:52 YR Gender:Male I/FU:I

Company Report #A0111749A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:05/31/00ISR Number: 3565167-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0118867A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:05/31/00ISR Number: 3565168-8Report Type:Periodic Company Report #A0111750A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Depressed Level Of	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Consciousness					
PER DAY/ ORAL		Dizziness		Tylenol Cold Medication (Formulation Unknown) (Tylenol And Cold Medication)	SS		

Date:05/31/00ISR Number: 3565169-XReport Type:Periodic Company Report #A0118881A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG							
UNKNOWN ORAL							

Date:05/31/00ISR Number: 3565170-6Report Type:Periodic Company Report #A0111814A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dermatitis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
300 MG/ TWICE		Pruritus	Professional				
PER DAY/ ORAL 2 WK							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565172-XReport Type:Periodic  
Age:61 YR Gender:Male I/FU:I

Company Report #A0118886A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Tremor					
PER DAY ORAL							

Date:05/31/00ISR Number: 3565174-3Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #A0118906A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Metrorrhagia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE			Professional				
PER DAY ORAL	10 DAY			Ortho-Tri-Cyclen	C		

Date:05/31/00ISR Number: 3565176-7Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0118912A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Headache					
PER DAY ORAL		Insomnia					

Date:05/31/00ISR Number: 3565178-0Report Type:Periodic  
Age:66 YR Gender:Female I/FU:I

Company Report #A0111815A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
300 MG/ TWICE			Professional				
PER DAY/ ORAL							

Date:05/31/00ISR Number: 3565179-2Report Type:Periodic Company Report #A0118913A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 TWICE PER		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY ORAL		Pruritus					
UNKNOWN	UNKNOWN			Tetracycline (Formulation Unknown) (Tetracycline)	SS		

Date:05/31/00ISR Number: 3565180-9Report Type:Periodic Company Report #A0111870A  
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/		Face Oedema	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
SINGLE DOSE/		Throat Tightness					
ORAL							

Date:05/31/00ISR Number: 3565182-2Report Type:Periodic Company Report #A0118925A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG TWICE		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL	4 WK						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565183-4Report Type:Periodic Company Report #A0112117A  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Urticaria					
PER DAY/ ORAL							

Date:05/31/00ISR Number: 3565185-8Report Type:Periodic Company Report #A0112119A  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Dizziness					
PER DAY/ ORAL		Flushing					

Date:05/31/00ISR Number: 3565186-XReport Type:Periodic Company Report #A0118926A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ill-Defined Disorder	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Nausea					
DAY ORAL		Vomiting					

Date:05/31/00ISR Number: 3565187-1Report Type:Periodic Company Report #A0112120A  
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Rash Erythematous					
PER DAY/ ORAL							

Urticaria

Date:05/31/00ISR Number: 3565188-3Report Type:Periodic Company Report #A0118949A  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Pruritus					
PER DAY ORAL		Urticaria					

Date:05/31/00ISR Number: 3565190-1Report Type:Periodic Company Report #A0112121A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Dry Mouth					
DAY/ ORAL		Fatigue					

Date:05/31/00ISR Number: 3565191-3Report Type:Periodic Company Report #A0118959A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG							
UNKNOWN ORAL	2 WK						



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565192-5Report Type:Periodic  
 Age:79 YR Gender:Female I/FU:I

Company Report #A0112122A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /PER							
DAY/ ORAL							

Date:05/31/00ISR Number: 3565194-9Report Type:Periodic  
 Age:23 YR Gender:Female I/FU:I

Company Report #A0118972A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Dry Mouth					
PER DAY ORAL		Hallucination					
		Nausea					
		Visual Field Defect					

Date:05/31/00ISR Number: 3565195-0Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #A0112123A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Fatigue					
DAY/ ORAL		Headache					
		Insomnia					
		Mental Impairment					
		Nausea					

Date:05/31/00ISR Number: 3565196-2Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0118976A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER							
DAY ORAL							

Date:05/31/00ISR Number: 3565197-4Report Type:Periodic Company Report #A0112124A  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Hyperhidrosis					
PER DAY/ ORAL							

Date:05/31/00ISR Number: 3565198-6Report Type:Periodic Company Report #A0118977A  
 Age:46 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Chest Discomfort					
PER DAY ORAL		Dyspnoea					
		Hypersensitivity					
		Tachycardia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565199-8Report Type:Periodic  
Age:63 YR Gender:Male I/FU:I

Company Report #A0112125A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE		Swelling					
PER DAY/ ORAL		Urticaria	Professional				
				Glucophage	C		
				Quinapril			
				Hydrochloride	C		
				Glibenclamide	C		

Date:05/31/00ISR Number: 3565200-1Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #A0112126A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Alcoholic Hangover					
DAY/ ORAL		Dizziness	Professional				
		Fatigue					
		Insomnia					
		Malaise					
		Nausea					
		Retching					

Date:05/31/00ISR Number: 3565201-3Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #A0118978A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Rash Erythematous					
PER DAY ORAL							

Date:05/31/00ISR Number: 3565202-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0112127A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							
DAY/ ORAL				Alprazolam	C		

Date:05/31/00ISR Number: 3565203-7Report Type:Periodic Company Report #A0118980A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:05/31/00ISR Number: 3565204-9Report Type:Periodic Company Report #A0112128A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flushing	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Night Sweats					
PER DAY/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565206-2Report Type:Periodic Company Report #A0112187A  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Myalgia					
PER DAY/ ORAL							

Date:05/31/00ISR Number: 3565207-4Report Type:Periodic Company Report #A0118981A  
 Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Crying	Professional				
PER DAY ORAL		Irritability					

Date:05/31/00ISR Number: 3565208-6Report Type:Periodic Company Report #A0112312A  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Diarrhoea					
PER DAY/ ORAL							

Date:05/31/00ISR Number: 3565210-4Report Type:Periodic Company Report #A0110653A  
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Bupropion			
150 MG/PER		Mental Disorder		Hydrochloride	PS		ORAL
DAY/ORAL							

Date:05/31/00ISR Number: 3565212-8Report Type:Periodic Company Report #A0110713A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL				Thyroxine Sodium	C		

Date:05/31/00ISR Number: 3565214-1Report Type:Periodic Company Report #A0110714A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia Eye Irritation	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Pharyngolaryngeal Pain					
PER DAY ORAL	2 WK	Rash Pruritic		Nortriptyline Prempro	C C		

Date:05/31/00ISR Number: 3565215-3Report Type:Periodic Company Report #A0110716A  
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea Retching	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Radiotherapy C

Date:05/31/00ISR Number: 3565218-9Report Type:Periodic  
Age:64 YR Gender:Male I/FU:I

Company Report #A0110727A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Abdominal Pain Nausea Retching	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565221-9Report Type:Periodic  
Age:68 YR Gender:Female I/FU:I

Company Report #A0110728A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Insomnia Tachycardia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565223-2Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0110849A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG TWICE PER DAY ORAL		Convulsion Movement Disorder Tremor	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Inhaler C

Date:05/31/00ISR Number: 3565226-8Report Type:Periodic  
Age:54 YR Gender:Male I/FU:I

Company Report #A0110942A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hyperglycaemia	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:05/31/00ISR Number: 3565227-XReport Type:Periodic Company Report #A0116543A  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dysuria Feeling Cold Penis Disorder	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL	6 WK						

Date:05/31/00ISR Number: 3565228-1Report Type:Periodic Company Report #A0110993A  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Pain Abdominal Pain Upper	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565230-XReport Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0116544A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY / ORAL	5 DAY	Dermatitis Dizziness Headache Hypertension	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565232-3Report Type:Periodic  
Age:52 YR Gender:Male I/FU:I

Company Report #A0116579A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dyspnoea Gingival Bleeding Gingivitis Sleep Disorder	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565235-9Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0116767A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY / ORAL		Anxiety Discomfort Insomnia Nervousness Nightmare	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565237-2Report Type:Periodic  
Age:77 YR Gender:Male I/FU:I

Company Report #A0116768A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Insomnia Nausea	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Paraesthesia					
/ ORAL		Psychomotor Hyperactivity					
		Tremor					

Date:05/31/00ISR Number: 3565239-6Report Type:Periodic Company Report #A0116769A  
 Age:32 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dermatitis Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3565242-6Report Type:Periodic Company Report #A0116770A  
 Age:45 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER		Anxiety	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
DAY / ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565246-3Report Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #A0116771A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Abdominal Discomfort Agitation	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565248-7Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0116773A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ UNK / ORAL		Face Oedema Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TRANSDERMAL	21 MG / UNK/			Nicotine Patch	SS		
TRANSDERMAL				Thyroxine Sodium	C		

Date:05/31/00ISR Number: 3565250-5Report Type:Periodic  
Age:21 YR Gender:Male I/FU:I

Company Report #A0116777A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Depersonalisation Depression Disorientation Dry Mouth Feeling Abnormal Hypersomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Psychiatric Symptom

Date:05/31/00ISR Number: 3565253-0Report Type:Periodic Company Report #A0116779A  
 Age:51 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL 8 DAY							

Date:05/31/00ISR Number: 3565258-XReport Type:Periodic Company Report #A0110994A  
 Age:43 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Depressed Mood Sedation	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL							

Date:05/31/00ISR Number: 3565259-1Report Type:Periodic Company Report #A0110995A  
 Age:61 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE		Dizziness Emotional Disorder	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY ORAL							
Feeling Abnormal							
Tremor							



Insomnia  
Tinnitus  
Consumer  
Bupropion  
Hydrochloride  
PS  
Glaxo Wellcome Inc  
ORAL  
150 MG /  
Tremor  
TWICE PER DAY  
/ ORAL

Date:05/31/00ISR Number: 3565264-5Report Type:Periodic Company Report #A0111000A  
Age:41 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety Depression	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Nervousness					

150 MG /  
TWICE PER  
DAY/ ORAL

Date:05/31/00ISR Number: 3565266-9Report Type:Periodic Company Report #A0111001A  
Age:59 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis Hypersensitivity	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

150 MG/ TWICE  
PER DAY /ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565267-0Report Type:Periodic  
 Age:42 YR Gender:Male I/FU:I

Company Report #A0111002A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY / ORAL		Dermatitis Ecchymosis Headache Pruritus Swelling Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565268-2Report Type:Periodic  
 Age:35 YR Gender:Male I/FU:I

Company Report #A0116792A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Abdominal Pain Diarrhoea	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565269-4Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0111003A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Drug Ineffective Irritability	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565270-0Report Type:Periodic Company Report #A0111004A  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /ORAL		Nausea Tinnitus Tremor					

Date:05/31/00ISR Number: 3565271-2Report Type:Periodic Company Report #A0116795A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE							
PER DAY /							
ORAL							

Date:05/31/00ISR Number: 3565272-4Report Type:Periodic Company Report #A0116796A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Photosensitivity Reaction	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565273-6Report Type:Periodic  
 Age:48 YR Gender:Male I/FU:I

Company Report #A0116797A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression Agitation	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Feeling Abnormal					
TWICE PER DAY		Irritability					
/ ORAL							

Date:05/31/00ISR Number: 3565275-XReport Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #A0116803A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal Feeling Cold	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3565277-3Report Type:Periodic  
 Age:74 YR Gender:Female I/FU:I

Company Report #A0116806A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth Weight Increased	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3565279-7Report Type:Periodic  
 Age:28 YR Gender:Female I/FU:I

Company Report #A0116808A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Agitation Dry Mouth	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565280-3Report Type:Periodic Company Report #A0116894A  
 Age:59 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Agitation Anxiety Headache	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565282-7Report Type:Periodic Company Report #A0116898A  
 Age:16 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage / ORAL		Pruritus Rash Erythematous Urticaria	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565701-6Report Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #A0116207A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Agitation	Professional				
PER DAY ORAL		Anxiety Depression Irritability Mood Altered Nervousness		Nicotine	C		

Date:05/31/00ISR Number: 3565704-1Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #A0116208A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Swelling	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:05/31/00ISR Number: 3565707-7Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0116209A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Dyspnoea					
PER DAY ORAL		Face Oedema Hypersensitivity Pruritus Urticaria					

Date:05/31/00ISR Number: 3565709-0Report Type:Periodic  
Age:51 YR Gender:Male I/FU:I

Company Report #A0116269A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Urticaria					
PER DAY ORAL							

Date:05/31/00ISR Number: 3565711-9Report Type:Periodic Company Report #A0116281A  
 Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL	3 WK	Constipation					
		Flatulence					

Date:05/31/00ISR Number: 3565715-6Report Type:Periodic Company Report #A0116440A  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Pyrexia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Swelling					
PER DAY ORAL		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565716-8Report Type:Periodic  
 Age:23 YR Gender:Female I/FU:I

Company Report #A0116459A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:05/31/00ISR Number: 3565717-XReport Type:Periodic  
 Age:58 YR Gender:Male I/FU:I

Company Report #A0116462A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Drug Ineffective					
PER DAY ORAL							

Date:05/31/00ISR Number: 3565718-1Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #A0116468A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Burning Sensation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Depression					
PER DAY ORAL		Dermatitis					
		Dissociation					
		Elevated Mood					
		Euphoric Mood					
		Hypersensitivity					
		Irritability					
		Lichen Planus					
		Oedema Genital					
		Pruritus					
		Urticaria					

Date:05/31/00ISR Number: 3565719-3Report Type:Periodic Company Report #A0116473A  
Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:05/31/00ISR Number: 3565722-3Report Type:Periodic Company Report #A0116475A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							
		Urticaria	Professional Company Representative				

Date:05/31/00ISR Number: 3565732-6Report Type:Periodic Company Report #A0116480A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
		Depression					
		Neoplasm Swelling					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565741-7Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0116490A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL	3 WK	Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565742-9Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0116493A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Tinnitus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565743-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0116494A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Fluid Retention	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565744-2Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #A0116529A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL	7 DAY	Dermatitis Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565745-4Report Type:Periodic Company Report #A0116530A  
Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:05/31/00ISR Number: 3565746-6Report Type:Periodic Company Report #A0116540A  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Pruritic	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							

Date:05/31/00ISR Number: 3565747-8Report Type:Periodic Company Report #A0113683A  
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							
Face Oedema							

Date:05/31/00ISR Number: 3565748-XReport Type:Periodic Company Report #A0113690A  
Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperuricaemia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL							





Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Anxiety Communication Disorder Feeling Abnormal	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Date:05/31/00ISR Number: 3565760-0Report Type:Periodic Company Report #A0113747A Age:42 YR Gender:Male I/FU:I							
150 MG / TWICE PER DAY / ORAL	3 WK	Asthenia Fatigue Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Nicotine	C		
Date:05/31/00ISR Number: 3565761-2Report Type:Periodic Company Report #A0113748A Age:71 YR Gender:Male I/FU:I							
150 MG / TWICE PER DAY / ORAL		Dissociation Dry Mouth Tremor	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Date:05/31/00ISR Number: 3565765-XReport Type:Periodic Company Report #A0113749A Age:64 YR Gender:Male I/FU:I							
150 MG / TWICE PER DAY / ORAL		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565771-5Report Type:Periodic  
 Age:34 YR Gender:Female I/FU:I

Company Report #A0113750A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE							
PER DAY/ ORAL							

Date:05/31/00ISR Number: 3565773-9Report Type:Periodic  
 Age:38 YR Gender:Male I/FU:I

Company Report #A0113751A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE							
PER DAY/ ORAL							
Glossodynia							
Tongue Disorder							

Date:05/31/00ISR Number: 3565776-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0113752A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER							
DAY/ ORAL							

Date:05/31/00ISR Number: 3565782-XReport Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0113753A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Middle Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							

Date:05/31/00ISR Number: 3565783-1Report Type:Periodic Company Report #A0113754A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER							
DAY/ ORAL	9	DAY					

Date:05/31/00ISR Number: 3565784-3Report Type:Periodic Company Report #A0113755A  
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:05/31/00ISR Number: 3565785-5Report Type:Periodic Company Report #A0113756A  
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lethargy	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER							
DAY/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565786-7Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0113758A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150 MG / ORAL	Lethargy	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565787-9Report Type:Periodic  
 Age:37 YR Gender:Female I/FU:I

Company Report #A0113765A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150 MG / PER DAY/ ORAL	Decreased Appetite Dizziness Feeling Abnormal Nausea Vomiting	Consumer	Zyban  Nicotine	PS  C	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565788-0Report Type:Periodic  
 Age:66 YR Gender:Female I/FU:I

Company Report #A0113768A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150 MG / TWICE PER DAY/ ORAL	Weight Increased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3565789-2Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0113900A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150 MG /	Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

TWICE PER

DAY/ ORAL

Date:05/31/00ISR Number: 3565790-9Report Type:Periodic Company Report #A0113099A  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

150 MG/TWICE

PER DAY/ORAL

Date:05/31/00ISR Number: 3565791-0Report Type:Periodic Company Report #A0113102A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

150 MG /

TWICE PER DAY

/ ORAL

Date:05/31/00ISR Number: 3565793-4Report Type:Periodic Company Report #A0113109A  
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cerumen Impaction	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

150 MG / ORAL

Feeling Abnormal  
Gingival Pain

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565794-6Report Type:Periodic Company Report #A0113125A  
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY / ORAL							

Date:05/31/00ISR Number: 3565795-8Report Type:Periodic Company Report #A0113130A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Unevaluable Event	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							

Date:05/31/00ISR Number: 3565796-XReport Type:Periodic Company Report #A0113368A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER DAY / ORAL							

Date:05/31/00ISR Number: 3565797-1Report Type:Periodic Company Report #A0113369A  
 Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Behaviour	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE PER DAY/ORAL MON Aggression Crying Drug Dependence							

Drug Ineffective  
Insomnia  
Psychiatric Symptom

Date:05/31/00ISR Number: 3565798-3Report Type:Periodic Company Report #A0113370A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY /							
ORAL							

Date:05/31/00ISR Number: 3565799-5Report Type:Periodic Company Report #A0113379A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL				Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	SS		ORAL
ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565800-9Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #A0113408A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Agitation					
PER DAY /		Dizziness					
ORAL		Insomnia		Lisinopril	C		
				Verapamil			
				Hydrochloride	C		
				Estropipate	C		
				Medroxyprogesterone			
				Ace.	C		

Date:05/31/00ISR Number: 3565801-0Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #A0113559A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3565802-2Report Type:Periodic  
Age:26 YR Gender:Male I/FU:I

Company Report #A0113573A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Insomnia					
DAY / ORAL							

Date:05/31/00ISR Number: 3565803-4Report Type:Periodic Company Report #A0113575A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE		Crying					
PER DAY /		Emotional Disorder					
ORAL	9 DAY						

Date:05/31/00ISR Number: 3565804-6Report Type:Periodic Company Report #A0113576A  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3565806-XReport Type:Periodic Company Report #A0113625A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Hypersensitivity	Professional				
TWICE PER DAY		Pruritus					
/ ORAL		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565807-1Report Type:Periodic  
 Age:32 YR Gender:Female I/FU:I

Company Report #A0113632A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Constipation					
PER DAY /		Feeling Hot And Cold					
ORAL		Insomnia					
		Menstruation Irregular					
		Tinnitus					
		Tremor					

Date:05/31/00ISR Number: 3565808-3Report Type:Periodic  
 Age:41 YR Gender:Male I/FU:I

Company Report #A0113645A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Disturbance In Attention					
PER DAY /		Dizziness					
ORAL		Fatigue					
		Feeling Abnormal					
		Irritability					
		Vision Blurred					

Date:05/31/00ISR Number: 3565809-5Report Type:Periodic  
 Age:42 YR Gender:Male I/FU:I

Company Report #A0113661A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Dysgeusia					
TWICE PER DAY			Professional				
/ ORAL							

Date:05/31/00ISR Number: 3565810-1Report Type:Periodic Company Report #A0113680A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3565922-2Report Type:Periodic Company Report #A0112333A  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3565925-8Report Type:Periodic Company Report #A0112334A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3565928-3Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0112354A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Cognitive Disorder	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Sleep Talking					
/ ORAL							

Date:05/31/00ISR Number: 3565930-1Report Type:Periodic  
 Age:34 YR Gender:Female I/FU:I

Company Report #A112357A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
UNKNOWN		Eye Movement Disorder					
/ORAL							

Date:05/31/00ISR Number: 3565932-5Report Type:Periodic  
 Age:31 YR Gender:Female I/FU:I

Company Report #A0112358A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Anger	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Decreased Appetite					
/ ORAL		Feeling Abnormal					
		Insomnia					
		Irritability					
		Muscle Rigidity					

Date:05/31/00ISR Number: 3565933-7Report Type:Periodic  
 Age:58 YR Gender:Female I/FU:I

Company Report #A0112359A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Condition Aggravated	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Confusional State	Professional				
/ ORAL		Drug Interaction					
50 MG / EIGHT		Dysgeusia		Desipramine Tablet	SS		ORAL
TIMES PER DAY		Flatulence					
ORAL		Nausea					
		Tremor		Lansoprazole	C		
		Ulcer		Theophylline	C		
				Salbutamol Sulphate	C		
				Beclomethasone			
				Dipropion	C		
				Diclofenac Sodium	C		
				Clopidogrel			
				Bisulphate	C		
				Conjugated Estrogens	C		
				Buspirone			
				Hydrochloride	C		
				Iodinated Glycerol	C		
				Thyroxine Sodium	C		
				Ticlopidine	C		

Date:05/31/00ISR Number: 3565935-0Report Type:Periodic Company Report #A0112360A  
Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Libido Decreased					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

/ ORAL

Date:05/31/00ISR Number: 3565939-8Report Type:Periodic Company Report #A0112361A  
 Age:40 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY		Depression Irritability	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

/ ORAL

Date:05/31/00ISR Number: 3565941-6Report Type:Periodic Company Report #A0112362A  
 Age:50 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY		Dysgeusia		Zyban	PS	Glaxo Wellcome Inc	ORAL

/ ORAL

Date:05/31/00ISR Number: 3565945-3Report Type:Periodic Company Report #A0112363A  
 Age:70 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY		Tremor		Zyban	PS	Glaxo Wellcome Inc	ORAL

/ ORAL

Date:05/31/00ISR Number: 3565948-9Report Type:Periodic Company Report #A0112364A  
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Pharyngolaryngeal Pain					
TWICE PER DAY							
/ ORAL				Tenoric	C		
				Alendronate Sodium	C		

Date:05/31/00ISR Number: 3565953-2Report Type:Periodic Company Report #A0112365A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Chest Discomfort					
TWICE PER DAY		Dermatitis					
/ ORAL		Diarrhoea		Ibuprofen	C		
		Oedema					
		Pruritus					
		Urticaria					

Date:05/31/00ISR Number: 3565955-6Report Type:Periodic Company Report #A0112366A  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Urticaria					
TWICE PER DAY							
/ ORAL							





150 MG /  
TWICE PER DAY  
/ ORAL  
Arthralgia  
Oedema Peripheral  
Consumer  
Zyban  
PS  
Glaxo Wellcome Inc  
ORAL

Date:05/31/00ISR Number: 3565975-1Report Type:Periodic Company Report #A0112398A  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /			Professional				
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3565980-5Report Type:Periodic Company Report #A0112399A  
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Night Sweats					
TWICE PER DAY		Tremor					
/ ORAL							

Date:05/31/00ISR Number: 3565981-7Report Type:Periodic Company Report #A0112400A  
Age:75 YR Gender:Female I/FU:I

Outcome	PT
	Agitation
	Asthenia
	Dyspnoea Exacerbated
	Excitability

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG /		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY						
/ ORAL						

Date:05/31/00ISR Number: 3565987-8Report Type:Periodic Company Report #A0114978A  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Dermatitis					
TWICE PER DAY		Pruritus					
/ ORAL				Arthritis Medication	C		

Date:05/31/00ISR Number: 3565988-XReport Type:Periodic Company Report #A0114981A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
ORAL							

Date:05/31/00ISR Number: 3565990-8Report Type:Periodic Company Report #A0115063A  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Insomnia	Professional				
TWICE PER DAY							

Rash Pruritic

/ ORAL

Date:05/31/00ISR Number: 3565991-XReport Type:Periodic Company Report #A0115070A

Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eyelid Oedema	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Face Oedema					
TWICE PER DAY		Rash Generalised					
/ ORAL		Urticaria					

Date:05/31/00ISR Number: 3565996-9Report Type:Periodic Company Report #A0115078A

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL			Professional Company Representative				

Date:05/31/00ISR Number: 3565998-2Report Type:Periodic Company Report #A0115079A

Age:51 YR Gender:Female I/FU:I

Outcome	PT
	Anxiety
	Feeling Abnormal
	Insomnia

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		Irritability Nightmare				
Dose	Duration		Report Source	Product	Role	Manufacturer
150 MG / PER			Consumer	Zyban	PS	Glaxo Wellcome Inc
DAY / ORAL	2 DAY					ORAL

Date:05/31/00ISR Number: 3566000-9Report Type:Periodic Company Report #A0115081A  
 Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Chest Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc
150 MG /						ORAL
TWICE PER DAY						
/ ORAL						

Date:05/31/00ISR Number: 3566001-0Report Type:Periodic Company Report #A0115111A  
 Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc
150 MG /		Hangover				ORAL
TWICE PER DAY						
/ ORAL	8 DAY					

Date:05/31/00ISR Number: 3566004-6Report Type:Periodic Company Report #A0115147A  
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Crying	Consumer	Zyban	PS	Glaxo Wellcome Inc
150 MG / PER						ORAL

DAY / ORAL

Date:05/31/00ISR Number: 3566007-1Report Type:Periodic Company Report #A0115180A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							

DAY / ORAL

Date:05/31/00ISR Number: 3566010-1Report Type:Periodic Company Report #A0115207A  
Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cluster Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Headache					
TWICE PER DAY		Nausea					
/ ORAL							

Date:05/31/00ISR Number: 3566014-9Report Type:Periodic Company Report #A0115212A  
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL		Personality Change		Oral Contraceptive	C		

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Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3566021-6Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0115214A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Flushing	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Professional Company Representative				

Date:05/31/00ISR Number: 3566024-1Report Type:Periodic  
Age:53 YR Gender:Male I/FU:I

Company Report #A0115217A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Drug Withdrawal Syndrome Dysgeusia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Insomnia Nervousness Night Sweats Nightmare Tension		Augmentin	C		

Date:05/31/00ISR Number: 3566026-5Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #A0115250A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL	4 WK	Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3566029-0Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #A0115267A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Pruritus					
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3566032-0Report Type:Periodic Company Report #A0115268A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Aggression	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Agitation	Professional	Oral Contraceptive			
		Drug Interaction		(Formulation			
		Mood Swings		Unknown) (Oral	SS		ORAL
ORAL							

Date:05/31/00ISR Number: 3566033-2Report Type:Periodic Company Report #A0115300A  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							



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Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3566034-4Report Type:Periodic  
 Age:48 YR Gender:Female I/FU:I

Company Report #A0115303A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Blood Pressure Increased	Professional				
DAY / ORAL		Feeling Abnormal		Lotensin Hct (Formulation Unknown) (Lotensin Hct)	SS		

Date:05/31/00ISR Number: 3566035-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0115304A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							

Date:05/31/00ISR Number: 3566039-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0112405A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
PER DAY/ORAL	3	WK					

Date:05/31/00ISR Number: 3566041-1Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0112407A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cough	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ORAL		Headache					
		Weight Increased					

Date:05/31/00ISR Number: 3566043-5Report Type:Periodic Company Report #A0112412A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accidental Overdose	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/THREE		Anxiety	Professional				
TIMES PER		Dizziness					
DAY/ORAL		Insomnia					
		Nausea					

Date:05/31/00ISR Number: 3566046-0Report Type:Periodic Company Report #A0112415A  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER							
DAY/ORAL							

Date:05/31/00ISR Number: 3566047-2Report Type:Periodic Company Report #A0112416A  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bone Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
PER DAY/ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3566061-7Report Type:Periodic Company Report #A0112417A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ORAL	1 WK	Mental Impairment	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3566063-0Report Type:Periodic Company Report #A0112421A  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/TWICE PER DAY/ORAL		Dry Mouth Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3566065-4Report Type:Periodic Company Report #A0112589A  
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ORAL	2 WK	Dizziness Influenza Nausea	Consumer	Zyban Irbesartan Chlorpheniramine Maleate	PS C C	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3566066-6Report Type:Periodic Company Report #A0112592A  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG/ORAL		Irritability	Consumer	Zyban Nicotine	PS SS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3566068-XReport Type:Periodic Company Report #A0112602A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Loss Of Consciousness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ORAL		Pruritus Urticaria					

Date:05/31/00ISR Number: 3566069-1Report Type:Periodic Company Report #A0112604A  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Hypersensitivity					
PER DAY/ORAL		Rash Papular Rash Pruritic					

Date:05/31/00ISR Number: 3566072-1Report Type:Periodic Company Report #A0112612A  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
PER DAY/ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3566074-5Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #A0112613A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nightmare	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Pain In Extremity					
PER DAY/ORAL							

Date:05/31/00ISR Number: 3566077-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0112614A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ORAL		Palpitations					

Date:05/31/00ISR Number: 3566079-4Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #A0112617A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rectal Haemorrhage	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE				Muscle Relaxant	C		
PER DAY/ORAL							

Date:05/31/00ISR Number: 3566081-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0112736A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ORAL							

Date:05/31/00ISR Number: 3566083-6Report Type:Periodic Company Report #A0112737A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Dysarthria					
PER DAY/ORAL		Speech Disorder		Ethanol	SS		ORAL
ORAL							

Date:05/31/00ISR Number: 3566084-8Report Type:Periodic Company Report #A0114964A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoaesthesia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE		Vision Blurred					
PER DAY /							
ORAL	5 DAY						

Date:05/31/00ISR Number: 3566087-3Report Type:Periodic Company Report #A0114965A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE		Dysphagia					
PER DAY /		Sleep Disorder					
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3566088-5Report Type:Periodic Company Report #A0112738A  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Erythema					
PER DAY/ORAL		Pruritus					
		Urticaria					

Date:05/31/00ISR Number: 3566091-5Report Type:Periodic Company Report #A0112739A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ORAL							

Date:05/31/00ISR Number: 3566092-7Report Type:Periodic Company Report #A0114968A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /ORAL		Pruritus					
		Urticaria					

Date:05/31/00ISR Number: 3566093-9Report Type:Periodic Company Report #A0112743A  
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER		Insomnia					
DAY/ORAL							

Date:05/31/00ISR Number: 3566094-0Report Type:Periodic Company Report #A0114975A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Decreased Appetite					
TWICE PER		Diarrhoea					
DAY/ ORAL		Flatulence					

Date:05/31/00ISR Number: 3566097-6Report Type:Periodic Company Report #A0114541A  
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Haematuria					
TWICE PER DAY		Insomnia					
/ ORAL							

Date:05/31/00ISR Number: 3566100-3Report Type:Periodic Company Report #A0114543A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL	1	MON					



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Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3566101-5Report Type:Periodic Company Report #A0114545A  
 Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /PER							
DAY / ORAL							

Date:05/31/00ISR Number: 3566103-9Report Type:Periodic Company Report #A0114549A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE		Depressed Level Of					
PER DAY /		Consciousness					
ORAL	7 DAY	Feeling Abnormal					
		Nervousness					
		Tremor					

Date:05/31/00ISR Number: 3566106-4Report Type:Periodic Company Report #A0114557A  
 Age:50 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /ORAL		Insomnia					

Date:05/31/00ISR Number: 3566108-8Report Type:Periodic Company Report #A114559A  
 Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accommodation Disorder	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							

TWICE PER DAY  
/ ORAL  
Dizziness  
Paraesthesia  
Sensation Of Pressure  
Syncope

Date:05/31/00ISR Number: 3566109-XReport Type:Periodic Company Report #A0114682A  
Age:32 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /PER DAY / ORAL		Dysgraphia Feeling Abnormal Feeling Drunk Insomnia Irritability Tremor	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3566111-8Report Type:Periodic Company Report #A0114683A  
Age:30 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ORAL		Crying Dry Mouth Feeling Abnormal Insomnia Major Depression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3566113-1Report Type:Periodic  
 Age:39 YR Gender:Male I/FU:I

Company Report #A0114684A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL		Bone Pain Dry Mouth Insomnia Irritability Psychomotor Hyperactivity	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3566115-5Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #A0114746A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /PER DAY / ORAL		No Adverse Drug Effect	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3566117-9Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0114838A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ ORAL		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3566119-2Report Type:Periodic  
 Age:33 YR Gender:Female I/FU:I

Company Report #A0114866A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3566123-4Report Type:Periodic Company Report #A0114869A  
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Derealisation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE		Flight Of Ideas					
PER DAY /		Gastric Disorder					
ORAL		Insomnia					
		Nightmare					

Date:05/31/00ISR Number: 3566125-8Report Type:Periodic Company Report #A0114870A  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspraxia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG		Tremor					
/VARIABLE							
DOSE/ ORAL							

Date:05/31/00ISR Number: 3566127-1Report Type:Periodic Company Report #A0114962A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							



150 MG /  
TWICE PER  
DAY / ORAL

Chest Pain  
Drug Ineffective  
Irritability

Consumer  
Zyban  
PS  
Glaxo Wellcome Inc  
ORAL

Date:05/31/00ISR Number: 3566147-7Report Type:Periodic Company Report #A0110371A  
Age:46 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Abdominal Pain Dizziness Flushing Gastrointestinal Disorder Ill-Defined Disorder Influenza Like Illness Pyrexia Urticaria	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3566150-7Report Type:Periodic Company Report #A0110563A  
Age:47 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dizziness Dry Mouth Insomnia Irritability	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3566152-0Report Type:Periodic  
Age:42 YR Gender:Female I/FU:F

Company Report #A0110565A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG / PER		Accommodation Disorder	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY / ORAL		Agitation					
		Disturbance In Attention					
		Dizziness					
		Mental Impairment					
		Nausea					
		Tremor					

Date:05/31/00ISR Number: 3566154-4Report Type:Periodic  
Age:26 YR Gender:Female I/FU:F

Company Report #A0110570A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Parosmia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3566159-3Report Type:Periodic  
Age:42 YR Gender:Female I/FU:F

Company Report #A0110701A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG / PER		Formication	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY / ORAL		Insomnia	Professional				

Date:05/31/00ISR Number: 3566162-3Report Type:Periodic  
Age:57 YR Gender:Female I/FU:F

Company Report #A0110704A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Chest Discomfort	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Chest Pain	Professional				
/ ORAL		Dizziness					
		Dry Mouth		Loratadine	C		
		Dry Throat		Gemfibrozil	C		
		Dysphagia		Cimetidine	C		
		Feeling Abnormal		Conjugated Estrogens	C		
				Salbutamol Sulphate	C		
				Vitamin E	C		
				Selenium	C		
				Vitamin B	C		
				Ascorbic Acid	C		

Date:05/31/00ISR Number: 3566164-7Report Type:Periodic Company Report #A0110707A  
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG /		Hyperventilation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Hypoaesthesia					
/ ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3566167-2Report Type:Periodic  
Age:73 YR Gender:Female I/FU:F

Company Report #A0110850A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Blood Pressure Increased Drug Ineffective	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Vicodin Cyclobenzaprine Hcl	C C		

Date:05/31/00ISR Number: 3566173-8Report Type:Periodic  
Age:45 YR Gender:Female I/FU:F

Company Report #A0110991A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG / ORAL 15 MG / 20 MG /		Agitation Drug Interaction Dysphonia Irritability Nervousness Tremor	Health Professional Company Representative	Zyban Thyroxine Sodium (Formulation Unknown) (Levothyroxine Sodium) Pravastatin Sodium (Formulation Uknown) (Pravastatin Sodium)	PS SS SS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3566175-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0055869A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Dermatitis Hypersensitivity Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3566176-3Report Type:Periodic Company Report #A0062919A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Pruritic	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							
DAY / ORAL				Alprazolam	C		

Date:05/31/00ISR Number: 3566178-7Report Type:Periodic Company Report #A0062920A  
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							
DAY / ORAL							

Date:05/31/00ISR Number: 3566214-8Report Type:Periodic Company Report #A0113901A  
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE		Tremor					
PER DAY/ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3566218-5Report Type:Periodic  
Age:20 YR Gender:Male I/FU:I

Company Report #A0113902A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Libido Decreased	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY /							
ORAL 8 WK							

Date:05/31/00ISR Number: 3566220-3Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #A0113903A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depressed Mood	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							
Irritability							

Date:05/31/00ISR Number: 3566225-2Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #A0113925A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE							
PER DAY/ ORAL							

Date:05/31/00ISR Number: 3566227-6Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #A0113936A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							

PER DAY/  
ORAL  
Constipation  
Dry Mouth  
Tremor

Date:05/31/00ISR Number: 3566228-8Report Type:Periodic Company Report #A0113937A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 ,G/ TWICE		Urticaria					
PER DAY /		Vision Blurred					
ORAL							

Date:05/31/00ISR Number: 3566231-8Report Type:Periodic Company Report #A0113938A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Rash Erythematous					
PER DAY/ ORAL							

Date:05/31/00ISR Number: 3566234-3Report Type:Periodic Company Report #A0113941A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Herpes Simplex	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY /							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3566236-7Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #A0113964A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL							

Date:05/31/00ISR Number: 3566239-2Report Type:Periodic  
Age:70 YR Gender:Female I/FU:I

Company Report #A0114126A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							

Date:05/31/00ISR Number: 3566242-2Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0114127A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ear Discomfort	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
	1	WK	Professional				
Tinnitus							

Date:05/31/00ISR Number: 3566247-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0114347A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							

Date:05/31/00ISR Number: 3566255-0Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #A0114348A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE		Depression					
PER DAY/ ORAL		Dry Mouth					
		Emotional Disorder					

Date:05/31/00ISR Number: 3566258-6Report Type:Periodic Company Report #A0114381A  
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Insomnia					
DAY /ORAL							

Date:05/31/00ISR Number: 3566262-8Report Type:Periodic Company Report #A0114390A  
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema Multiforme	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL			Professional Company Representative				



150 MG /  
TWICE PER  
DAY / ORAL

Chest Pain  
Dermatitis  
Dyspnoea  
Hypoaesthesia  
Swelling

Consumer Zyban PS Glaxo Wellcome Inc ORAL

Date:05/31/00ISR Number: 3566275-6Report Type:Periodic Company Report #A0114538A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /TWICE		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER ORAL/ORAL		Sedation					
		Tremor					
		Vision Blurred					

Date:05/31/00ISR Number: 3588334-4Report Type:Periodic Company Report #A0116900A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /TWICE		Dizziness	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY /							
ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3588335-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0116936A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL 3 WK		Pyrexia	Health Professional Company Representative	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3588336-8Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #A0116940A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Dizziness Dizziness Postural Euphoric Mood Feeling Abnormal Hypokinesia Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3588337-XReport Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0117055A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / SEE TEXT / ORAL		Delusion Irritability Suicide Attempt	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3588338-1Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #A0117149A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3588339-3Report Type:Periodic Company Report #A0117178A  
 Age:63 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Dyspnoea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3588340-XReport Type:Periodic Company Report #A0117179A  
 Age:40 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG /		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY/ ORAL							
Overdose							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3588341-1Report Type:Periodic  
Age:78 YR Gender:Male I/FU:I

Company Report #A0117200A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL	5 DAY	Benign Prostatic Hyperplasia Condition Aggravated Dysuria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3588342-3Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #A0117206A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Agitation Costochondritis Dysgeusia Insomnia Nausea	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3588343-5Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #A0117209A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3588344-7Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #A0117220A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Insomnia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Tinnitus	Professional				
/ ORAL	5	WK					

Date:05/31/00ISR Number: 3588345-9Report Type:Periodic Company Report #A0117226A  
 Age:71 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Insomnia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY			Professional				
/ ORAL				Felodipine	C		
				Hydrochlorothiazide	C		

Date:05/31/00ISR Number: 3588346-0Report Type:Periodic Company Report #A0117230A  
 Age:46 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY							
/ ORAL	3	WK					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3588347-2Report Type:Periodic  
 Age:51 YR Gender:Female I/FU:I

Company Report #A0117232A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Body Temperature					
DAY / ORAL		Decreased					
		Feeling Abnormal					
		Feeling Cold					
		Insomnia					
		Irritability					

Date:05/31/00ISR Number: 3588348-4Report Type:Periodic  
 Age:75 YR Gender:Male I/FU:I

Company Report #A0117233A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Rash Pruritic					
TWICE PER DAY							
/ ORAL	1 WK			Nicotine	C		

Date:05/31/00ISR Number: 3588349-6Report Type:Periodic  
 Age:39 YR Gender:Female I/FU:I

Company Report #A0117283A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disturbance In Attention	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Drug Dependence					
TWICE PER DAY							
/ ORAL							

Date:05/31/00ISR Number: 3588350-2Report Type:Periodic Company Report #A0117286A  
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
				Ibuprofen	C		

Date:05/31/00ISR Number: 3588351-4Report Type:Periodic Company Report #A0117299A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
		Drug Ineffective					
		Headache					
		Insomnia					
		Nausea					

Date:05/31/00ISR Number: 3588352-6Report Type:Periodic Company Report #A0117320A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL 2 WK							
		Confusional State					
		Drug Ineffective					
		Insomnia					
		Mental Impairment					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3588353-8Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #A0117337A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL	12 DAY						

Date:05/31/00ISR Number: 3588377-0Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #A0119212A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arrhythmia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Date:05/31/00ISR Number: 3588379-4Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #A0119215A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache Irritability	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Tremor					
DAY / ORAL							

Date:05/31/00ISR Number: 3588380-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0119342A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							

Date:05/31/00ISR Number: 3588383-6Report Type:Periodic Company Report #A0119347A  
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Apathy Dyspnoea Exacerbated	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Feeling Abnormal					
DAY/ ORAL		Therapeutic Response Unexpected					

Date:05/31/00ISR Number: 3588385-XReport Type:Periodic Company Report #A0119400A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypertension Pruritus	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
1 TABLET /		Rash Generalised					
TWICE PER							
DAY/ ORAL							

Date:05/31/00ISR Number: 3588387-3Report Type:Periodic Company Report #A0119413A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Dry Mouth	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Insomnia					
TWICE PER DAY							
/ ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3588390-3Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0119420A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Depression Irritability Mood Altered	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3588393-9Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0119423A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Dermatitis Exfoliative Hypersensitivity	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3588395-2Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #A0119426A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL		Drug Ineffective Nausea Oral Intake Reduced Vomiting	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3588397-6Report Type:Periodic  
Age:77 YR Gender:Male I/FU:I

Company Report #A0119432A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Constipation	Consumer	Bupropion			

150 MG /  
TWICE PER DAY  
/ ORAL  
500 MG / PER  
DAY / ORAL

Dry Mouth  
Insomnia

Hydrochloride PS Glaxo Wellcome Inc ORAL

Levofloxacin Tablet SS ORAL

Date:05/31/00ISR Number: 3588399-XReport Type:Periodic Company Report #A0119438A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

150 MG / ORAL

Date:05/31/00ISR Number: 3588402-7Report Type:Periodic Company Report #A0119439A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

150 MG /  
TWICE PER DAY  
/ ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3588403-9Report Type:Periodic  
Age:76 YR Gender:Female I/FU:I

Company Report #A0119441A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Difficulty In Walking Dizziness Gait Disturbance Sensation Of Heaviness	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3588404-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0119495A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL	1 WK	Abnormal Behaviour Drug Ineffective	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3588405-2Report Type:Periodic  
Age:37 YR Gender:Male I/FU:I

Company Report #A0119576A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Anxiety Depressed Mood Depression Drug Level Changed Exercise Electrocardiogram Abnormal Flushing Headache Heart Rate Abnormal Hypertension Hypothyroidism Insomnia Panic Disorder Pyrexia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Tachycardia  
Tremor

Date:05/31/00ISR Number: 3588406-4Report Type:Periodic Company Report #A0119758A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							

Date:05/31/00ISR Number: 3588407-6Report Type:Periodic Company Report #A0119849A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea Dizziness	Other	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							
TRANSDERMAL	14 MG/	Nausea		Nicotine Patch	SS		
TRANSDERMAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/00ISR Number: 3588408-8Report Type:Periodic  
Age:52 YR Gender:Male I/FU:I

Company Report #A0120199A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Flatulence	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3588409-XReport Type:Periodic  
Age:54 YR Gender:Male I/FU:F

Company Report #A0070637A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / SINGLE DOSE / ORAL		Accommodation Disorder Agitation Anxiety Confusional State Headache Hyperaesthesia Nervousness Paraesthesia Psychotic Disorder Tension Tinnitus	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/31/00ISR Number: 3588411-8Report Type:Periodic  
Age:61 YR Gender:Female I/FU:F

Company Report #A0103803A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY		Agitation Constipation Insomnia	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

/ ORAL

Date:06/02/00ISR Number: 3507890-5Report Type:Expedited (15-DaCompany Report #A005730

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Required	Abdominal Distension	Health	Zoloft	PS	Pfizer	
Intervention to	Dissociative Identity	Professional			Pharmaceuticals Inc	
Prevent Permanent	Disorder		Bupropion	SS		
Impairment/Damage	Nausea					
	Nervousness					
	Tremor					

Date:06/02/00ISR Number: 3507892-9Report Type:Expedited (15-DaCompany Report #A0088169A

Age:40 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Chest Pain
Initial or Prolonged	Face Oedema
	Fatigue
	Hyperchlorhydria
	Hypothyroidism
	Liver Function Test
	Abnormal
	Palpitations
	Paraesthesia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Pericardial Effusion Swelling					
		Urinary Tract Infection	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER DAY/ ORAL							

Date:06/05/00ISR Number: 3508735-XReport Type:Expedited (15-DaCompany Report #A0117763A  
Age:64 YR Gender:Female I/FU:F

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Ageusia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
Dose		Decreased Appetite	Professional				
Disability		Headache	Company Representative Other	Terbinafine (Formulation Unknown) (Terbinafine)	SS		ORAL
150 MG TWICE PER DAY ORAL				Aspirin	C		
250 MG PER DAY ORAL				Armour Thyroid	C		

Date:06/05/00ISR Number: 3508739-7Report Type:Expedited (15-DaCompany Report #A0119802A  
Age:8 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Arthralgia	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
Dose		Conjunctivitis	Professional				
Required		Erythema Multiforme		Methylphenidate Hcl	C		
100 MG TWICE		Joint Swelling		Adderall	C		
Intervention to PER DAY ORAL		Myalgia					
Prevent Permanent Impairment/Damage		Serum Sickness					
		Urticaria					

Date:06/05/00ISR Number: 3509248-1Report Type:Expedited (15-DaCompany Report #A0118008A  
Age:58 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - ORAL	Drug Interaction	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged	Hallucination	Professional Other	Zolpidem Tartrate (Formulation Unknown) (Zolpidem Tartrate)	SS		
			Alprazolam (Formulation Unknown) (Alprazolam)	SS		
			Temazepam (Formulation Unknown) (Temazepam)	SS		

Date:06/05/00ISR Number: 3509249-3Report Type:Expedited (15-DaCompany Report #A0121104A  
Age:50 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Death 150 MG/ ORAL	Drug Interaction	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
	Myocardial Infarction		Sildenafil Citrate Tablet (Sildenafil Citrate)	SS		ORAL

SINGLE DOSE.

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL

Co-Trimoxazole C

Date:06/05/00ISR Number: 3509251-1Report Type:Expedited (15-DaCompany Report #A0119185A  
Age:58 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / ORAL Initial or Prolonged			Difficulty In Walking	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Multiple Sclerosis	Prednisone	C		
			Health Professional				

Date:06/07/00ISR Number: 3508637-9Report Type:Direct Company Report #USP 081288  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Medication Error	Wellbutrin 100mg Wellbutrin Sr 100 (Bupirion 100 Sr)	PS SS	Glaxo Pharm Glaxo Pharm	

Date:06/08/00ISR Number: 3510563-6Report Type:Expedited (15-DaCompany Report #A007267  
Age:18 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 10.00 MG Initial or Prolonged TOTAL; DAILY; Required ORAL			Bone Marrow Depression	Glucotrol Xl	PS	Pfizer Inc	ORAL
			Diabetes Mellitus				
			Inadequate Control				
Intervention to 200.00 MG Prevent Permanent TOTAL; BID Impairment/Damage 100.00 MG TOTAL; QID			Drug Interaction	Wellbutrin	SS		
			Fatigue				
			Herpes Zoster	Ritalin	SS		
			Leukopenia				

Urine Glucose False  
Positive  
Vomiting

Date:06/08/00ISR Number: 3510643-5Report Type:Expedited (15-DaCompany Report #A0120487A  
Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG ORAL		Biopsy Skin Abnormal	Literature	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Eye Disorder Scar Stevens-Johnson Syndrome Toxic Epidermal Necrolysis Visual Disturbance	Health Professional				

Date:06/08/00ISR Number: 3510667-8Report Type:Expedited (15-DaCompany Report #USA/00/01444/LAS  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 250 MG, ONCE A DAY, ORAL		Ageusia Drug Interaction	Health Professional	Lamisil	PS	Novartis Pharmaceuticals Corp	ORAL
150 MG, ONCE A DAY, ORAL;				Wellbutrin	SS		ORAL
				Aspirin (Acetylsalicylic			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Acid) C  
 Thyroid C  
 Prempo (Provella-14) C

Date:06/09/00ISR Number: 3511199-3Report Type:Expedited (15-DaCompany Report #A0117940A  
 Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG /TWICE PER DAY/ ORAL		Asthenia Bronchospasm Burning Sensation	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Chest Pain Dizziness Drug Hypersensitivity Dyspnoea Eye Irritation Eyelid Oedema Haemorrhage Irritability Oedema Pain Palpitations Pruritus Rash Erythematous Rash Vesicular Toxic Shock Syndrome Urticaria		Carisoprodol Diazepam	C C		

Date:06/09/00ISR Number: 3511496-1Report Type:Expedited (15-DaCompany Report #A0121649A  
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Alanine Aminotransferase Increased	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Arthralgia Bronchospasm Dermatitis Eyelid Oedema Rash Generalised	Professional	Echinacea Ginseng Lignocaine Fusidate Sodium	C C C C		

Urticaria

Date:06/09/00ISR Number: 3516348-9Report Type:Periodic Company Report #209927  
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia		Accutane	PS	Hlr Technology	ORAL
2 PER DAY		Condition Aggravated					
ORAL		Confusional State		Serzone (Nefazodone			
		Depression		Hydrochloride)	SS		ORAL
2 PER DAY		Dry Skin					
ORAL		Headache		Wellbutrin			
		Lip Dry		(Bupropion			
		Musculoskeletal Stiffness		Hydrochloride)	SS		ORAL
ORAL		Nausea					
		Schizophrenia					
		Suicidal Ideation					
		Thinking Abnormal					
		Vomiting					
		Weight Decreased					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/12/00ISR Number: 3511675-3Report Type:Expedited (15-DaCompany Report #A0070194A

Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Amnesia	Consumer	Bupropion			
Other		Cognitive Disorder		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER		Convulsion					
DAY/ORAL		Drug Ineffective		Ibuprofen	C		
		Encephalopathy		General Anesthetic	C		
		Feeling Abnormal		Surgery	C		

Date:06/14/00ISR Number: 3513457-5Report Type:Direct

Company Report #

Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardio-Respiratory Arrest		Morphine Pca	PS		
PCA 1MG Q 8		Coma					
MIN				Bupropine 25mg	SS		
75MG QD				Diffenlyderm	C		
				Dartax	C		
				Dubansetere			
				Hydarcharde	C		
				Trarodore	C		
				Tigas	C		
				Cefraolin	C		
				Bethameehol	C		
				Proxirine	C		
				Polycreboph	C		
				Cromolyin	C		
				Primarin	C		
				Ketorolac	C		
				Tums	C		
				Bashosc	C		
				Hydrocodone/Apap	C		
				Fentanyl	C		
				Versied	C		
				Propofool	C		

Date:06/14/00ISR Number: 3514915-XReport Type:Expedited (15-DaCompany Report #A0121644A  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Cerebral Artery Embolism	Health Professional Company Representative	Wellbutrin Warfarin Sodium	PS C	Glaxo Wellcome Inc	

Date:06/14/00ISR Number: 3514916-1Report Type:Expedited (15-DaCompany Report #A0121923A  
Age:39 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability 150 MG PER DAY ORAL	Dizziness Fear Feeling Abnormal Hiccups	Foreign Health Professional Company Representative	Zyban Amitriptyline	PS C	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/14/00ISR Number: 3514918-5Report Type:Expedited (15-DaCompany Report #A0121921A

Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL		Anaphylactic Shock	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Dyspnoea					
		Hypersensitivity		Blood Pressure			
		Leukocytosis		Medication	C		
		Localised Oedema		Allegra-D	C		
		Pharyngolaryngeal Pain		Budesonide	C		
		Pyrexia					
		Urticaria					

Date:06/15/00ISR Number: 3513785-3Report Type:Direct

Company Report #

Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100MG QID ORAL	90 DAY	Drug Effect Decreased		Teva Usa Bupropion	PS	Teva Usa	ORAL
				Zyprexa	C		
				..	C		

Date:06/15/00ISR Number: 3514777-0Report Type:Expedited (15-DaCompany Report #A0122022A

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Retinal Degeneration	Health Professional	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL

Date:06/15/00ISR Number: 3514778-2Report Type:Expedited (15-DaCompany Report #A0122021A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Retinal Degeneration	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
ORAL			Professional				

Date:06/15/00ISR Number: 3514780-0Report Type:Expedited (15-DaCompany Report #B0082488A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Retinal Detachment	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
Intervention to		Retinal Exudates	Study				
TWICE PER DAY							
Prevent Permanent		Retinal Oedema	Health				
/ ORAL							
Impairment/Damage		Retinitis	Professional				
		Vasculitis					
		Visual Disturbance					

Date:06/19/00ISR Number: 3515684-XReport Type:Expedited (15-DaCompany Report #A0117734A

Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Colitis Ulcerative	Foreign	Bupropion			
		Diarrhoea	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
			Professional				
PER DAY/ORAL				Mesalazine	C		
				Cyclobenzaprine Hcl	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/19/00ISR Number: 3515685-1Report Type:Expedited (15-DaCompany Report #A0121698A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depressed Level Of	Literature	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL		Consciousness	Health				
		Grand Mal Convulsion	Professional				
		Intentional Misuse					
		Intentional Self-Injury					
		Lethargy					
		Status Epilepticus					

Date:06/19/00ISR Number: 3515686-3Report Type:Expedited (15-DaCompany Report #A0122018A  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Pressure	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Fluctuation					
Initial or Prolonged		Coma		Wellbutrin			
PER DAY/ORAL		Condition Aggravated		Tablet-Controlled			
		Emphysema		Released (Bupropion			
		Oxygen Saturation		Hydrochloride)	SS		ORAL
150 MG/TWICE		Decreased					
PER DAY/ORAL		Pneumonia		Levofloxacin	C		
				Alendronate Sodium	C		
				Frusemide	C		
				Metoprolol Succinate	C		
				Oestradiol	C		
				Verapamil	C		
				Fenofibrate	C		
				Potassium Chloride	C		
				Tylenol No. 3	C		

Date:06/20/00ISR Number: 3515908-9Report Type:Direct  
Age:5 MON Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SR 150 MG BID		Abnormal Behaviour Complications Of Maternal Exposure To Therapeutic		Wellbutrin Burroughs Well	PS	Burroughs Well	ORAL
PO		Drugs Convulsion Fear Tremor		Diflucan Nystatin Cream Neb Tx	C C C		

Date:06/20/00ISR Number: 3516367-2Report Type:Expedited (15-DaCompany Report #203784  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 250 MG 1 PER 12 HOUR ORAL		Arthralgia Arthritis	Study Health	Ticlid	PS	Syntex (Usa) Inc	ORAL
4.5 MG 1 PER 12 HOUR ORAL		Dermatitis Pyrexia	Professional	Sibrafiban (Sibrafiban)	SS		ORAL
150 MG DAILY 1 PER 12 HOUR ORAL				Wellbutrin (Ibuproion Hydrochloride) Placebo (Placebo)	SS SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/22/00ISR Number: 3517395-3Report Type:Direct  
Age:13 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 2 WK Initial or Prolonged	Nausea Pruritus Pyrexia Rash Erythematous Rash Papular Urticaria		Wellbutrin	PS		

Date:06/22/00ISR Number: 3518369-9Report Type:Expedited (15-DaCompany Report #D0008602A  
Age:35 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL Initial or Prolonged	Amaurosis Fugax Condition Aggravated	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/23/00ISR Number: 3518642-4Report Type:Expedited (15-DaCompany Report #A0122304A  
Age:64 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL Initial or Prolonged	Blood Magnesium Decreased Grand Mal Convulsion	Health Professional Other	Wellbutrin Sr Tarka Hydrochlorothiazide	PS C C	Glaxo Wellcome Inc	ORAL

Date:06/23/00ISR Number: 3518645-XReport Type:Expedited (15-DaCompany Report #A0122473A  
Age:67 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY /	Brain Stem Infarction Dysarthria	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

ORAL	4	MON	Dysphagia	Company			
			Hypoaesthesia Oral	Representative	Glipizide	C	
			Nervous System Disorder		Rosiglitazone	C	
			Tongue Paralysis		Antihypertensive	C	

Date:06/23/00ISR Number: 3518649-7Report Type:Expedited (15-DaCompany Report #A0122591A  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Study	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL			Health Professional				

Date:06/23/00ISR Number: 3518653-9Report Type:Expedited (15-DaCompany Report #A0122595A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Suicide Attempt	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL			Professional Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/23/00ISR Number: 3518706-5Report Type:Expedited (15-DaCompany Report #A0120940A  
 Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache	Foreign	Bupropion			
		Neck Pain	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Night Sweats	Professional				
PER DAY/ORAL		Pain					
		Paraesthesia					
		Visual Disturbance					

Date:06/26/00ISR Number: 3519482-2Report Type:Expedited (15-DaCompany Report #A0122595A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Suicidal Ideation	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL	8 DAY	Suicide Attempt	Professional				
			Company				
			Representative				

Date:06/26/00ISR Number: 3519486-XReport Type:Expedited (15-DaCompany Report #A0122538A  
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cardiovascular Disorder	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
Initial or Prolonged		Coma	Professional				
PER DAY/ ORAL		Lethargy		Opiate (Opiate)	SS		
		Neuroleptic Malignant		Benzodiazepines	SS		
		Syndrom					
		Overdose					
		Pupil Fixed					
		Rhabdomyolysis					
		Stupor					
		Toxicologic Test Abnormal					

Date:06/26/00ISR Number: 3519488-3Report Type:Expedited (15-DaCompany Report #A0122740A  
Age:75 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Abdominal Pain	Foreign	Bupropion			
Initial or Prolonged	Cholelithiasis	Consumer	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE						
	Dyspnoea					
PER DAY/ORAL						
	Pruritus					
	Urticaria					

Date:06/26/00ISR Number: 3519745-0Report Type:Expedited (15-DaCompany Report #B0082531A  
Age:59 YR Gender:Female I/FU:I

Outcome	PT
Life-Threatening	Aspartate
Hospitalization -	Aminotransferase
Initial or Prolonged	Increased
	Atrioventricular Block
	Complete
	Blood Creatine
	Phosphokinase Increased
	Blood Creatine
	Phosphokinase Mb

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Increased Bradycardia Cardiac Disorder Hypokinesia	Report Source	Product	Role	Manufacturer	Route
150 MG/ ORAL		Myocardial Infarction Supraventricular  Tachycardia	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:06/29/00ISR Number: 3521889-4Report Type:Direct  
Age:38 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1 OR 2 A Other ORAL		Anxiety Dizziness  Fatigue  Insomnia Joint Swelling Lymphocyte Count Decreased Memory Impairment Musculoskeletal Stiffness Myalgia Vision Blurred Visual Disturbance		Zyban / Wellbutrin Glaxo/ Wellcome	PS	Glaxco / Wellcome	ORAL

Date:06/29/00ISR Number: 3522237-6Report Type:Expedited (15-DaCompany Report #A0066944A  
Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG/ TWICE Initial or Prolonged PER DAY/ ORAL Disability Required Intervention to 100 MG /		Agitation  Bone Pain  Chest Discomfort Chest Pain Coronary Artery Occlusion	Health  Professional	Wellbutrin Sr   Wellbutrin Tablet (Bupropion Hydrochloride)	PS   SS	Glaxo Wellcome Inc	ORAL   ORAL

Prevent Permanent TWICE PER Impairment/Damage DAY/ ORAL	Dermatitis  Disorientation  Dyspnoea Fatigue Flatulence Increased Appetite  Pharyngolaryngeal Pain Pruritus Weight Increased	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	SS	ORAL
150 MG/ ORAL		Clonazepam Aspirin Sertraline Hydrochloride Semisodium Valproate	C C C C	

Date:06/29/00ISR Number: 3522338-2Report Type:Expedited (15-DaCompany Report #A0066944A  
 Age:68 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Agitation
Initial or Prolonged	Chest Discomfort
Disability	Chest Pain
Required	Coronary Artery
Intervention to	Atherosclerosis
Prevent Permanent	Coronary Artery Occlusion
Impairment/Damage	Dermatitis
	Disorientation
	Dyspnoea



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG/ TWICE PER DAY/ ORAL		Fatigue Increased Appetite Pain Pharyngolaryngeal Pain	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
		Pruritus	Professional				
100 MG/ TWICE PER DAY/ ORAL		Weight Increased		Wellbutrin Tablet (Bupropion Hydrochloride)	SS		ORAL
150 MG/ ORAL				Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	SS		ORAL
				Clonazepam	C		
				Aspirin	C		
				Sertraline			
				Hydrochloride	C		
				Semisodium Valproate	C		

Date:06/30/00ISR Number: 3522972-XReport Type:Expedited (15-DaCompany Report #A0120078A  
Age:58 YR Gender:Male I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG TWICE PER DAY ORAL		Hospitalization - Initial or Prolonged	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Convulsion	Professional				
		Disorientation		Clonazepam	C		
		Disturbance In Attention		Paroxetine			
		Grand Mal Convulsion		Hydrochloride	C		
		Mood Swings		Perphenazine	C		
		Visual Acuity Reduced					

Date:06/30/00ISR Number: 3522983-4Report Type:Expedited (15-DaCompany Report #A0122805A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Multiple Sclerosis Neurological Symptom	Health Professional Company Representative	Wellbutrin Sr	PS	Glaxo Wellcome Inc	

Date:07/03/00ISR Number: 3524436-6Report Type:Expedited (15-DaCompany Report #PRIUSA2000006683  
Age:32 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Literature Health Professional	Haldol	PS	Rw Johnson Pharmaceutical Research Institute Div Ortho Pharm	ORAL
ORAL				Amitriptyline (Amitriptyline)	SS		ORAL
ORAL				Bupropion (Amfebutamone)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/03/00ISR Number: 3524825-XReport Type:Expedited (15-DaCompany Report #B0083476A  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG / TWICE PER DAY / ORAL		Headache Malignant Hypertension	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:07/03/00ISR Number: 3524829-7Report Type:Expedited (15-DaCompany Report #A0123050A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs	Foreign Study Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:07/03/00ISR Number: 3524831-5Report Type:Expedited (15-DaCompany Report #A0122980A  
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG / TWICE PER DAY / ORAL		Cardiac Arrest Grand Mal Convulsion Myocardial Infarction Pulse Absent	Health Professional Company Representative	Wellbutrin Sr   Diazepam Ranitidine Hydrochloride Rosiglitazone Glibenclamide Metoprolol Tartrate Enalapril Maleate Atorvastatin Calcium	PS   C C C C C C	Glaxo Wellcome Inc	ORAL

Allopurinol C  
Fenofibrate C  
Nicotinic Acid C

Date:07/05/00ISR Number: 3524824-8Report Type:Expedited (15-DaCompany Report #A0123011A  
Age:62 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG / TWICE PER DAY / ORAL	Abnormal Behaviour Hallucination Hyperventilation Paranoia	Consumer	Bupropion Hydrochloride  Oxycodone Hydrochloride	PS  C	Glaxo Wellcome Inc	ORAL

Date:07/06/00ISR Number: 3525179-5Report Type:Direct Company Report #  
Age:27 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ? Initial or Prolonged	Convulsion	Health Professional	Wellbutrin Alcohol	PS C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/06/00ISR Number: 3525629-4Report Type:Expedited (15-DaCompany Report #HQ8104303JUL2000  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Cyanosis	Health Professional	Effexor	PS	Wyeth Ayerst Laboratories Inc	ORAL
25 MG ORAL							
		Drug Interaction Respiratory Arrest		Wellbutrin - Slow Release	SS		ORAL
300 MG IN THE							
AM AND 150 MG							
IN THE PM							
(DOSE WAS							
INCREASED)							

Date:07/11/00ISR Number: 3527248-2Report Type:Direct Company Report #  
Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Dyspnoea Pain		Zyban 150 Mg Glaxo-Wellcome	PS	Glaxo-Wellcome	ORAL
2 TABLE DAILY							
ORAL		Tachycardia					

Date:07/11/00ISR Number: 3527791-6Report Type:Expedited (15-DaCompany Report #JRFUSA2000004633  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Health	Duragesic	PS	Alza Corp	
TRANSDERMAL	TRANSD						
			Professional	Sertraline (Sertraline)	SS		
				Olanzapine (Olanzapine)	SS		
				Bupropion			

(Amfebutamone) SS  
Codeine (Codeine) SS

Date:07/13/00ISR Number: 3528993-5Report Type:Expedited (15-DaCompany Report #A0083735A  
Age:23 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150 MG/ TWICE PER DAY/ ORAL	Anaphylactoid Reaction Arthralgia Dyspnoea Face Oedema Joint Stiffness Oedema Peripheral Pruritus Serum Sickness Urticaria	Study Health Professional	Wellbutrin Sr Medroxyprogesterone Ace.	PS C	Glaxo Wellcome Inc	ORAL

Date:07/13/00ISR Number: 3528995-9Report Type:Expedited (15-DaCompany Report #A0117940A  
Age:47 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Asthenia Blister Bronchospasm Burning Sensation Chest Pain Cyst

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE		Dizziness Dyspnoea Ear Haemorrhage	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ORAL		Erythema	Professional				
		Eye Disorder		Carisoprodol	C		
		Eye Irritation		Diazepam	C		
		Eyelid Oedema					
		Folliculitis					
		Haemorrhage					
		Hypersensitivity					
		Irritability					
		Localised Infection					
		Pain					
		Palpitations					
		Pruritus					
		Rash Pustular					
		Swelling					
		Toxic Shock Syndrome					
		Urticaria					

Date:07/13/00ISR Number: 3529023-1Report Type:Expedited (15-DaCompany Report #A0116641A  
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Circulatory Collapse	Health				
PER DAY ORAL		Coronary Artery Disease	Professional				
		Myocardial Infarction					
		Ventricular Fibrillation					

Date:07/13/00ISR Number: 3529026-7Report Type:Expedited (15-DaCompany Report #A0122980A  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							

PER DAY ORAL

Grand Mal Convulsion

Professional

Myocardial Infarction  
Palpitations  
Pulse Absent  
Syncope

Company  
Representative

Diazepam C  
Ranitidine  
Hydrochloride C  
Rosiglitazone C  
Glibenclamide C  
Metoprolol Tartrate C  
Enalapril Maleate C  
Atorvastatin Calcium C  
Allopurinol C  
Fenofibrate C  
Nicotinic Acid C

Date:07/14/00ISR Number: 3529926-8Report Type:Expedited (15-DaCompany Report #B0083707A  
Age:29 YR Gender:Male I/FU:I

Outcome PT  
Required Arthralgia  
Intervention to Blood Creatine  
Prevent Permanent Phosphokinase Increased  
Impairment/Damage Blood Lactate  
Dehydrogenase Increased  
Mouth Ulceration

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Myalgia Urticaria White Blood Cell Count					
		Increased	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
/THREE TIMES			Professional				
PER DAY/ ORAL							

Date:07/14/00ISR Number: 3529929-3Report Type:Expedited (15-DaCompany Report #B0082488A  
Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Retinal Disorder					
Required Intervention to 150 MG /TWICE		Retinal Exudates	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Prevent Permanent PER DAY/ ORAL Impairment/Damage		Retinal Oedema	Professional				
		Retinal Vasculitis Retinitis Visual Disturbance					

Date:07/17/00ISR Number: 3530293-4Report Type:Direct Company Report #  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema		Zyban	PS		

Date:07/17/00ISR Number: 3530321-6Report Type:Direct Company Report #  
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Platelet Count Decreased		Zyban 150 Mg	PS		ORAL
1T PO BID							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aggression	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
40 RABLET		Crying	Health				
/SINGLE DOSE/		Depressed Level Of	Professional				
ORAL		Consciousness		Zyban Tablet- Zyban			
		Erectile Dysfunction		(Bupropion			
ORAL		Fall		Hydrochloride)	SS		ORAL
		Intentional Misuse		Digoxin	C		
		Nervousness		Captopril	C		
		Suicidal Ideation		Frusemide	C		
		Visual Disturbance					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dyskinesia	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Fatigue	Professional				
PER DAY/ORAL		Multiple Sclerosis	Company	Oral Contraceptive	C		
		Muscle Disorder	Representative				
		Neurological Symptom					
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/17/00ISR Number: 3530744-5Report Type:Expedited (15-DaCompany Report #A0123857A  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG/ ORAL	Drug Level Above	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
		Therapeutic Drug Toxicity					

Date:07/17/00ISR Number: 3583408-6Report Type:Periodic Company Report #2000SUS0406  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Effect Decreased Drug Interaction	Health Professional	Sustiva	PS	Dupont Pharmaceuticals Co	ORAL
PO				Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL
PO				Combivir (Zidovudine/Lamivudi ne)	C		
				Paxil (Paroxetine Hydrochloride)	C		
				Depakote (Valproate Semisodium)	C		
				Lipitor (Atorvastatin Calcium)	C		

Date:07/18/00ISR Number: 3531310-8Report Type:Expedited (15-DaCompany Report #B0084242A  
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1 TABLET / TWICE PER DAY	Arthralgia Depression Dizziness	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

/ UNKNOWN

Headache  
Insomnia  
Neck Pain  
Tachycardia  
Tremor  
Visual Disturbance  
Vomiting

Date:07/19/00ISR Number: 3534897-4Report Type:Expedited (15-DaCompany Report #A0123997A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death UNK ORAL Congenital Anomaly		Complications Of Maternal Exposure To Therapeutic Drugs Hydrocephalus Induced Labour Stillbirth	Health Professional	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/21/00ISR Number: 3533529-9Report Type:Expedited (15-DaCompany Report #A0122740A

Age:75 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY ORAL	Abdominal Pain	Foreign Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
INTRAVENOUS INTRAVENOUS 24 HR	Dyspnoea Pruritus Urticaria	Professional	Cephzolin Sodium	SS		

Date:07/21/00ISR Number: 3533530-5Report Type:Expedited (15-DaCompany Report #D0009000A

Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - ORAL Initial or Prolonged Other	Retinal Artery Thrombosis	Foreign Health Professional Company Representative	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:07/21/00ISR Number: 3533534-2Report Type:Expedited (15-DaCompany Report #A0122304A

Age:64 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - ORAL 6 MON Initial or Prolonged	Blood Magnesium Decreased Grand Mal Convulsion	Health Professional Other	Wellbutrin Sr Tarka Hydrochlorothiazide	PS C C	Glaxo Wellcome Inc	ORAL

Date:07/24/00ISR Number: 3534226-6Report Type:Expedited (15-DaCompany Report #A0123751A

Age:56 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL	Chest X-Ray Abnormal  Dermatitis  Dysphagia Face Oedema Headache Hypersensitivity Mucous Membrane Disorder Nausea Proteinuria Pruritus Pyrexia Rash Maculo-Papular Tachycardia	Health  Professional  Company Representative	Wellbutrin    Omeprazole	PS    C	Glaxo Wellcome Inc	ORAL
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Date:07/24/00ISR Number: 3534227-8Report Type:Expedited (15-DaCompany Report #A0123746A  
Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 225 MG PER DAY ORAL	1 YR	Death	Health  Professional  Company Representative	Wellbutrin    Adderall (Formulation Unknown) (Adderall)	PS    SS	Glaxo Wellcome Inc	ORAL
20 MG IN THE MORNING	4 MON						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/24/00ISR Number: 3534232-1Report Type:Expedited (15-DaCompany Report #A0105636A

Age:17 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - SEE TEXT	Abdominal Pain	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged ORAL	Blood Creatine	Professional				
Other	Phosphokinase Increased Grand Mal Convulsion Hallucination Intentional Misuse Mental Impairment Sedation Sinus Tachycardia Suicide Attempt					

Date:07/24/00ISR Number: 3534249-7Report Type:Expedited (15-DaCompany Report #A0120791A

Age:20 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG THREE Initial or Prolonged TIMES PER DAY	Abnormal Behaviour	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL	Agitation	Professional				
	Confusional State					
	Delusion		Baclofen	C		
	Drug Level Above Therapeutic		Oxycodone Hydrochloride	C		
	Hallucination Paranoia Psychotic Disorder					

Date:07/24/00ISR Number: 3534892-5Report Type:Expedited (15-DaCompany Report #D0009020A

Age:58 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening UNKNOWN ORAL	Asthenia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Hospitalization - Blood Glucose Increased  
 Initial or Prolonged Dehydration  
 Fatigue  
 Ketoacidosis  
 Pollakiuria  
 Thirst  
 Weight Decreased

Hydrochlorothiazide C  
 Thyroxine Sodium C  
 Sitosterol C  
 Triamterene C

Date:07/24/00ISR Number: 3534894-9Report Type:Expedited (15-DaCompany Report #A0120136A  
 Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL		Dehydration	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
		Ovarian Cyst	Professional				
		Vomiting		Citalopram Hydrobromide	C		

Date:07/24/00ISR Number: 3536131-8Report Type:Direct Company Report #  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Gastric Hypertonia Tremor		Wellbutrin Sr 150mg Tablet	PS		ORAL
1T PO BID				Flexeril	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Motrin C

Date:07/25/00ISR Number: 3534553-2Report Type:Direct  
 Age:39 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Amnesia		Welbutrin	PS		
300 MG							
		Cognitive Disorder		Claritin	SS		
1 SR							
		Speech Disorder					

Date:07/25/00ISR Number: 3535059-7Report Type:Expedited (15-DaCompany Report #240967  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anxiety	Health	Klonopin	PS	Hoffmann La Roche	
Initial or Prolonged		Chest Pain	Professional			Inc	ORAL
ORAL							
		Convulsion	Other	Wellbutrin Sr			
		Drug Withdrawal		(Bupropion			
		Convulsions		Hydrochloride)	SS		ORAL
100 MG 1 PER							
		Drug Withdrawal Syndrome					
DAY ORAL							
		Electroencephalogram		Zolepidem (Zolpidem			
		Abnormal		Tartrate)	SS		ORAL
10 MG 1 PER							
		Grand Mal Convulsion					
DAY ORAL							
		Headache		Thyroxine Sodium			
		Spinal Fracture		(Levothyroxine			
				Sodium)	C		
				Loestrin (Ethinyl			
				Estradiol/Norethindr			
				one Acetate)	C		
				Quinapril			
				Hydrochloride			
				(Quinapril			
				Hydrochloride)	C		

Outcome	PT
Disability	Amnesia
	Anxiety
	Apathy
	Asthenia
	Breast Fibrosis
	Breast Hyperplasia
	Breast Mass
	Cognitive Disorder
	Convulsion
	Decreased Appetite
	Depression
	Disturbance In Attention
	Drug Ineffective
	Encephalopathy
	Fatigue
	Feeling Abnormal
	Fibrocystic Breast
	Disease
	Hodgkin'S Disease Nodular
	Sclerosis Stage

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Unspecified

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL		Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
			General Anesthetic (General Anesthetic)	SS		
			Ibuprofen	C		
			Surgery	C		
			Loperamide Hydrochloride	C		

Date:07/27/00ISR Number: 3536092-1Report Type:Direct  
Age:36 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1 QD F3 THEN 1 PO BID		Chest Pain		Zyban 150mg	PS		ORAL
		Dizziness					
		Dyspnoea Electrocardiogram Abnormal Orthostatic Hypotension		Motrin	C		

Date:07/27/00ISR Number: 3536508-0Report Type:Expedited (15-DaCompany Report #A0103180A  
Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG/THREE Initial or Prolonged TIMES PER DAY/ORAL		Angioneurotic Oedema	Health Professional	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
		Cardiac Arrest					
		Cough					
		Dysphagia Dyspnoea		Ace Inhibitor (Formulation			

Laryngeal Oedema  
Lymphocytosis  
Post Procedural  
Complication  
Throat Tightness  
Tongue Oedema  
Tracheal Oedema

Unknown)  
Pravastatin  
Lorazepam  
Aspirin  
Ibuprofen  
SS  
C  
C  
C  
C

Date:07/27/00ISR Number: 3536511-0Report Type:Expedited (15-DaCompany Report #A0124404A  
Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Burning Sensation  Decreased Activity Difficulty In Walking Pain Pruritus Swelling Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/28/00ISR Number: 3537865-1Report Type:Expedited (15-DaCompany Report #A0124525A

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - SEE TEXT/ Initial or Prolonged ORAL	Bradycardia	Foreign	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
	Complications Of Maternal Exposure To Therapeutic Drugs Neonatal Apnoeic Attack Oxygen Saturation Decreased Pregnancy	Study Health Professional				

Date:07/28/00ISR Number: 3537932-2Report Type:Expedited (15-DaCompany Report #A0120776A

Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - UNKNOWN Initial or Prolonged PER DAY	Bipolar Ii Disorder 200 MG TWICE Condition Aggravated	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	
UNKNOWN 100 MG UNKNOWN ORAL	Headache Muscle Twitching Petit Mal Epilepsy Rash Pruritic		Lamictal Tablet (Lamotrigine)	SS		ORAL
	Sinusitis Stevens-Johnson Syndrome Toxic Epidermal Necrolysis Tremor Urinary Incontinence		Clonazepam Omeprazole Dalmane Citalopram Hydrobromide Trazodone Compazine Sucralfate Multivitamin Olanzapine	C C C C C C C C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
UNKNOWN ORAL							
		Dehydration	Health	Ethanol (Formulation			
		Heat Stroke	Professional	Unknown) (Alcohol)	SS		
UNKNOWN	UNKNOWN						

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Cardiac Failure	Consumer	Pondimin	PS	Ah Robins Co	ORAL
Initial or Prolonged INTERMITTENTL		Cardiac Valve Disease		Phentermine	SS		
Other Y		Chest Pain					
				Prozac	SS		
20 TO 40 MG							
DAILY							
50 MG TWICE				Wellbutrin	SS		
DAILY							
100 MG DAILY				Zoloft	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/28/00ISR Number: 3541035-0Report Type:Periodic  
Age:43 YR Gender:Female I/FU:F

Company Report #8-97286-020L

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	20 MG THREE TIMES A DAY, ORAL	Cardiac Valve Disease	Consumer	Ponderex	PS	Ah Robins Co	ORAL
		Palpitations					
		Pulmonary Hypertension					
	15-30 MG DAILY, ORAL			Phentermine	SS		ORAL
	30 MG DAILY, ORAL			Redux	SS		ORAL
	100-150 MG TWICE DAILY, ORAL			Wellbutrin	SS		ORAL

Date:07/28/00ISR Number: 3541089-1Report Type:Periodic  
Age:47 YR Gender:Female I/FU:F

Company Report #8-97318-014L

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	60-120 MG Initial or Prolonged DAILY, ORAL	Cardiac Valve Disease	Consumer	Ponderex	PS	Ah Robins Co	ORAL
Other	75-225 MG DAILY	Pulmonary Hypertension		Effexor	SS		
	20-60 MG DAILY			Prozac	SS		
	150-300 MG			Wellbutrin	SS		

DAILY

50 MG DAILY

Zoloft SS

Amitriptyline C  
Cylert C  
Cytomel C  
Estraderm C  
Ritalin C

Date:07/31/00ISR Number: 3538605-2Report Type:Expedited (15-DaCompany Report #A0122099A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ SEE Initial or Prolonged TEXT/ ORAL		Agitation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Amnesia	Health				
		Overdose Suicide Attempt	Professional	Hydromorphone Hcl (Formulation Unknown) (Hydromorphone Hcl)	SS		ORAL
4 MG/ SEE TEXT/ ORAL				Cocaine	C		

Date:07/31/00ISR Number: 3538912-3Report Type:Expedited (15-DaCompany Report #B0084888A

Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death AS DIRECTED/ORAL		Sudden Death	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Health				
			Professional	Pravastatin Frusemide Aspirin Omeprazole Ramipril	C C C C C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/02/00ISR Number: 3540219-5Report Type:Expedited (15-DaCompany Report #000403-LX957  
 Age:57 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Discomfort	Health	Ambien	PS	Lorex	
Initial or Prolonged	Drug Interaction	Professional			Pharmaceuticals	ORAL
PO	Hallucination		Alprazolam	SS		
			Temazepam	SS		
			Wellbutrin	SS		ORAL
PO						

Date:08/07/00ISR Number: 3544684-9Report Type:Expedited (15-DaCompany Report #B0085017A  
 Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Epilepsy	Foreign	Bupropion			
Initial or Prolonged	Fatigue	Other	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL						
Required	Memory Impairment					
Intervention to	Nervous System Disorder					
Prevent Permanent	Status Epilepticus					
Impairment/Damage						

Date:08/07/00ISR Number: 3544832-0Report Type:Expedited (15-DaCompany Report #D0009329A  
 Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Hemiparesis		Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL	9 DAY					
Initial or Prolonged	Tinnitus					

Date:08/07/00ISR Number: 3544833-2Report Type:Expedited (15-DaCompany Report #B0085154A  
 Age:67 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					

Hospitalization - 150 MG/ORAL Initial or Prolonged	Cardiac Arrest Cardiac Enzymes Increased Circulatory Collapse Ventricular Fibrillation	Foreign	Zyban Phenytoin Carbamazepine	PS C C	Glaxo Wellcome Inc	ORAL
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Date:08/14/00ISR Number: 3550028-9Report Type:Expedited (15-DaCompany Report #A0125473A  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG TWICE PER DAY ORAL		Anxiety Bite	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Blood Potassium Decreased Blood Sodium Decreased Ecchymosis Fall Grand Mal Convulsion Head Injury Headache Memory Impairment Mouth Haemorrhage Musculoskeletal Stiffness Myalgia Tongue Disorder Tongue Oedema		Diltiazem Hydrochloride Thyroxine Sodium Prempro Maxzide Ranitidine Hydrochloride	C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/14/00ISR Number: 3550031-9Report Type:Expedited (15-DaCompany Report #B0085539A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Motion Sickness	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL		Vertigo Visual Acuity Reduced	Health Professional Company Representative				

Date:08/14/00ISR Number: 3550057-5Report Type:Expedited (15-DaCompany Report #A0122473A

Age:67 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150 MG (TWICE Initial or Prolonged PER DAY) ORAL 4 MON		Brain Stem Infarction	Health Professional Company Representative	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
		Dysarthria					
		Dysphagia		Glipizide	C		
		Hypoaesthesia		Rosiglitazone	C		
		Neurological Symptom Tongue Disorder Tongue Paralysis		Antihypertensive	C		

Date:08/14/00ISR Number: 3550060-5Report Type:Expedited (15-DaCompany Report #A0121921A

Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 150 MG (TWICE PER DAY) ORAL		Anaphylactic Shock Angioneurotic Oedema	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Dermatitis					
		Dyspnoea		Blood Pressure Medication	C		
		Hypersensitivity		Alleggra-D	C		
		Localised Oedema		Budesonide	C		
		Pharyngolaryngeal Pain Pruritus Pyrexia					

Urticaria  
White Blood Cell Count  
Increased

Date:08/14/00ISR Number: 3550113-1Report Type:Expedited (15-DaCompany Report #A0125213A  
Age:53 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 200 MG / Initial or Prolonged TWICE PER DAY/ ORAL	Balance Disorder Difficulty In Walking Fall Gait Disturbance Tremor	Health Professional	Wellbutrin  Valproic Acid Lithium Salt Venlafaxine Hydrochloride Clonazepam Glipizide Medroxyprogesterone Ace. Oestradiol	PS   C C C C C C C	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/14/00ISR Number: 3550171-4Report Type:Expedited (15-DaCompany Report #A0119646A  
Age:52 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY Required / ORAL	Anxiety Asthenia Chest Pain	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
Intervention to Prevent Permanent Impairment/Damage	Confusional State Depression Fatigue Flushing Headache Hypertension Insomnia Panic Disorder Without Agoraphobia Pruritus Psychotic Disorder Tinnitus		Fenofibrate Nicotine	C C		

Date:08/15/00ISR Number: 3551188-6Report Type:Expedited (15-DaCompany Report #AMANTADI2000-00169  
Age:94 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 100 MG QOTHRD PO 75 MG BID PO	Abnormal Behaviour Agitation Confusional State Dizziness Insomnia Memory Impairment Neurological Symptom Neurotoxicity Orthostatic Hypotension Pain In Extremity Peripheral Coldness Peripheral Embolism Restlessness	Literature Health Professional	Symmetrel Bupropriion Lasix Kci Quinaglute Vitamin B12 Feosol Multivitamin	PS SS C C C C C C	Endo Pharmaceuticals Inc	ORAL ORAL

Syncope  
Tremor

Date:08/16/00ISR Number: 3551544-6Report Type:Expedited (15-DaCompany Report #A0125519A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	
Disability		Asthenia					
		Gait Disturbance					
		Ill-Defined Disorder					
		Injury					
		Pain					

Date:08/16/00ISR Number: 3551560-4Report Type:Expedited (15-DaCompany Report #A0122805A  
Age:42 YR Gender:Female I/FU:F

Outcome  
Other  
Required  
Intervention to  
Prevent Permanent

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE		Coordination Abnormal	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL		Fatigue	Professional				
		Gait Disturbance Multiple Sclerosis Muscle Spasms Neurological Symptom Tremor	Company Representative	Oral Contraceptive	C		

Date:08/18/00ISR Number: 3553758-8Report Type:Expedited (15-DaCompany Report #240967  
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - ORAL		Anxiety Body Temperature	Health Professional	Clonopin	PS	Hoffmann La Roche Inc	ORAL
Initial or Prolonged Disability		Increased Chest Pain Drug Withdrawal	Other	Wellbutrin Sr (Bupropion Hydrochloride)	SS		ORAL
300 MG DAILY, 1 PER DAY; ORAL		Convulsions Drug Withdrawal Syndrome					
10 MG 1 PER DAY; ORAL		Electroencephalogram Abnormal Grand Mal Convulsion		Zolepidem (Zolpidem Tartrate)	SS		ORAL
		Headache Medication Error Personality Disorder Spinal Compression Fracture Spinal Fracture White Blood Cell Count Increased		Synthroid (Levothyroxine Sodium) Loestrin (Ethinyl Estradiol/Norethindr one Acette) Accupril (Quinapril Hydrochloride) Bcp (Oral Contraceptive Nos)	C C C C		

Ambien (Zolpidem  
Tartrate) C  
Celexa (Citalopram) C

Date:08/18/00ISR Number: 3588957-2Report Type:Periodic Company Report #S00-USA-00534-01  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Asthenia Malaise	Consumer	Celexa	PS	Forest Laboratories Inc	ORAL
20 MG PO				Wellbutrin(Amfebutam one Hydrochloride)	SS		
200 MG QD PO				Desyrel (Trazodone Hydrochloride)	C		
				Klonopin (Clonazepam)	C		
				Neurontin (Gabapentin)	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/18/00ISR Number: 3591383-3Report Type:Periodic Company Report #S00-USA-00489-01  
 Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Health Professional	Celexa	PS	Forest Laboratories Inc	ORAL
40 MG QD PO				Atenolol	SS		
				Verapamil	SS		
				Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL
300 MG QD PO				Ritalin (Methylphenidate Hydrochloride)	C		

Date:08/21/00ISR Number: 3554654-2Report Type:Expedited (15-DaCompany Report #A0125734A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bradycardia Cardiac Disorder Depressed Level Of Consciousness Hypotension Overdose	Health Professional Company Representative	Wellbutrin Olanzapine (Formulation Unknown) (Olanzapine)	PS SS	Glaxo Wellcome Inc	

Date:08/21/00ISR Number: 3554655-4Report Type:Expedited (15-DaCompany Report #A0125460A  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL							
Hospitalization - Initial or Prolonged		Fall Wrist Fracture	Professional Company Representative	Clozapine Lisinopril Docusate Sodium Amlodipine	C C C C		

Date:08/22/00ISR Number: 3554813-9Report Type:Direct  
Age:47 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	1 PO BID	Hepatitis	Health	Wellbutrin 75	PS		ORAL
Required Intervention to Prevent Permanent Impairment/Damage		Liver Function Test Abnormal	Professional	Ambien Fiortal/Codeine Morphine ... Cimetidine Ventolin	C C C C C C		

Date:08/22/00ISR Number: 3555915-3Report Type:Expedited (15-DaCompany Report #B0085873A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG/TWICE	Drug Interaction	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - PER DAY/ORAL	Initial or Prolonged			Phenytoin Tablet (Phenytoin)	SS		ORAL
PER DAY/ORAL				Carbamazepine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/22/00ISR Number: 3555918-9Report Type:Expedited (15-DaCompany Report #B0085514A  
Age:62 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL	Gastrointestinal Disorder	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
	Inflammatory Bowel Disease		Ciprofloxacin	C		
	Mucosal Erosion		Amoxicillin	C		
	Rectal Haemorrhage		Augmentin	C		
			Prednisolone	C		

Date:08/22/00ISR Number: 3555919-0Report Type:Expedited (15-DaCompany Report #B0084724A  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - AS Initial or Prolonged DIRECTED/ORAL	Alopecia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
	Blood Pressure Fluctuation	Consumer	Allopurinol	C		
	Dermatitis Exfoliative					
	Erythema					
	Hypothermia					
	Malaise					
	Pigmentation Disorder					
	Skin Inflammation					

Date:08/22/00ISR Number: 3555932-3Report Type:Expedited (15-DaCompany Report #9947976  
Age:79 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 50.00 MG Required TOTAL: DAILY: Intervention to ORAL	Angiopathy	Consumer	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
	Arthralgia					
	Condition Aggravated					
	Delirium					

Prevent Permanent	Diarrhoea	Aricept	SS	ORAL
10.00 MG				
Impairment/Damage	Drooling			
TOTAL: DAILY:				
	Fall			
ORAL				
	Hallucination	Wellbutrin	SS	
	Hip Fracture	Iron	C	
	Insomnia	Lanoxin	C	
	Pneumonia	Vitamin E	C	
		Baby Aspirin	C	
		Centrum Silver	C	
		Unknown		
		Laxative/Stool		
		Softener	C	
		Sinemet Cr	C	
		Glucosamine	C	
		Vitamin B12	C	

Date:08/23/00ISR Number: 3556113-XReport Type:Direct Company Report #  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Decreased Appetite		Wellbutrin 150mg Sr	PS		ORAL
150MG PO BID						
Initial or Prolonged	Dizziness					
	Fall					
	Head Injury					
	Lethargy					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/24/00ISR Number: 3557611-5Report Type:Expedited (15-DaCompany Report #044-0073-M0000016

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Interaction	Foreign Health Professional	Dilantin	PS	Parke Davis Div Warner Lambert Co	ORAL
1 TABLET(S)							
(DAILY), PER							
ORAL							
150 MG				Zyban (Amfebutamone Hydrochloride)	SS		ORAL
(DAILY), PER							
ORAL				(Carbamazepine)	C		

Date:08/24/00ISR Number: 3557660-7Report Type:Expedited (15-DaCompany Report #PRIUSA2000006683

Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Haldol	PS	Rw Johnson Pharmaceutical Research Institute Div Ortho Pharm	ORAL
Hospitalization - Initial or Prolonged		Electrocardiogram Qrs Complex Prolonged Grand Mal Convulsion					
ORAL							
ORAL		Intentional Misuse		Amitriptyline	SS		ORAL
ORAL		Intentional Self-Injury		Bupropion (Amfebutamone)	SS		ORAL

Date:08/24/00ISR Number: 3557738-8Report Type:Expedited (15-DaCompany Report #A0124755A

Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Life-Threatening 200 MG TWICE Hospitalization - PER DAY ORAL Initial or Prolonged	Amnesia Convulsion Insomnia Intentional Misuse Loss Of Consciousness Suicide Attempt Tympanic Membrane	Health Professional	Wellbutrin  Venlafaxine Hydrochloride (Formulation Unknown) (Venlafaxine	PS  SS	Glaxo Wellcome Inc	ORAL
AT NIGHT	Perforation Weight Increased		Olanzapine (Formulation Unknown) (Olanzapine) Ibuprofen (Formulation Unknown) (Ibuprofen) Percocet	SS SS SS C		

Date:08/24/00ISR Number: 3557817-5Report Type:Expedited (15-DaCompany Report #A0125417A  
Age: Gender:Not Specified/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
Other		Convulsion	Professional	Tramadol			
UNKNOWN ORAL			Company Representative	Hydrochloride	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/25/00ISR Number: 3558980-2Report Type:Expedited (15-DaCompany Report #A021954  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to ORAL		Decreased Appetite Difficulty In Walking	Consumer Health	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
Prevent Permanent ORAL Impairment/Damage		Emotional Disorder	Professional	Wellbutrin	SS		ORAL
		Gait Disturbance		Tnp-470 (Experimental Drug)	C		
		Halo Vision		Taxol	C		
		Hypoaesthesia		Flecainide	C		
		Insomnia		Ritalin	C		
		Neuropathy Peripheral		Restoril	C		
		Paraesthesia					
		Tearfulness					
		Tunnel Vision					
		Vision Blurred					
		Vomiting					
		Weight Decreased					

Date:08/28/00ISR Number: 3560528-3Report Type:Expedited (15-DaCompany Report #B0086215A  
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death ORAL		Circulatory Collapse	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Myalgia	Consumer				
		Myocardial Infarction					

Date:08/28/00ISR Number: 3560530-1Report Type:Expedited (15-DaCompany Report #B0086401A  
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / PER Initial or Prolonged DAY / ORAL		Crepitations	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Cyanosis					
		Depressed Level Of Consciousness		Paroxetine	C		
				Temazepam	C		

Dyspnoea  
Left Ventricular Failure  
Respiratory Failure

Aldactide C  
Combivent C  
Movicol C

Date:08/28/00ISR Number: 3560537-4Report Type:Expedited (15-DaCompany Report #A0126045A  
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Epilepsy	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG/AT		Sudden Death	Professional				
NIGHT/ORAL							

Carbamazepine C  
Sertraline  
Hydrochloride C  
Methylphenidate Hcl C

Date:08/28/00ISR Number: 3560740-3Report Type:Expedited (15-DaCompany Report #B0086031A  
Age:43 YR Gender:Female I/FU:I

Outcome	PT
Other	Depression Headache Oedema Peripheral Physical Assault

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Suicide Attempt

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1		Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TABLET/TWICE						
PER DAY/ ORAL						

Date:08/29/00ISR Number: 3561543-6Report Type:Expedited (15-DaCompany Report #A0113937A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Arthralgia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
Intervention to		Serum Sickness	Professional				
PER DAY/ORAL							
Prevent Permanent		Urticaria					
Impairment/Damage		Vision Blurred					

Date:08/29/00ISR Number: 3561657-0Report Type:Expedited (15-DaCompany Report #A0104240A  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Abnormal Behaviour	Study	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
400 MG/ PER							
Hospitalization -		Aggression	Health				
DAY/ ORAL							
Initial or Prolonged		Mania	Professional	Paracetamol	C		
Required		Physical Assault					
Intervention to		Suicidal Ideation					
Prevent Permanent							
Impairment/Damage							

Date:08/30/00ISR Number: 3562025-8Report Type:Expedited (15-DaCompany Report #D0009801A  
Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening TWICE PER DAY Hospitalization - / ORAL Initial or Prolonged		Cheyne-Stokes Respiration  Dysarthria  Respiratory Depression Sedation	Foreign  Health  Professional	Zyban    Allopurinol Digitoxin Torasemide Molsidomine Dipyridamole Potassium Chloride Phenprocoumon Heparin Sodium	PS    C C C C C C C C	Glaxo Wellcome Inc	ORAL

Date:08/30/00ISR Number: 3562120-3Report Type:Expedited (15-DaCompany Report #US0013200  
Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 225 MG DAILY  20 MG IN THE MORNING	1 YR  4 MON	Death	Health  Professional	Wellbutrin  Adderall	PS  SS	Shire Richwood Inc	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/00ISR Number: 3565698-9Report Type:Periodic  
Age:63 YR Gender:Male I/FU:I

Company Report #A0116206A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE			Professional				
PER DAY ORAL				Triamcinoline	C		

Date:08/31/00ISR Number: 3565708-9Report Type:Periodic  
Age:62 YR Gender:Male I/FU:I

Company Report #A0116210A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Heat Rash	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Pruritus					
DAY ORAL				Etodolac	C		
				Doxazosin Mesylate	C		
				Aspirin	C		

Date:09/01/00ISR Number: 3562866-7Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Circulatory Collapse		Remeron	PS		ORAL
Required							
60MG PO Q HS		Pulse Absent		Wellbutrin Sr	SS		ORAL
Intervention to							
150MG PO TID				Neurontin	SS		ORAL
Prevent Permanent							
600MG 1 PO							
Impairment/Damage							
QAM AMD 2 PO							
Q HS							
				Lamictal	SS		ORAL
200MG PO BID							
				Cytomel	SS		
25MCG PO QD							

Date:09/01/00ISR Number: 3564170-XReport Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #A0116991A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / PER Initial or Prolonged DAY / ORAL Required Intervention to Prevent Permanent Impairment/Damage	Convulsion	Study Health Professional	Zyban  Conjugated Estrogens Antioxidants	PS  C C	Glaxo Wellcome Inc	ORAL

Date:09/01/00ISR Number: 3564171-1Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0121024A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG ORAL	Bronchospasm Dizziness  Eyelid Oedema Face Oedema Hypersensitivity Intentional Misuse Nausea Oedema Peripheral Pharyngeal Oedema	Health Professional	Bupropion Hydrochloride  Dicyclomine Hydrochloride Ethanol	PS  C C	Glaxo Wellcome Inc	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/01/00ISR Number: 3564172-3Report Type:Periodic  
Age:31 YR Gender:Male I/FU:I

Company Report #A0121050A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Urticaria	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL	2 WK		Professional				

Date:09/01/00ISR Number: 3564173-5Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #A0121379A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
Initial or Prolonged		Hypertension					
TWICE PER							
DAY/ ORAL							

Date:09/01/00ISR Number: 3564175-9Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #A0122019A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyskinesia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /			Professional				
TWICE PER DAY							
/ ORAL	1 YR						

Date:09/01/00ISR Number: 3564177-2Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0123999A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							

Initial or Prolonged Constipation  
TWICE PER  
DAY/ ORAL  
Dizziness  
Hypoaesthesia  
Insomnia  
Paraesthesia  
Pollakiuria

Vitamin C

Date:09/01/00ISR Number: 3564178-4Report Type:Periodic Company Report #A0124881A  
Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG / TWICE PER DAY /ORAL	3 WK	Arthralgia Oedema Peripheral Rash Pruritic Urticaria	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:09/01/00ISR Number: 3564180-2Report Type:Periodic Company Report #A0124945A  
Age:26 YR Gender:Female I/FU:I

Outcome  
Other  
PT  
Arthralgia  
Candidiasis  
Decreased Activity  
Dermatitis  
Difficulty In Walking  
Erythema  
Face Oedema  
Insomnia  
Oedema Peripheral

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Pruritus Urticaria				
Dose	Duration		Report Source	Product	Role	Manufacturer
150 MG /			Consumer	Zyban	PS	Glaxo Wellcome Inc
TWICE PER						ORAL
DAY/ ORAL				Medroxyprogesterone Ace.	C	

Date:09/01/00ISR Number: 3564181-4Report Type:Periodic Company Report #A0124983A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Zyban	PS	Glaxo Wellcome Inc	

Date:09/01/00ISR Number: 3564184-XReport Type:Periodic Company Report #A0109719A  
 Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - ORAL		Akinesia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged Disability		Dystonia	Professional	Venlafaxine Hydrochloride	C		
Other Required Intervention to Prevent Permanent Impairment/Damage							

Date:09/01/00ISR Number: 3564186-3Report Type:Periodic Company Report #A0116361A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anaphylactic Shock	Health	Zyban	PS	Glaxo Wellcome Inc	

Initial or Prolonged      Dermatitits      Professional  
Face Oedema  
Hypersensitivity  
Oedema Peripheral  
Urticaria  
Vomiting

Date:09/01/00ISR Number: 3564189-9Report Type:Periodic      Company Report #A0104586A  
Age:60 YR      Gender:Male      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Convulsion					
TWICE PER DAY		Weight Decreased					
/ ORAL							

Date:09/01/00ISR Number: 3564190-5Report Type:Periodic      Company Report #A0109688A  
Age:22 YR      Gender:Female      I/FU:I

Outcome	PT
Other	Back Pain
	Depression
	Fatigue
	Hypoaesthesia



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Migraine						
Nausea						
Vision Blurred						
Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG /		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY						
/ ORAL						

Date:09/01/00ISR Number: 3564192-9Report Type:Periodic Company Report #A0109834A  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Arthritis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL						
Initial or Prolonged	Joint Stiffness					

Date:09/01/00ISR Number: 3564193-0Report Type:Periodic Company Report #A0112059A  
 Age:37 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Bronchospasm	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /						
Initial or Prolonged	Dermatitis	Professional				
TWICE PER DAY						
Required	Dyspnoea	Company				
/ ORAL						
Intervention to	Face Oedema	Representative				
Prevent Permanent	Hyperhidrosis					
Impairment/Damage	Hypersensitivity					
	Pruritus					
	Rash Papular					
	Throat Tightness					
	Tongue Oedema					
	Urticaria					

Date:09/01/00ISR Number: 3564196-6Report Type:Periodic Company Report #A0115264A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Chest Pain	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Dyspnoea Pruritus Rash Pruritic					

Date:09/01/00ISR Number: 3564198-XReport Type:Periodic Company Report #A0116901A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Consumer	Zyban	PS	Glaxo Wellcome Inc	

Date:09/01/00ISR Number: 3577814-3Report Type:Periodic Company Report #A0121399A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 MG / TWICE PER DAY / ORAL		Agitation Frequent Bowel Movements Muscle Spasms	Consumer	Lotronex  Citalopram Hydrobromide (Formulation Unknown) (Citalopram	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL				Hydrobromide)	SS		
				Wellbutrin			
				Tablet-Controlled			
				Release (Bupropion			
				Hydrochloride)	SS		ORAL

Date:09/05/00ISR Number: 3565455-3Report Type:Expedited (15-DaCompany Report #A0126869A  
 Age:58 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100 MG/ TWICE Initial or Prolonged PER DAY/	Dizziness Headache Orthostatic Hypotension	Consumer	Lamictal	PS	Glaxo Wellcome Inc	
150 MG/ TWICE PER DAY/ PO			Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	SS		ORAL
300 MG / PER DAY/			Amitriptyline Hcl (Formulation Unknown) (Amitriptyline Hcl)	SS		
			Clonazepam Thyroxine Sodium Mirtazapine Zolpidem Tartrate	C C C C		

Date:09/08/00ISR Number: 3568263-2Report Type:Direct  
 Age:10 YR Gender:Male I/FU:I Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required Intervention to SEE IMAGE Prevent Permanent	Abnormal Behaviour Convulsion Hyperhidrosis	Health Professional	Wellbutrin 150mg + 75mg Glaxo Wellcome Acetaminophen	PS C	Glaxo Wellcome	

Impairment/Damage	Lethargy	Milk Of Magnesia	C
	Loose Associations	Maalox Suspension	C
	Loss Of Consciousness	Hydroxyzine Pamoate	C
	Nervousness	Hydroxyzine	C
		Lithobid	C
		Eskalith	C
		Depakote	C

Date:09/11/00ISR Number: 3569205-6Report Type:Expedited (15-DaCompany Report #A0126585A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pulmonary Hypertension	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
TWICE PER			Professional				
DAY/ ORAL				Nefazodone Hydrochloride	C		

Date:09/11/00ISR Number: 3569344-XReport Type:Expedited (15-DaCompany Report #B0086888A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /SEE							
TEXT/ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/11/00ISR Number: 3569345-1Report Type:Expedited (15-DaCompany Report #B0086885A  
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG / AS DIRECTED/ ORAL		Asthma Bronchospasm Drug Interaction Dyspnoea	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
50 MG /PER DAY / ORAL				Ramipril (Formulation Unknown) (Ramipril)	SS		ORAL

Date:09/11/00ISR Number: 3569346-3Report Type:Expedited (15-DaCompany Report #B0086215A  
Age:67 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death ORAL		Circulatory Collapse Myalgia Myocardial Infarction	Foreign Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:09/11/00ISR Number: 3569347-5Report Type:Expedited (15-DaCompany Report #B0086827A  
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / PER Initial or Prolonged DAY / ORAL		Myocardial Ischaemia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Omeprazole Diclofenac	C C		

Date:09/11/00ISR Number: 3569348-7Report Type:Expedited (15-DaCompany Report #B0083476A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG /TWICE Initial or Prolonged PER DAY /		Headache	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL		Hemianopia	Health				
		Malignant Hypertension	Professional				
		Vision Blurred		Diane-35	C		

Date:09/11/00ISR Number: 3569349-9Report Type:Expedited (15-DaCompany Report #B0085154A  
Age:67 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG /TWICE Hospitalization - PER DAY /		Cardiac Arrest	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged ORAL		Cardiac Enzymes Increased	Consumer				
ORAL		Circulatory Collapse					
		Drug Interaction Myalgia		Phenytoin (Phenytoin)	SS		ORAL
		Ventricular Fibrillation		Carbamazepine	C		

Date:09/11/00ISR Number: 3569350-5Report Type:Expedited (15-DaCompany Report #B0085873A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG /TWICE Hospitalization - PER DAY /		Drug Interaction	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged ORAL							
				Phenytoin Tablet			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

PER DAY/  
ORAL  
(Phenytoin) SS ORAL

Carbamazepine C

Date:09/12/00ISR Number: 3570356-0Report Type:Expedited (15-DaCompany Report #A0127152A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Level Above Therapeutic Overdose	Health Professional	Wellbutrin Sertraline Hydrochloride (Sertraline Hydrochloride)	PS	Glaxo Wellcome Inc	
					SS		

Date:09/12/00ISR Number: 3570586-8Report Type:Expedited (15-DaCompany Report #B0086864A  
Age:75 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening ORAL		Acute Respiratory	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - Initial or Prolonged		Distress Syndrome Bronchospasm Dyspnoea Hyperhidrosis Inspiratory Capacity Decreased	Health Professional	Prednisolone Beclomethasone Dipropion Salmeterol Xinafoate Gallopamil Hydrochloride Nitroglycerin Dipyridamole	C C C C C C		

Date:09/12/00ISR Number: 3570587-XReport Type:Expedited (15-DaCompany Report #D0009801A  
Age:73 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150 MG / Hospitalization - TWICE PER DAY		Agitation Cerebral Ischaemia	Foreign Health	Zyban	PS	Glaxo Wellcome Inc	ORAL

Initial or Prolonged / ORAL  
Cheyne-Stokes Respiration Professional

Dysarthria  
Nervous System Disorder  
Respiratory Depression  
INTRA VENOUS SINGLE DOSE /  
Sedation  
INTRA VENOUS

Dipyridamole  
Injection 45 Mg  
(Dipyridamole) SS  
  
Allopurinol C  
Digitoxin C  
Torasemide C  
Molsidomine C  
Potassium Chloride C  
Phenprocoumon C  
Heparin Sodium C

Date:09/12/00ISR Number: 3570588-1Report Type:Expedited (15-DaCompany Report #B0086711A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Respiratory Alkalosis Tetany	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/12/00ISR Number: 3571006-XReport Type:Expedited (15-DaCompany Report #A0125417A

Age:39 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150 MG (TWICE Other PER DAY) ORAL Required Intervention to Prevent Permanent Impairment/Damage	Grand Mal Convulsion	Health  Professional  Company Representative	Wellbutrin Sr   Tramadol Hydrochloride Atorvastatin Calcium	PS   C C	Glaxo Wellcome Inc	ORAL

Date:09/13/00ISR Number: 3570377-8Report Type:Direct

Company Report #

Age:39 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150MG BID  ORAL	Dermatitis Erythema Multiforme  Eyelid Oedema  Myalgia Oedema Rash Generalised Tongue Oedema		Zyban 150mg Glaxo Wellcome  Nicotine Patch	PS  C	Glaxo Wellcome	ORAL

Date:09/13/00ISR Number: 3570903-9Report Type:Direct

Company Report #

Age:35 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose BID PO 3 WK	Urticaria		Zyban	PS		ORAL

Date:09/14/00ISR Number: 3570904-0Report Type:Direct

Company Report #

Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Creatine Phosphokinase Increased Blood Creatinine Increased Chromaturia Coma Dialysis Grand Mal Convulsion Intentional Misuse Rhabdomyolysis		Wellbutrin	PS		

Date:09/14/00ISR Number: 3571728-0Report Type:Expedited (15-DaCompany Report #10519379  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Balance Disorder Dizziness Drug Interaction	Foreign Health Professional	Serzone	PS	Bristol Myers Squibb Co Pharmaceutical Research Institute	ORAL

100  
MILLIGRAMS,  
2/1 DAY ORAL

150 MILLIGRAMS, 2/1 DAY ORAL				Zyban (Bupropion Hcl)	SS		ORAL
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Cipramil (Citalopram

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C  
 Predsol  
 (Prednisolone) C

Date:09/14/00ISR Number: 3571752-8Report Type:Expedited (15-DaCompany Report #A0127330A  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1 TABLET/ SEE Initial or Prolonged TEXT/ ORAL		Overdose	Health Professional	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL

Date:09/15/00ISR Number: 3572872-4Report Type:Expedited (15-DaCompany Report #B0087305A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Alcohol Increased Completed Suicide Drug Abuser	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	

Date:09/15/00ISR Number: 3572873-6Report Type:Expedited (15-DaCompany Report #B0087742A  
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG /AS DIRECTED/ ORAL		Aphonia Dysphonia Squamous Cell Carcinoma Vocal Cord Disorder	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:09/15/00ISR Number: 3572874-8Report Type:Expedited (15-DaCompany Report #B0087339A  
 Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Dyspnoea Exacerbated	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Left Ventricular Failure Myocardial Infarction					

Date:09/15/00ISR Number: 3572875-XReport Type:Expedited (15-DaCompany Report #B0087727A  
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL		Atrial Fibrillation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Bradycardia					
		Electrocardiogram St Segment Elevation Nodal Arrhythmia Pericardial Effusion Pericarditis Sick Sinus Syndrome		Bendroflurazide	C		

Date:09/18/00ISR Number: 3574699-6Report Type:Expedited (15-DaCompany Report #A0125270A  
Age:48 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Activated Partial Thromboplastin Time Prolonged

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Level Above Therapeutic	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Grand Mal Convulsion International Normalised Ratio Increased Pneumonia	Foreign Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
60 MG / PER DAY / ORAL		Sepsis Therapeutic Agent Toxicity		Paroxetine Hydrochloride (Paroxetine Hydrochloride)	SS		ORAL
.125 MG / PER DAY /				Lanoxin (Digoxin)	SS		
2.5 MG / PER DAY / ORAL				Warfarin Sodium (Warfarin Sodium)	SS		ORAL
				Zafirlukast	C		
				Prednisone	C		
				Salmeterol + Fluticasone	C		
				Beclomethasone Dipropion.	C		
				Prochlorperazine	C		
				Alprazolam	C		
				Codeine Phosphate	C		
				Enalapril Maleate	C		
				Lorazepam	C		
				Nitroglycerin	C		
				Salbutamol Sulphate	C		
				Temazepam	C		
				Hycodan	C		
				Amitriptyline	C		
				Ipratropium Bromide	C		
				Salbutamol Sulphate	C		

Date:09/18/00ISR Number: 3574701-1Report Type:Expedited (15-DaCompany Report #A0127464A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Foreign	Bupropion			
Other		Depression	Consumer	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Dysgeusia					
DAY / ORAL		Self Mutilation		Multivitamin	C		
		Suicidal Ideation		Hypericum	C		
				Ethanol	C		

Date:09/18/00ISR Number: 3574703-5Report Type:Expedited (15-DaCompany Report #B0084724A  
Age:54 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Alopecia
Initial or Prolonged	Dermatitis Exfoliative
Disability	Eosinophilia
Required	Hypersensitivity
Intervention to	Hypothermia
Prevent Permanent	Malaise
Impairment/Damage	Pigmentation Disorder
	Rash Erythematous
	Skin Inflammation

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Skin Lesion				
		Viral Infection				
Dose	Duration		Report Source	Product	Role	Manufacturer
			Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc
150 MG /			Professional			ORAL
TWICE PER DAY						
/ ORAL				Allopurinol	C	

Date:09/19/00ISR Number: 3573345-5Report Type:Direct  
 Age:40 YR Gender: I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Grand Mal Convulsion		Zyban 150 Mgs Glaxo Wellcome	PS	Glaxo Wellcome	ORAL
Initial or Prolonged							
1 TAB TWICE							
Disability							
DAILY ORAL							

Date:09/21/00ISR Number: 3576886-XReport Type:Expedited (15-DaCompany Report #A0122712A  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Condition Aggravated	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
Initial or Prolonged		Pyoderma Gangrenosum	Professional				
PER DAY/ ORAL							
Required				Azathioprine	C		
Intervention to				Metoprolol	C		
Prevent Permanent				Prednisone	C		
Impairment/Damage				Nortriptyline Hcl	C		
				Acyclovir	C		
				Fluoxetine			
				Hydrochloride	C		

Date:09/21/00ISR Number: 3576891-3Report Type:Expedited (15-DaCompany Report #A0127813A  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Potassium Decreased	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Convulsion					
DAY/ORAL		Fall		Potassium Chloride	C		
		Fatigue		Multivitamin	C		
		Grand Mal Convulsion					
		Headache					
		Muscle Rigidity					
		Pain					
		Skin Discolouration					

Date:09/21/00ISR Number: 3576925-6Report Type:Expedited (15-DaCompany Report #B0087347A  
Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fall	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL		Femoral Neck Fracture	Other	Paracetamol	C		
				Dihydrocodeine			
				Tartrate	C		
				Indomethacin	C		
				Iron Sulfate	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/22/00ISR Number: 3578342-1Report Type:Expedited (15-DaCompany Report #B0087889A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
SEE TEXT /		Suicide Attempt					
ORAL							
				Dihydrocodeine+Paracetam.	C		
				Cimetidine	C		
				Diazepam	C		

Date:09/22/00ISR Number: 3578343-3Report Type:Expedited (15-DaCompany Report #B0087719A  
Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Gastrointestinal	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Angiodysplasia					
Initial or Prolonged		Haemorrhagic					
PER DAY /		Haemoglobin Decreased		Aspirin	C		
ORAL				Ramipril	C		

Date:09/22/00ISR Number: 3578344-5Report Type:Expedited (15-DaCompany Report #B0087304A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Optic Nerve Disorder	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Vision Blurred	Study				
Initial or Prolonged		Visual Acuity Reduced	Health				
TWICE PER DAY		Visual Disturbance	Professional	Citalopram			
/ ORAL				Hydrobromide	C		

Date:09/25/00ISR Number: 3579498-7Report Type:Expedited (15-DaCompany Report #2000AP04229  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 50 MG DAILY		Chest Pain Drug Interaction	Foreign Health Professional	Tenormin	PS	Astrazeneca Pharmaceuticals Lp	ORAL
PO 150 MG BID PO			Other	Bupropion Hydrochloride	SS		ORAL

Date:09/25/00ISR Number: 3579650-0Report Type:Expedited (15-DaCompany Report #2000030151GB  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 500 MG , TID, ORAL		Cerebral Artery Thrombosis	Foreign Health Professional	Cyklokapron	PS	Pharmacia And Upjohn Co	ORAL
150 MG, QD, ORAL			Other	Bupropion (Amfebutamone)	SS		ORAL
				Fluoxetine (Fluoxetine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/26/00ISR Number: 3580490-7Report Type:Expedited (15-DaCompany Report #A0127997A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG/ TWICE	Completed Suicide	Foreign	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
	PER DAY/ORAL	Drug Level Above	Health				
		Therapeutic	Professional	Olanzapine	SS		
		Intentional Misuse	Other	Fluoxetine	SS		
		Toxicologic Test Abnormal		Lorazepam	C		
				Haloperidol	C		

Date:09/26/00ISR Number: 3580558-5Report Type:Expedited (15-DaCompany Report #A0127864A

Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG /	Amnesia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged	TWICE DAY /	Hypotension					
	ORAL	Pneumonia					
		Sedation		Nifedipine			
		Speech Disorder		(Formulation			
		Urine Analysis Abnormal		Unknown)			
				(Nifedipine)	SS		
				Atenolol			
				(Formulation			
				Unknown) (Atenolol)	SS		
				Gabapentin			
				(Formulation			
				Unknown)			
				(Gabapentin)	SS		

2 YR

Date:09/28/00ISR Number: 3581577-5Report Type:Direct

Company Report #

Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Other Swelling Zyban 150mg Glaxo PS Glaxo  
 OROPHARINGEAL 150MG 3  
 Urticaria  
 300MGF OR 16  
 OROPHARINGEAL

Prilosec C  
 Diovan C  
 Pulmicort C  
 Rhinocort C

Date:09/28/00ISR Number: 3582972-0Report Type:Expedited (15-DaCompany Report #A0114445A  
 Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Cognitive Disorder	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
		Depersonalisation Electroencephalogram Abnormal Emotional Disorder Feeling Abnormal Feeling Hot Headache Nervous System Disorder Pyrexia Vomiting Weight Decreased		Lamotrigine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/28/00ISR Number: 3583029-5Report Type:Expedited (15-DaCompany Report #A0128110A

Age:36 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY/ ORAL	Adrenal Neoplasm Blood Catecholamines Increased Epinephrine Increased Hypertension	Health Professional	Wellbutrin Sr  Antidepressant Blood Pressure Medication	PS  C C	Glaxo Wellcome Inc	ORAL

Date:09/28/00ISR Number: 3583031-3Report Type:Expedited (15-DaCompany Report #A0128188A

Age:63 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ORAL	Aggression Amnesia Atrial Fibrillation Constipation Coordination Abnormal Depression Difficulty In Walking Disorientation Dysarthria Fatigue Insomnia Libido Decreased Loss Of Consciousness Malaise Restlessness Sexual Dysfunction Sinus Congestion Speech Disorder Thyroid Disorder Urinary Tract Infection	Consumer	Wellbutrin Sr  Warfarin Sodium Lorazepam Zolpidem Tartrate Temazepam Liothyronine Sodium Gabapentin	PS  C C C C C C	Glaxo Wellcome Inc	ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Foreign	Bupropion			
		Metabolic Acidosis	Health	Hydrochloride	PS	Glaxo Wellcome Inc	
50 TABLET		Overdose	Professional				

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain Upper	Consumer	Zyban 150mg Glaxo			
		Dyspnoea		Wellcome Inc	PS	Glaxo Wellcome Inc	ORAL
WEEK 1 1		Urticaria					
TABLET WEEK 2							
2 TABLET							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/29/00ISR Number: 3584460-4Report Type:Expedited (15-DaCompany Report #A0070194A

Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG/ PER	Amnesia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY/ ORAL		Anger	Professional				
		Anxiety		General Anesthetic			
		Anxiety Disorder		(Formulation			
		Asthenia		Unknown) (General			
		Brain Hypoxia		Anesthetic)	SS		
		Breast Fibrosis		Ibuprofen	C		
		Breast Hyperplasia		Loperamide			
		Breast Mass		Hydrochloride	C		
		Cognitive Disorder					
		Confusional State					
		Convulsion					
		Decreased Activity					
		Decreased Appetite					
		Depressed Mood					
		Depression					
		Disturbance In Attention					
		Drug Ineffective					
		Dysphoria					
		Early Morning Awakening					
		Encephalopathy					
		Fatigue					
		Fear					
		Feeling Abnormal					
		Fibrocystic Breast					
		Disease					
		Headache					
		Hodgkin'S Disease Nodular					
		Sclerosis Stage					
		Unspecified					
		Hypochondriasis					
		Initial Insomnia					
		Loss Of Libido					
		Mental Impairment					
		Mood Swings					
		Nausea					
		Needle Biopsy Site					
		Unspecified Abnormal					
		Suicidal Ideation					
		Thinking Abnormal					

Vomiting

Date:09/29/00ISR Number: 3584462-8Report Type:Expedited (15-DaCompany Report #A0128346A  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erectile Dysfunction	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
ORAL	2 MON	Semen Abnormal Spermatozoa Abnormal		Olanzapine (Formulation Unknown) (Olanzapine)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/29/00ISR Number: 3584468-9Report Type:Expedited (15-DaCompany Report #A0128376A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG/ SEE Hospitalization - TEXT/ ORAL Initial or Prolonged		Brain Herniation	Professional				
		Convulsion Dyspnoea Overdose Pupils Unequal Suicide Attempt Vaginal Haemorrhage Vasospasm	Company Representative				

Date:10/02/00ISR Number: 3585611-8Report Type:Expedited (15-DaCompany Report #B0088136A  
Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL		Asthenia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Circulatory Collapse	Health				
		Confusional State Eye Rolling Headache Loss Of Consciousness Pallor Salivary Hypersecretion	Professional				

Date:10/02/00ISR Number: 3585757-4Report Type:Expedited (15-DaCompany Report #A0125460A  
Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death PO		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - Initial or Prolonged		Fall Wrist Fracture	Professional Company Representative	Clozapine Lisinopril Docusate Sodium	C C C		

Date:10/02/00ISR Number: 3585759-8Report Type:Expedited (15-DaCompany Report #A0054084A  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Deafness Neurosensory Dermatitis	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE PER DAY/ PO		Dizziness Ear Discomfort Ear Disorder Hypersensitivity Labyrinthitis Nausea Nystagmus Pruritus Tympanic Membrane Disorder Urticaria Vertigo Vomiting					



Hospitalization - 150 MG /TWICE Initial or Prolonged PER DAY/ ORAL	Multiple Sclerosis Optic Nerve Disorder Vision Blurred Visual Acuity Reduced Visual Disturbance	Foreign Study Health Professional	Zyban  Citalopram Hydrobromide	PS  C	Glaxo Wellcome Inc	ORAL
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Date:10/02/00ISR Number: 3585863-4Report Type:Expedited (15-DaCompany Report #A0128480A  
Age:83 YR Gender:Female I/FU:I

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG/TWICE Initial or Prolonged PER DAY/ORAL	Hip Fracture Leukocytosis Liver Function Test Abnormal Respiratory Failure	Health Professional	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
			Alprazolam	C		
			Pulmocare	C		
			Vicodin	C		
			Digoxin	C		
			Megestrol Acetate	C		
			Heparin	C		
			Promod Supplement	C		
			Ranitidine			
			Hydrochloride	C		
			Frusemide	C		
			Potassium Chloride	C		
			Ferrous Sulfate	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/02/00ISR Number: 3585938-XReport Type:Expedited (15-DaCompany Report #B0088136A  
Age:20 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG, TWICE PER DAY, ORAL	Asthenia Circulatory Collapse Confusional State Dystonia Eye Rolling Headache Loss Of Consciousness Pallor Salivary Hypersecretion	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:10/02/00ISR Number: 3585939-1Report Type:Expedited (15-DaCompany Report #B0088158A  
Age:82 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Bundle Branch Block Left Supraventricular Tachycardia Tachycardia	Foreign	Bupropion Hydrochloride Ipratropium Bromide Fluticasone Propionate Terbutaline Losartan Potassium Aspirin	PS C C C C C	Glaxo Wellcome Inc	ORAL

Date:10/02/00ISR Number: 3586124-XReport Type:Expedited (15-DaCompany Report #WAES 00091109  
Age:22 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged SUBCUTANEOUS SC Other	Agitation Back Pain Coma Confusional State Depression	Health Professional	Flexeril Betaseron Unk Wellbutrin Unk Depakote Unk Ativan	PS SS SS SS C	Merck And Co Inc	

Diplopia  
 Drug Interaction  
 Drug Level Below  
 Therapeutic  
 Epilepsy  
 Grand Mal Convulsion  
 Hypokalaemia  
 Multiple Sclerosis  
 Peptic Ulcer  
 Pyrexia  
 Respiratory Failure  
 Scoliosis  
 Urinary Tract Infection

Potassium Chloride C

Date:10/03/00ISR Number: 3586823-XReport Type:Expedited (15-DaCompany Report #B0088627A  
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ORAL		Dermatitis	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Psoriasis		Calcipotriene	C		
				Fluticasone	C		
				Propionate	C		
				Allopurinol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/03/00ISR Number: 3586824-1Report Type:Expedited (15-DaCompany Report #B0088634A  
Age:69 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/PER Initial or Prolonged DAY/ORAL	Angina Pectoris	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Gaviscon	C		
			Nizatidine	C		
			Aspirin	C		
			Nifedipine	C		

Date:10/03/00ISR Number: 3586825-3Report Type:Expedited (15-DaCompany Report #B0088630A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/SEE Initial or Prolonged TEXT/ORAL	Neutropenia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
	Tonsillitis					
			Birth Control	C		

Date:10/03/00ISR Number: 3586917-9Report Type:Expedited (15-DaCompany Report #A0128274A  
Age:39 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG / TWICE PER DAY / ORAL	Anxiety	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
	Feeling Abnormal					
	Feeling Hot					
	Hyperaesthesia		Nicotine	C		
	Irritability					
	Nervousness					
	Nightmare					
	Oedema					
	Pruritus					
	Skin Discolouration					

Urticaria

Date:10/04/00ISR Number: 3586657-6Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Depression		Wellbutrin Sr	PS	Glaxo Wellcome	

Date:10/05/00ISR Number: 3589083-9Report Type:Expedited (15-DaCompany Report #A0125213A  
 Age:53 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 200 MG / Initial or Prolonged TWICE PER DAY	Balance Disorder Difficulty In Walking	Health Professional	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
/ ORAL	Fall Parkinsonian Gait Tremor		Valproic Acid Lithium Salt Venlafaxine Hydrochloride Clonazepam Glipizide Medroxyprogesterone Ace. Oestradiol	C C C C C C C C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/06/00ISR Number: 3590708-2Report Type:Expedited (15-DaCompany Report #A0127152A  
Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional	Wellbutrin Sertraline Hydrochloride (Sertraline Hydrochloride)	PS	Glaxo Wellcome Inc	
					SS		

Date:10/06/00ISR Number: 3590803-8Report Type:Expedited (15-DaCompany Report #D0010641A  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Arrhythmia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Headache Tachycardia	Health Professional				

Date:10/06/00ISR Number: 3590804-XReport Type:Expedited (15-DaCompany Report #B0086470A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Disability PER DAY/ ORAL		Atrial Fibrillation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Required Intervention to Prevent Permanent Impairment/Damage		Nausea	Professional	Combivent Beclomethasone Dipropion	C		
					C		

Date:10/06/00ISR Number: 3598782-4Report Type:Periodic Company Report #USA015005  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation	Consumer	Meridia	PS	Knoll Pharmaceutical	

10 MG OD PO	Central Nervous System	Other		Co Sub Basf Corp	ORAL
	Stimulation		Meridia	SS	ORAL
10 MG OD PO	Migraine		Wellbutrin	SS	ORAL
150 MG OD PO	Nausea				
	Therapeutic Response				
	Unexpected				

Date:10/10/00ISR Number: 3591925-8Report Type:Expedited (15-DaCompany Report #A0128493A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Arrest	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL			Professional				

Date:10/10/00ISR Number: 3592070-8Report Type:Expedited (15-DaCompany Report #D0010662A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Epilepsy	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL			Health				
Initial or Prolonged			Professional				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/10/00ISR Number: 3592695-XReport Type:Expedited (15-DaCompany Report #A0128801A  
 Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
200 MG/TWICE		Tremor	Professional				
PER DAY/ ORAL		Tunnel Vision		Citalopram Hydrobromide (Citalopram Hydrobromide)	SS		
60 MG/PER DAY				Lithium Salt Perphenazine	C C		

Date:10/11/00ISR Number: 3592416-0Report Type:Direct Company Report #  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Chest Pain		Accutane 40 Mg Roche	PS	Roche	
40 MG 1 A DAY		Constipation Crying Headache Hypoaesthesia		Zyban 150 Mg Catalytica For Glaxo Wellcome	SS	Catalytica For Glaxo Wellcome	
150MG 2 A DAY		Major Depression Musculoskeletal Stiffness Neck Pain Panic Attack Speech Disorder Suicidal Ideation Trismus					

Date:10/13/00ISR Number: 3605050-0Report Type:Expedited (15-DaCompany Report #A0130183A  
 Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Foreign	Wellbutrin	PS	Glaxo Wellcome Inc	

Atropine  
 (Formulation  
 Unknown) (Atropine) SS  
 Chloral Hydrate  
 (Formulation  
 Unknown) (Chloral  
 Hydrate) SS  
 Daizepam  
 (Formulation  
 Unknown) (Diazepam) SS

Date:10/13/00ISR Number: 3605051-2Report Type:Expedited (15-DaCompany Report #B0088745A  
 Age:49 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - ORAL	Duration Chest Pain	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged Disability	Electrocardiogram T Wave Inversion Myocardial Infarction	Health Professional	Ipratropium Bromide Salbutamol Sulphate Aspirin Bendrofluazide Prednisolone Beclomethasone Dipropion	C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/00ISR Number: 3595759-XReport Type:Expedited (15-DaCompany Report #A0128376A

Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG/ SEE	Brain Herniation	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - TEXT/ ORAL	Initial or Prolonged	Convulsion	Professional				
		Intubation	Company				
		Overdose	Representative				
		Pupils Unequal					
		Respiratory Failure					
		Suicide Attempt					
		Vasospasm					

Date:10/16/00ISR Number: 3595760-6Report Type:Expedited (15-DaCompany Report #A0070194A

Age:39 YR Gender:Female I/FU:F

Outcome	PT
Disability	Affective Disorder
	Amnesia
	Anger
	Anxiety
	Apathy
	Asthenia
	Biopsy Breast Abnormal
	Breast Hyperplasia
	Breast Mass
	Cognitive Disorder
	Confusional State
	Convulsion
	Decreased Appetite
	Depressed Mood
	Depression
	Disturbance In Attention
	Drug Ineffective
	Early Morning Awakening
	Encephalopathy
	Fatigue
	Fear
	Feeling Abnormal
	Fibroadenoma Of Breast
	Fibrocystic Breast
	Disease

Headache  
Hodgkin'S Disease Nodular  
Sclerosis Stage  
Unspecified  
Hypochondriasis  
Hypoxia  
Initial Insomnia  
Loss Of Libido  
Memory Impairment  
Mental Disorder  
Mental Impairment  
Mood Swings  
Nausea  
Nervous System Disorder  
Petit Mal Epilepsy  
Post Procedural  
Complication  
Suicidal Ideation  
Sweat Gland Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Thinking Abnormal Vomiting	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL			Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
				General Anesthetic (Formulation Unknown) (General Anesthetic)	SS		
				Ibuprofen	C		
				Surgery	C		
				Loperamide Hydrochloride	C		
				Dextrose + Lactated Ringers	C		
				Tylenol No. 3	C		
				Hyoscyamine Sulphate	C		
				Bupivacaine Hdydrochloride	C		
				Hydrocodone + Paracetamol	C		
				Propofol	C		
				Lignocaine	C		
				Fentanyl	C		
				Midazolam	C		
				Suxamethonium Chloride	C		
				Oxygen	C		
				Droperidol	C		
				Nitrous Oxide	C		

Date:10/16/00ISR Number: 3595783-7Report Type:Expedited (15-DaCompany Report #A0121698A  
 Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ ORAL		Convulsion	Literature	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged Other		Depressed Level Of Consciousness Grand Mal Convulsion Intentional Misuse	Health Professional				

Lethargy  
Status Epilepticus  
Suicide Attempt

Date:10/16/00ISR Number: 3595784-9Report Type:Expedited (15-DaCompany Report #A0122538A  
Age:54 YR Gender:Female I/FU:F

Outcome	PT
Life-Threatening	Cardiovascular Disorder
Hospitalization -	Coma
Initial or Prolonged	Hypoxic Encephalopathy
Disability	Lethargy
Required	Neuroleptic Malignant
Intervention to	Syndrome
Prevent Permanent	Overdose
Impairment/Damage	Renal Failure Acute
	Rhabdomyolysis
	Staring
	Stupor

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Urine Analysis Abnormal

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
			Opiate (Formulation Unknown) (Opiate) Benzodiazepines (Formulation Unknown) (Benzodiazepines)	SS SS		

Date:10/16/00ISR Number: 3595785-0Report Type:Expedited (15-DaCompany Report #A0093694A  
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ PER Initial or Prolonged DAY/ ORAL Disability	1 MON	Depression Grand Mal Convulsion Pain Spinal Fracture	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:10/16/00ISR Number: 3595788-6Report Type:Expedited (15-DaCompany Report #A0129437A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100 MG/ PER DAY/ ORAL	10 DAY	Abortion Spontaneous	Study Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:10/16/00ISR Number: 3595926-5Report Type:Expedited (15-DaCompany Report #B0089470A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							
				Ibuprofen	C		
				Dextroprop. +	C		
				Paracetamol	C		
				Dihydrocodeine+Parac	C		
				etam.			

Date:10/16/00ISR Number: 3595929-0Report Type:Expedited (15-DaCompany Report #B0089165A  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL		Pregnancy	Health				
			Professional				

Date:10/16/00ISR Number: 3595944-7Report Type:Expedited (15-DaCompany Report #B0089357A  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anxiety	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL		Athetosis	Health				
		Fear	Professional	Mefenamic Acid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Macrodex

C

Date:10/16/00ISR Number: 3595946-0Report Type:Expedited (15-DaCompany Report #B0089436A

Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL		Feeling Abnormal	Health Professional Company Representative				

Date:10/16/00ISR Number: 3595948-4Report Type:Expedited (15-DaCompany Report #B0089550A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Atrioventricular Block	Foreign	Zyban	PS	Glaxo Wellcome Inc	
2 MON		Second Degree	Health Professional	Diazepam Beclomethasone Dipropion	C C		

Date:10/16/00ISR Number: 3595952-6Report Type:Expedited (15-DaCompany Report #B0088136A

Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150 MG/ Initial or Prolonged TWICE PER DAY		Asthenia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
/ ORAL		Circulatory Collapse	Health				
		Confusional State	Professional				
		Eye Rolling Headache Loss Of Consciousness Pallor Salivary Hypersecretion					

Tonic Convulsion

Date:10/16/00ISR Number: 3595954-XReport Type:Expedited (15-DaCompany Report #B0088627A  
 Age:54 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ ORAL	Dermatitis	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged	Psoriasis		Calcipotriene	C		
			Fluticasone	C		
			Propionate	C		
			Allopurinol	C		

Date:10/16/00ISR Number: 3595956-3Report Type:Expedited (15-DaCompany Report #B0084724A  
 Age:54 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Alopecia
Initial or Prolonged	Blood Pressure
Disability	Fluctuation
Required	Dermatitis Exfoliative
Intervention to	Eosinophilia
Prevent Permanent	Erythema
Impairment/Damage	Hypersensitivity

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Hypothermia Malaise Rash Erythematous Skin Inflammation Skin Lesion Viral Infection	Foreign Health Professional	Zyban  Allopurinol	PS  C	Glaxo Wellcome Inc	ORAL

Date:10/16/00ISR Number: 3595958-7Report Type:Expedited (15-DaCompany Report #B0089466A  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG/ ORAL		Asthma	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Salbutamol Sulphate Beclomethasone Dipropion. Ipratropium Bromide Fluoxetine	C C C C		

Date:10/16/00ISR Number: 3595961-7Report Type:Expedited (15-DaCompany Report #B0082531A  
Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150 MG/ ORAL Hospitalization - Initial or Prolonged		Aspartate Aminotransferase Increased Atrioventricular Block Complete Blood Creatine Phosphokinase Increased Blood Creatine Phosphokinase Mb Increased Bradycardia Hypokinesia Myocardial Infarction Supraventricular	Foreign Literature Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Tachycardia

Date:10/16/00ISR Number: 3595963-0Report Type:Expedited (15-DaCompany Report #B0088823A  
Age:46 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL	Constipation Intestinal Obstruction	Foreign Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:10/16/00ISR Number: 3595965-4Report Type:Expedited (15-DaCompany Report #B0089068A  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death ORAL	Headache Ruptured Cerebral Aneurysm Subarachnoid Haemorrhage	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/00ISR Number: 3595966-6Report Type:Expedited (15-DaCompany Report #B0089121A  
Age:37 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ PER	Arthralgia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged DAY/ ORAL	Haematuria					
	Rash Maculo-Papular					

Date:10/16/00ISR Number: 3595967-8Report Type:Expedited (15-DaCompany Report #B0089166A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL	Aggression	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged	Emotional Disorder Fatigue Hallucination Nightmare Sleep Walking Stress	Consumer				

Date:10/16/00ISR Number: 3595968-XReport Type:Expedited (15-DaCompany Report #D0010841A  
Age:24 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150 MG/ PER	Amnesia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - DAY/ ORAL	Cardiac Arrest	Health				
Initial or Prolonged Required	Electroencephalogram Abnormal	Professional				
Intervention to Prevent Permanent Impairment/Damage	Epilepsy					

Date:10/16/00ISR Number: 3595970-8Report Type:Expedited (15-DaCompany Report #D0010722A  
Age:23 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL	Multiple Sclerosis	Foreign  Health  Professional Company Representative	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:10/16/00ISR Number: 3595974-5Report Type:Expedited (15-DaCompany Report #B0089112A  
Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL	Chest Pain  Circulatory Collapse  Hyperhidrosis Syncope Vasovagal	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:10/16/00ISR Number: 3595976-9Report Type:Expedited (15-DaCompany Report #B0089069A  
Age:62 YR Gender:Male I/FU:I

Outcome  
Death  
Hospitalization -  
  
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Initial or Prolonged

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL			Brain Stem Infarction	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Circulatory Collapse		Diclofenac Allopurinol	C C		

Date:10/16/00ISR Number: 3596416-6Report Type:Expedited (15-DaCompany Report #A0128949A  
Age:28 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		1 WK	Abortion Spontaneous	Foreign  Study Health Professional	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL

Date:10/17/00ISR Number: 3595457-2Report Type:Expedited (15-DaCompany Report #A0129577A  
Age:50 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG TWICE PER DAY			Erectile Dysfunction		Zyban	PS	Glaxo Wellcome	ORAL
			Hypoaesthesia					
			Triple Vessel Bypass Graft		Cardiac Medication	SS		

Date:10/17/00ISR Number: 3595454-7Report Type:Expedited (15-DaCompany Report #A0129318A  
Age:14 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG PER DAY			Dermatitis		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged			Dyspnoea Eyelid Oedema		Wellbutrin No Concurrent	C	Glaxo Wellcome	ORAL

Malaise  
Medication Error  
Pruritus

Medications

C

Date:10/17/00ISR Number: 3595459-6Report Type:Expedited (15-DaCompany Report #B0089482A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG TWICE PER DAY	20 DAY	Left Ventricular Failure Oedema		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL

No Concurrent  
Medication

C

Date:10/18/00ISR Number: 3597615-XReport Type:Expedited (15-DaCompany Report #A0129318A  
Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG/PER Initial or Prolonged DAY/ORAL		Dermatitis Eyelid Oedema	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Malaise  
Medication Error  
Pruritus  
Respiratory Disorder

Bupropion  
Hydrochloride

C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/18/00ISR Number: 3598141-4Report Type:Expedited (15-DaCompany Report #A0129577A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG/TWICE PER DAY/ORAL		Erectile Dysfunction Hypoaesthesia  Sexual Dysfunction  Triple Vessel Bypass Graft	Foreign Consumer	Bupropion Hydrochloride   Cardiac Medication (Formulation Unknown) (Cardiac Medication)	PS    SS	Glaxo Wellcome Inc	ORAL

Date:10/18/00ISR Number: 3598158-XReport Type:Expedited (15-DaCompany Report #B0089482A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to 150 MG/TWICE Prevent Permanent PER DAY/ORAL Impairment/Damage		Left Ventricular Failure Oedema	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:10/20/00ISR Number: 3598180-3Report Type:Expedited (15-DaCompany Report #A0128376A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 150MG SEE Hospitalization - TEXT 1 DAY Initial or Prolonged		Agitation  Blood Gases Abnormal Brain Herniation Brain Oedema Bundle Branch Block Left Cerebrovascular Accident Coma Confusional State Convulsion		Wellbutrin   Zyban Tylenol Pm	PS   SS C	Glaxo Wellcome	ORAL

Dyspnoea  
 Extensor Plantar Response  
 Hemiparesis  
 Hypokalaemia  
 Lung Infiltration  
 Muscle Twitching  
 Mydriasis  
 Overdose  
 Pulmonary Oedema  
 Pupillary Reflex Impaired  
 Pupils Unequal  
 Respiratory Disorder  
 Respiratory Failure  
 Sinus Tachycardia  
 Suicide Attempt

Date:10/20/00ISR Number: 3598887-8Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain Upper		Zyban	PS		ORAL
150 MG PO BID 2	MON	Diarrhoea		Evista	C		
		Dyspepsia					
		Medication Error					
		Vomiting					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/20/00ISR Number: 3598928-8Report Type:Direct  
 Age:25 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - BID	PT		Zyban 150mg	PS		
Initial or Prolonged	Discomfort		Dicyclomine	C		

Date:10/20/00ISR Number: 3598182-7Report Type:Expedited (15-DaCompany Report #A0129538A  
 Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	PT		Wellbutrin Concurrent Medications	PS C	Glaxo Wellcome	

Date:10/20/00ISR Number: 3598188-8Report Type:Expedited (15-DaCompany Report #B0088713A  
 Age:26 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability 150MG TWICE PER DAY	PT		Zyban	PS	Glaxo Wellcome	ORAL
2 WK	Asthenia Blood Creatine Phosphokinase Increased Myositis Oedema Peripheral Pain					

Date:10/20/00ISR Number: 3598191-8Report Type:Expedited (15-DaCompany Report #B0089511A  
 Age:55 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150MG TWICE	PT		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
	Cardiac Arrest Diabetes Mellitus					

Ventricular Tachycardia

PER DAY	32	DAY				
75MG PER DAY			Aspirin	C		ORAL
50MG PER DAY			Atenolol	C		ORAL
40MG PER DAY			Frusemide	C		ORAL
10MG PER DAY			Lisinopril	C		ORAL
25MG PER DAY			Spironolactone	C		ORAL
600MG TWICE			Gemfibrozil	C		ORAL
PER DAY						

Date:10/23/00ISR Number: 3599909-0Report Type:Expedited (15-DaCompany Report #A0129538A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Electrocardiogram Qt Corrected Interval Prolonged	Health Professional Company Representative	Wellbutrin Sr	PS	Glaxo Wellcome Inc	

Date:10/23/00ISR Number: 3599911-9Report Type:Expedited (15-DaCompany Report #A0128376A  
 Age:40 YR Gender:Female I/FU:F

Outcome	PT
Death Hospitalization - Initial or Prolonged	Abnormal Behaviour Acute Respiratory Distress Syndrome

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / SEE TEXT / ORAL		Agitation Blood Gases Abnormal Brain Herniation Bundle Branch Block Left Cerebrovascular Accident	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
SEE TEXT / ORAL		Coma Convulsion Dry Throat Dyspnoea Extensor Plantar Response	Company Representative	Bupropion Hydrochloride Tablet - Zyban (Bupropion Hydrochloride)	SS		ORAL
		Hemiparesis Hypertension Lip Dry Lung Infiltration Miosis Muscle Twitching Mydriasis Overdose Pulmonary Oedema Pupillary Reflex Impaired Pupils Unequal Respiratory Disorder Respiratory Failure Sinus Tachycardia Suicide Attempt Vaginal Haemorrhage Vasospasm		Extra Strength Tylenol Pm	C		

Date:10/23/00ISR Number: 3600401-5Report Type:Expedited (15-DaCompany Report #B0089511A  
Age:55 YR Gender:Male I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Cardiac Arrest Diabetes Mellitus Ventricular Tachycardia	Foreign Other	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Aspirin	C
Atenolol	C
Frusemide	C
Lisinopril	C
Spirocholactone	C
Gemfibrozil	C

Date:10/23/00ISR Number: 3600402-7Report Type:Expedited (15-DaCompany Report #B0088713A  
 Age:26 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Asthenia	Foreign	Bupropion			
150 MG /		Blood Creatine	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Phosphokinase Increased	Professional				
/ ORAL	2 WK	Myositis	Other				
		Oedema Peripheral					
		Pain In Extremity					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/23/00ISR Number: 3599071-4Report Type:Expedited (15-DaCompany Report #A0129781A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Laryngeal Oedema		Lamictal	SS	Glaxo Wellcome	ORAL
		Mania					

Date:10/24/00ISR Number: 3599704-2Report Type:Expedited (15-DaCompany Report #B0089739A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Chest Pain		Bupropion			
		Chronic Obstructive Airways Disease		Hydrochloride Combivent	PS C	Glaxo Wellcome	ORAL
		Exacerbated Electrocardiogram T Wave Inversion					

Date:10/24/00ISR Number: 3600705-6Report Type:Expedited (15-DaCompany Report #JRFUSA2000004633  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death TRANSDERMAL	TRANSD	Coma	Literature	Duragesic	PS	Alza Corp	
		Drug Toxicity	Health Professional	Codeine (Codeine)	SS		
				Sertraline (Sertraline)	SS		
				Olanzapine (Olanzapine)	SS		
				Bupropion (Amfebutamone)	SS		

Date:10/24/00ISR Number: 3601103-1Report Type:Expedited (15-DaCompany Report #A0129781A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Laryngeal Oedema	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL						
	Mania	Professional Company	Lamictal Unspecified Tablet (Lamotrigine)	SS		ORAL
ORAL		Representative				

Date:10/24/00ISR Number: 3601897-5Report Type:Expedited (15-DaCompany Report #B0089987A  
Age:24 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG /TWICE Initial or Prolonged PER DAY/ORAL	Circulatory Collapse Hypoglycaemia Hypoglycaemic Coma	Foreign	Zyban  Perindopril Human Insulin	PS  C C	Glaxo Wellcome Inc	ORAL

Date:10/24/00ISR Number: 3599698-XReport Type:Expedited (15-DaCompany Report #A0129782A  
Age:22 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Bite
Initial or Prolonged	Convulsion Electroencephalogram

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Abnormal Intentional Misuse Suicide Attempt Toxicologic Test Abnormal	Report Source	Product	Role	Manufacturer	Route
2250MG SINGLE DOSE	1 DAY			Zyban	PS	Glaxo Wellcome	ORAL
150MG PER DAY	2 WK			Zyban	SS	Glaxo Wellcome	ORAL
1 DAY				Prenatal Vitamins Marijuana	C C		

Date:10/24/00ISR Number: 3599708-XReport Type:Expedited (15-DaCompany Report #B0089987A  
Age:24 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG TWICE Initial or Prolonged PER DAY			Circulatory Collapse		Bupropion	PS	Glaxo Wellcome	ORAL
2MG PER DAY			Hypoglycaemic Coma		Perindopril	C		ORAL
					Human Insulin	C		

Date:10/24/00ISR Number: 3599709-1Report Type:Expedited (15-DaCompany Report #B0090105A  
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Respiratory Distress		Zyban	PS	Glaxo Wellcome	

Date:10/25/00ISR Number: 3601561-2Report Type:Expedited (15-DaCompany Report #A0129782A  
Age:22 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Bite	Health	Bupropion			

Initial or Prolonged 2250	Complications Of Maternal	Professional	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
	Exposure To Therapeutic					
MG/SINGLE	Drugs					
DOSE/ORAL	Convulsion		Zyban Tablet - Zyban			
	Electroencephalogram		(Bupropion			
	Abnormal		Hydrochloride)	SS		ORAL
150 MG/PER	Intentional Misuse					
DAY/ORAL	Maternal Use Of Illicit		Prenatal	C		
	Drugs		Cannabis	C		
	Suicide Attempt					
	Tongue Disorder					
	Toxicologic Test Abnormal					

Date:10/25/00ISR Number: 3601934-8Report Type:Expedited (15-DaCompany Report #B0090105A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Respiratory Distress	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/25/00ISR Number: 3601936-1Report Type:Expedited (15-DaCompany Report #B0089739A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Angina Pectoris	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Cardiac Disorder Chest Pain Condition Aggravated Electrocardiogram T Wave Inversion Respiratory Disorder	Health Professional	Combivent	C		

Date:10/25/00ISR Number: 3600343-5Report Type:Expedited (15-DaCompany Report #A0129656A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG TWICE PER DAY		Chills Erythema Infection Pruritus Pyrexia Rash Maculo-Papular Skin Discolouration Skin Exfoliation Tremor		Zyban	PS	Glaxo Wellcome	ORAL

Date:10/25/00ISR Number: 3600349-6Report Type:Expedited (15-DaCompany Report #B0089929A  
Age:67 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG TWICE PER DAY	7 WK	Convulsion Eye Irritation Insomnia Keratoconjunctivitis Sicca		Zyban Cimetidine Temazepam	PS C C	Glaxo Wellcome	

Micturition Disorder

Date:10/25/00ISR Number: 3600350-2Report Type:Expedited (15-DaCompany Report #B0089984A  
 Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris		Bupropion	PS	Glaxo Wellcome	ORAL
150MG TWICE							
PER DAY	5	DAY					
				Atenolol + Chlorthalidone	C		ORAL
				Doxazosin	C		ORAL
2MG PER DAY							

Date:10/25/00ISR Number: 3600351-4Report Type:Expedited (15-DaCompany Report #B0089998A  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Oculogyration		Bupropion	PS	Glaxo Wellcome	ORAL
150MG PER DAY	2	DAY					
20MG PER DAY				Fluoxetine	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/26/00ISR Number: 3601007-4Report Type:Expedited (15-DaCompany Report #A0123011A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Hallucination		Zyban	PS	Glaxo Wellcome	ORAL
150MG TWICE		Hyperventilation					
PER DAY	7	DAY					
Overdose				Oxycontin	SS		ORAL
20MG TWICE		Paranoia					
PER DAY				Amantadine	C		ORAL
100MG THREE							
TIMES PER DAY							

Date:10/26/00ISR Number: 3601021-9Report Type:Expedited (15-DaCompany Report #B0089784A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Amnesia		Zyban	PS	Glaxo Wellcome	ORAL
150MG TWICE		Grand Mal Convulsion					
PER DAY	5	DAY					
Hypersensitivity				Losartan	C		ORAL
1TAB PER DAY		Thrombocythaemia		Bromazepam	C		ORAL
1TAB PER DAY							

Date:10/30/00ISR Number: 3602848-XReport Type:Expedited (15-DaCompany Report #A0130183A  
Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Toxicity		Bupropion	PS	Glaxo Wellcome	
UNKNOWN				Chloral Hydrate	SS		
UNKNOWN				Atropine	SS		

Diazepam

SS

UNKNOWN

Date:10/30/00ISR Number: 3602854-5Report Type:Expedited (15-DaCompany Report #A0130633A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris		Bupropion	PS	Glaxo Wellcome	ORAL
150MG PER DAY		Arrhythmia		Venlafaxine	SS		
UNKNOWN	375MG PER DAY						

Date:10/30/00ISR Number: 3602856-9Report Type:Expedited (15-DaCompany Report #A0130638A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris		Bupropion	PS	Glaxo Wellcome	ORAL
		Arrhythmia		Venlafaxine	SS		
UNKNOWN							

Date:10/30/00ISR Number: 3602865-XReport Type:Expedited (15-DaCompany Report #B0088745A  
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged RESPIRATORY Disability (INHALATION)		Chest Pain Electrocardiogram T Wave Inversion Myocardial Infarction		Zyban Salbutamol Aspirin Bendrofluazide Prednisolone Beclomethasone	PS C C C C	Glaxo Wellcome Glaxo Wellcome Glaxo Wellcome	ORAL ORAL ORAL
RESPIRATORY (INHALATION)				Ipratropium	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/30/00ISR Number: 3603787-0Report Type:Expedited (15-DaCompany Report #A0125519A

Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Disability		Arrhythmia	Professional				
		Atherosclerosis					
		Coronary Artery Disease					
		Fibrosis					
		Hepatic Steatosis					
		Ill-Defined Disorder					
		Injury					
		Oedema					
		Pain					
		Pancreatic Disorder					
		Pulmonary Congestion					
		Pulmonary Oedema					

Date:10/30/00ISR Number: 3603788-2Report Type:Expedited (15-DaCompany Report #A0123011A

Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Hallucination	Health	Bupropion			
		Hyperventilation	Professional	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Overdose					
		Paranoia					
				Oxycodone			
				Hydrochloride			
				(Formulation			
				Unknown) (Oxycodone			
				Hydrochloride)	SS		ORAL
				Amantadine	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SEE IMAGE		Agitation Disorientation	Consumer	Effexor Xr	PS	Wyeth Ayerst Laboratories	ORAL
ORAL		Hallucination Hyperhidrosis Lethargy		Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL
		Syncope Thinking Abnormal		Synthroid (Levothyroxien Sodium)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 150 MG/ PER Intervention to DAY/ ORAL Prevent Permanent Impairment/Damage		Oculogyration	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Fluoxetine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/30/00ISR Number: 3604400-9Report Type:Expedited (15-DaCompany Report #B0089984A  
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Angina Pectoris	Foreign	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Condition Aggravated					
Intervention to		Coronary Artery Occlusion		Atenolol +			
Prevent Permanent				Chlorthalidone	C		
Impairment/Damage				Doxazosin	C		

Date:10/30/00ISR Number: 3604401-0Report Type:Expedited (15-DaCompany Report #B0089929A  
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Eye Irritation	Foreign	Zyban	PS	Glaxo Wellcome Inc	
150 MG/ TWICE		Insomnia	Health				
PER DAY/	7 WK	Keratoconjunctivitis	Professional	Cimetidine	C		
		Sicca		Temazepam	C		
		Micturition Disorder					

Date:10/30/00ISR Number: 3604408-3Report Type:Expedited (15-DaCompany Report #B0089784A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Amnesia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Drug Hypersensitivity	Health				
Initial or Prolonged		Grand Mal Convulsion	Professional	Losartan Potassium	C		
PER DAY/ ORAL		Thrombocythaemia		Bromazepam	C		

Date:10/30/00ISR Number: 3602873-9Report Type:Expedited (15-DaCompany Report #B0090537A  
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	300MG PER DAY 5 WK	Fall		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Femur Fracture					

Date:10/31/00ISR Number: 3603414-2Report Type:Expedited (15-DaCompany Report #A0120776A  
Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	7 DAY	Bipolar Disorder		Lamictal	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	200MG TWICE	Condition Aggravated		Wellbutrin	SS	Glaxo Wellcome	
	PER DAY	Dermatitis					
	1MG AS	Headache		Klonopin	C		
REQUIRED		Muscle Twitching					
		Petit Mal Epilepsy		Prilosec	C		
		Pruritus		Dalmane	C		
		Sinusitis		Celexa	C		
		Stevens-Johnson Syndrome		Trazodone	C		
		Toxic Epidermal		Compazine	C		
		Necrolysis		Carafate	C		
		Tremor		Vitamins	C		
		Urinary Incontinence		Zyprexa	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/31/00ISR Number: 3603596-2Report Type:Direct  
Age:16 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS		
Other		Convulsion		Tetracycline	C		
BID				Ery Solution	C		
				Ritalin	C		

Date:10/31/00ISR Number: 3604492-7Report Type:Direct  
Age:41 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Akathisia		Zyban (Bupropion			
Other		Amnesia		Hcl) Sustained-			
		Anxiety		Released Tablets 150			
150 MG ORALLY		Confusional State		Mg	PS		ORAL
PER DAY		Disturbance In Attention					
		Epistaxis					
		Movement Disorder					
		Speech Disorder					
		Tardive Dyskinesia					
		Tinnitus					
		Vision Blurred					

Date:10/31/00ISR Number: 3605053-6Report Type:Expedited (15-DaCompany Report #B0090537A  
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Fall	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL	5 WK						
Initial or Prolonged		Feeling Abnormal	Consumer				
		Femur Fracture					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris	Foreign	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
150 MG /PER		Arrhythmia					
DAY/ORAL							
				Venlafaxine Hydrochloride (Formulation Unknown) (Venlafaxine	SS		
375 MG/PER							
DAY							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris	Foreign	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
ORAL		Arrhythmia					
				Venlafaxine Hydrochloride (Formulation Unknown) (Venlafaxine	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/31/00ISR Number: 3606776-5Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #AMANTADI2000-00258

Outcome	PT	Report Source	Product	Role	Manufacturer	Route	
Dose Duration Life-Threatening Hospitalization - BID PO Initial or Prolonged 150 MG DLY PO	Aggression Coma Confusional State Convulsion Drug Interaction Drug Level Below Therapeutic Ecchymosis Fall Fatigue Headache Heart Rate Increased Memory Impairment Movement Disorder Muscle Rigidity Nausea Oxygen Saturation Decreased Pallor Respiratory Rate Decreased Speech Disorder	Distributor	Symmetrel  Wellbutrin Sr Wellcome  Semisodium Valproate	  Glaxo SS C	PS  Glaxo Wellco	Endo Pharmaceuticals Inc  Glaxo Wellco	ORAL  ORAL

Date:10/31/00ISR Number: 3603416-6Report Type:Expedited (15-DaCompany Report #A0130226A  
Age:41 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100MG PER DAY 90 DAY Initial or Prolonged	Abdominal Tenderness Albuminuria Blood Creatine Phosphokinase Increased Chest Wall Pain Chromaturia Haematocrit Decreased Haemoglobinuria Hepatic Enzyme Increased		Wellbutrin  Methadone	PS  SS	Glaxo Wellcome  Glaxo Wellcome	ORAL

Hepatitis C  
Hypercalcaemia  
Hypokalaemia  
Leukopenia  
Rhabdomyolysis  
White Blood Cells Urine  
Positive

Date:10/31/00ISR Number: 3603417-8Report Type:Expedited (15-DaCompany Report #A0130639A  
Age:22 YR Gender:Female I/FU:I

Outcome PT  
Other Angioneurotic Oedema  
Arthralgia  
Joint Swelling  
Leukocytosis  
Myalgia  
Pyrexia  
Skin Discolouration  
Urinary Tract Infection

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Urticaria

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG TWICE PER DAY			Zyban	PS	Glaxo Wellcome	ORAL
			Oral Contraceptive	C		

Date:11/01/00ISR Number: 3604014-0Report Type:Expedited (15-DaCompany Report #A0070194A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	24 DAY	Activities Of Daily Living Impaired		Zyban	PS	Glaxo Wellcome	ORAL
1 DAY		Amnesia		Lidocaine	C		
1 DAY		Anger		General Anesthesia	C		
INTRA VENOUS	1 DAY	Anorexia		Surgery	C		
		Anxiety Disorder		D5 L.R	C		
INTRA VENOUS	1 DAY	Apathy		Levsin	C		
		Asthenia		Demerol	C		
1 DAY		Cognitive Disorder		Diprivan	C		
1 DAY		Confusional State		Midazolam	C		
1 DAY		Convulsion		Nitrous Oxide	C		
1 DAY		Depression		Droperidol	C		
1 DAY		Diarrhoea		Zofran	C	Glaxo Wellcome	
1 DAY		Disturbance In Attention		Desflurane	C		
1 DAY		Drug Ineffective		Succinylcholine	C	Glaxo Wellcome	
1 DAY		Early Morning Awakening		Fentanyl	C		
		Encephalopathy		Hydrocodone + Acetaminophen	C		

1	DAY	Fatigue	Marcaine	C	
		Fear	Tylenol #3	C	ORAL
		Feeling Abnormal	Imodium	C	
		Feeling Guilty	Motrin	C	
		Fibroadenoma Of Breast	Oxygen	C	

1 DAY

Headache  
Hypochondriasis  
Initial Insomnia  
Loss Of Libido  
Mood Swings  
Nausea  
Suicidal Ideation  
Thinking Abnormal  
Vomiting

Date:11/01/00ISR Number: 3604036-XReport Type:Expedited (15-DaCompany Report #B0090510A  
Age:45 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG PER DAY 5 DAY	Angina Unstable		Bupropion	PS	Glaxo Wellcome	ORAL
	100MG PER DAY			Diclofenac	C		ORAL
	250MG PER DAY 6 DAY			Amoxicillin	C		ORAL

Date:11/01/00ISR Number: 3604025-5Report Type:Expedited (15-DaCompany Report #B0090063A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	2TAB PER DAY 14 DAY	Cerebral Artery Embolism		Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/01/00ISR Number: 3604032-2Report Type:Expedited (15-DaCompany Report #B0090433A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Ischaemia		Zyban	PS	Glaxo Wellcome	ORAL
150MG AS		Sudden Death					
DIRECTED							

Date:11/01/00ISR Number: 3604035-8Report Type:Expedited (15-DaCompany Report #B0090507A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Dermatitis Exfoliative		Bupropion	PS	Glaxo Wellcome	ORAL
150MG TWICE		Vasculitis					
PER DAY				Dextropropoxyphene + Paracetamol	C		ORAL
				Flucloxacillin	C		ORAL
250MG FOUR							
TIMES PER DAY	8	DAY					
				Betamethasone Valerate + Salicylic Acid	C		
TOPICAL							

Date:11/02/00ISR Number: 3605180-3Report Type:Expedited (15-DaCompany Report #A0127926A  
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Ageusia	Health	Zyban	PS	Glaxo Wellcome	ORAL
150MG TWICE		Anxiety	Professional				
Initial or Prolonged		Condition Aggravated		Paxil	C		ORAL
PER DAY	8	Depression		Enteric Coated Asa	C		ORAL
325MG PER DAY		Dyspnoea		Unspecified			

	Feeling Abnormal	Immunosuppressants	C	
	Hypertension	Cyclosporin	C	
125MG TWICE				
	Panic Attack			
PER DAY				
	Tearfulness	Ramipril	C	ORAL
12.5MG PER				
DAY	Tobacco Abuse			
	Tremor	Pravachol	C	
40MG PER DAY				
		Losec	C	ORAL
20MG PER DAY				
		Oxazepam	C	ORAL
15MG AS				
REQUIRED				
		Oxazepam	C	ORAL
30MG AT NIGHT				
		Hytrin	C	ORAL
1MG PER DAY				
		Lasix	C	ORAL
60MG PER DAY				
		Atrovent	C	Glaxo Wellcome
RESPIRATORY				
(INHALATION)	2PUFF FOUR			
TIMES PER DAY				
		Prednisone	C	ORAL
5MG PER DAY				
		Septra	C	Glaxo Wellcome
1TAB PER DAY				
		Cozaar	C	ORAL
25MG PER DAY				

Date:11/02/00ISR Number: 3605987-2Report Type:Expedited (15-DaCompany Report #A0120776A  
Age:44 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Headache
Initial or Prolonged	Muscle Twitching
	Petit Mal Epilepsy
	Rash Pruritic
	Sinusitis
	Stevens-Johnson Syndrome

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Adverse Event	Report Source	Product	Role	Manufacturer	Route
UNKNOWN	200 MG /	Toxic Epidermal Necrolysis Tremor Urinary Incontinence	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	
TWICE PER DAY/ UNKNOWN				Lamictal Tablet (Lamotrigine)	SS		ORAL
100 MG / ORAL				Clonazepam	C		
				Omeprazole	C		
				Dalmane	C		
				Citalopram Hydrobromide	C		
				Trazadone	C		
				Prochlorperazine	C		
				Sucralfate	C		
				Multivitamin	C		
				Olanzapine	C		

Date:11/02/00ISR Number: 3605989-6Report Type:Expedited (15-DaCompany Report #A0130226A  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG / PER Initial or Prolonged DAY / ORAL		Abdominal Pain Upper	Health Professional	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
		Albuminuria					
		Blood Creatine Phosphokinase Increased Chest Wall Pain Chromaturia Haematocrit Decreased Haemoglobinuria Hepatic Enzyme Increased Hepatic Function Abnormal Hepatitis C Hypocalcaemia Hypokalaemia Leukopenia		Methadone Hydrochloride (Formulation Unknown) (Methadone Hydrochloride)	SS		

Rhabdomyolysis  
White Blood Cells Urine  
Positive

Date:11/02/00ISR Number: 3606471-2Report Type:Expedited (15-DaCompany Report #A0130639A  
Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema Arthralgia	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE		Joint Swelling					
PER DAY/ ORAL		Leukocytosis Myalgia Pruritus Pyrexia Skin Discolouration Urinary Tract Infection Urticaria		Oral Contraceptive	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/02/00 ISR Number: 3605188-8 Report Type:Expedited (15-DaCompany Report #B0089839A  
 Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 13 DAY Initial or Prolonged Other		C-Reactive Protein Increased Dermatitis Myopathy Polyarthrititis		Zyban	PS	Glaxo Wellcome	ORAL

Date:11/03/00 ISR Number: 3606861-8 Report Type:Expedited (15-DaCompany Report #A0070194A  
 Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150 MG / PER DAY / ORAL		Amnesia Anaesthetic Complication	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Anger Anxiety Anxiety Disorder Apathy Asthenia Brain Contusion Cognitive Disorder Confusional State Convulsion Crying Decreased Activity Decreased Appetite Depressed Mood Depression Disturbance In Attention Drug Ineffective Dysphoria Encephalopathy Fatigue Fear Feeling Abnormal Headache Hypochondriasis Hypoxia		General Anesthetic (Formulation Unknown) (General Anesthetic) Ibuprofen Loperamide Hydrochloride Dextrose + Lactated Ringers Tylenol No. 3 Hyoscyamine Sulphate Bupivacaine Hydrochloride Pethidine Hydrochloride Hydrocodone + Paracetamol Propofol Lignocaine Fentanyl Midazolam Suxamethonium Chloride Desflurane	SS C		

Insomnia  
Loss Of Libido  
Memory Impairment  
Mood Swings  
Nausea  
Suicidal Ideation  
Thinking Abnormal

Ondansetron  
Hydrochloride C  
Oxygen C  
Droperidol C  
Nitrous Oxide C

Date:11/03/00ISR Number: 3607169-7Report Type:Expedited (15-DaCompany Report #B0090507A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability		Dermatitis Exfoliative Vasculitis	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY / ORAL				Dextroprop. + Paracetamol Flucloxacillin Betamet.Val.+Salicyl	C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ic Acid C

Date:11/03/00ISR Number: 3607171-5Report Type:Expedited (15-DaCompany Report #B0090510A  
Age:45 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG / PER	Angina Unstable	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
DAY / ORAL				Diclofenac	C		
				Amoxicillin	C		

Date:11/03/00ISR Number: 3607172-7Report Type:Expedited (15-DaCompany Report #B0090433A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG / AS	Myocardial Ischaemia Sudden Death	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
DIRECTED /							
ORAL							

Date:11/03/00ISR Number: 3605721-6Report Type:Expedited (15-DaCompany Report #D0011443A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Difficulty In Walking Pain In Extremity		Zyban	PS	Glaxo Wellcome	ORAL

Date:11/06/00ISR Number: 3606520-1Report Type:Expedited (15-DaCompany Report #A0128801A  
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200MG TWICE		Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
PER DAY	YR	Drug Interaction					
60MG PER DAY		Tremor		Celexa	SS		
		Tunnel Vision		Lithium Trilafon	C C	Glaxo Wellcome	

Date:11/06/00ISR Number: 3607154-5Report Type:Expedited (15-DaCompany Report #B0090063A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL	14 DAY	Cerebral Artery Embolism	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:11/06/00ISR Number: 3607766-9Report Type:Expedited (15-DaCompany Report #B0089839A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL Required Intervention to Prevent Permanent Impairment/Damage		C-Reactive Protein Increased Dermatitis Myopathy Polyarthritis	Foreign Health Professional Company Representative	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/06/00ISR Number: 3607768-2Report Type:Expedited (15-DaCompany Report #A0127926A

Age:56 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG (TWICE PER DAY) ORAL	Anxiety Condition Aggravated  Depression  Dysgeusia Dyspnoea Feeling Abnormal Hypertension Panic Attack Tearfulness Tremor	Foreign Health  Professional	Bupropion Hydrochloride   Immunosuppressive Paroxetine Hydrochloride Prednisone Cyclosporin Ipratropium Bromide Ramipril Frusemide Co-Trimoxazole Aspirin Pravastatin Sodium Terazosin Hydrochloride Omeprazole Oxazepam Losartan Potassium	PS    C C C C C C C C C C C C C C C C	Glaxo Wellcome Inc	ORAL

Date:11/06/00ISR Number: 3607833-XReport Type:Expedited (15-DaCompany Report #D0011443A

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG, ORAL	Difficulty In Walking  Pain In Extremity	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:11/06/00ISR Number: 3606525-0Report Type:Expedited (15-DaCompany Report #B0086032A

Age:59 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Hospitalization - Arrhythmia Zyban PS Glaxo Wellcome ORAL  
 1TAB PER DAY 2 DAY  
 Initial or Prolonged Atrial Fibrillation  
 Cardiomegaly  
 Dyspnoea  
 Echocardiogram Abnormal  
 Hypertension  
 Pulmonary Oedema

Date:11/06/00ISR Number: 3606533-XReport Type:Expedited (15-DaCompany Report #B0090441A  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Multiple Sclerosis		Bupropion			
		Nervous System Disorder		Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG TWICE							
PER DAY							
				Loratadine	C		ORAL
10MG PER DAY							
				Flucloxacillin	C		ORAL
250MG FOUR							
TIMES PER DAY 6	DAY						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/00ISR Number: 3607293-9Report Type:Expedited (15-DaCompany Report #A0066944A  
 Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG TWICE Initial or Prolonged PER DAY		Agitation		Wellbutrin	PS	Glaxo Wellcome	ORAL
Disability Other		Chest Discomfort					
		Chest Pain		Wellbutrin	SS	Glaxo Wellcome	ORAL
		Coronary Artery Atherosclerosis		Klonopin	C		
		Coronary Artery Occlusion		Aspirin	C		
				Zoloft	C		
2 YR		Dermatitis		Depakote	C		
		Disorientation					
		Drug Ineffective					
		Dyspnoea					
		Fatigue					
		Headache					
		Hyperhidrosis					
		Increased Appetite					
		Myalgia					
		Pharyngolaryngeal Pain					
		Pruritus					
		Weight Increased					

Date:11/07/00ISR Number: 3608290-XReport Type:Direct Company Report #  
 Age:16 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other BID		Convulsion		Wellbutrin	PS		
				Tetracycline	C		
				Ery Solution	C		
				Ritalin	C		

Date:11/07/00ISR Number: 3608310-2Report Type:Expedited (15-DaCompany Report #A0128801A  
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other 200 MG /	Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
TWICE PER	Drug Interaction	Professional				
DAY/ ORAL	1 YR	Tremor				
		Tunnel Vision	Citalopram Hydrobromide (Formulation Unknown) (Citalopram Hydrobromide)	SS		
UNKNOWN	60 MG / PER					
DAY/ UNKNOWN			Lithium Salt Perphenazine	C C		

Date:11/07/00ISR Number: 3608445-4Report Type:Expedited (15-DaCompany Report #B0086032A  
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1 TABLET/ PER		Arrhythmia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged DAY/ ORAL		Atrial Fibrillation	Health				
		Cardiomegaly	Professional				
		Dyspnoea					
		Echocardiogram Abnormal					
		Hypertension					
		Pulmonary Oedema					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/00ISR Number: 3608446-6Report Type:Expedited (15-DaCompany Report #B0090441A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Multiple Sclerosis	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
Intervention to							
PER DAY/ ORAL							
Prevent Permanent				Loratadine	C		
Impairment/Damage				Flucloxacillin	C		

Date:11/08/00ISR Number: 3608263-7Report Type:Expedited (15-DaCompany Report #A0130681A  
Age:21 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Drug Abuser		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG PER DAY 2	MON						
Initial or Prolonged		Fall		Alcohol	C		
Other		Grand Mal Convulsion		No Concurrent			
		Humerus Fracture		Medications	C		
		Intentional Misuse		Marijuana	C		
		Stress					
		Suicide Attempt					

Date:11/08/00ISR Number: 3608273-XReport Type:Expedited (15-DaCompany Report #B0090959A  
Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous		Zyban	PS	Glaxo Wellcome	

Date:11/08/00ISR Number: 3608274-1Report Type:Expedited (15-DaCompany Report #B0091014A  
Age:63 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Extrasystoles		Bupropion			
		Palpitations		Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG TWICE							

PER DAY

Thyroxine C Glaxo Wellcome ORAL

75UG PER DAY

Date:11/09/00ISR Number: 3608768-9Report Type:Expedited (15-DaCompany Report #B0091004A

Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG PER DAY			Cerebrovascular Accident Condition Aggravated	Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
75MG PER DAY			Hemiparesis	Aspirin	C		ORAL
RESPIRATORY (INHALATION)	100UG PER DAY	Attack	Transient Ischaemic	Salbutamol	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)	20UG PER DAY			Ipratropium	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)	25UG PER DAY			Salmeterol	C	Glaxo Wellcome	

Date:11/09/00ISR Number: 3608770-7Report Type:Expedited (15-DaCompany Report #B0091092A

Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG TWICE			Vision Blurred Visual Field Defect	Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
PER DAY	29 DAY						

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

500MG PER DAY		Ascorbic Acid	C	ORAL
1CAP PER DAY	YR	Multivitamins	C	ORAL

Date:11/09/00ISR Number: 3609604-7Report Type:Direct Company Report #  
 Age:12 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 1 QD ORAL	Grand Mal Convulsion		Concerta 18mg	PS		ORAL
2 BID ORAL	Insomnia		Wellbutrin Sr 100mg	SS		ORAL
	Medication Error					

Date:11/09/00ISR Number: 3610188-8Report Type:Expedited (15-DaCompany Report #A0066944A  
 Age:68 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL	Agitation	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged Disability Required 100 MG / Intervention to TWICE PER DAY Prevent Permanent / ORAL Impairment/Damage	Chest Discomfort Chest Pain Coronary Artery Atherosclerosis Coronary Artery Occlusion	Professional	Wellbutrin Tablet (Bupropion Hydrochloride)	SS		ORAL
	Dermatitis		Clonazepam	C		
	Disorientation		Aspirin	C		
	Drug Ineffective		Sertraline			
	Dyspnoea		Hydrochloride	C		
	Fatigue		Semisodium Valproate	C		
	Headache					
	Hyperhidrosis					
	Increased Appetite					
	Pharyngolaryngeal Pain					
	Pruritus					
	Weight Increased					

Date:11/09/00ISR Number: 3608758-6Report Type:Expedited (15-DaCompany Report #A0131125A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Congenital Anomaly			Caesarean Section	Bupropion	PS	Glaxo Wellcome	ORAL
150MG TWICE							
PER DAY	5	WK	Complications Of Maternal				
			Exposure To Therapeutic				
			Drugs				
			Congenital Aortic Valve				
			Incompetence				
			Congenital Aortic Valve				
			Stenosis				

Date:11/09/00ISR Number: 3608767-7Report Type:Expedited (15-DaCompany Report #B0090989A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Migraine With Aura	Bupropion			
150MG PER DAY	41	DAY	Visual Disturbance	Hydrochloride	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/09/00ISR Number: 3608769-0Report Type:Expedited (15-DaCompany Report #B0091015A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG TWICE PER DAY	3 DAY	Chest Pain Hypokalaemia Tachycardia		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG PER DAY				Aspirin	C		ORAL
10MG PER DAY				Enalapril	C		ORAL
7.5MG PER DAY				Zopiclone	C		ORAL
100MG PER DAY				Calcium Carbonate + Disodium Etidronate	C		ORAL
135MG THREE TIMES PER DAY				Atenolol	C		ORAL
150MG PER DAY				Mebeverine	C		ORAL
				Ranitidine	C	Glaxo Wellcome	ORAL

Date:11/13/00ISR Number: 3609968-4Report Type:Expedited (15-DaCompany Report #B0091033A  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG TWICE PER DAY	15 DAY	Brain Stem Infarction Cerebellar Infarction Drug Interaction		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
50MG PER DAY		Headache Hypertension		Catecholamine Reuptake Inhibitor	SS		
				Atenolol	SS		ORAL

Date:11/13/00ISR Number: 3609992-1Report Type:Expedited (15-DaCompany Report #B0086470A  
Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Atrial Fibrillation		Zyban	PS	Glaxo Wellcome	ORAL
150MG TWICE							
Disability		Hyperhidrosis					
PER DAY	11 DAY						
Other		Malaise		Combivent	C		
RESPIRATORY							
(INHALATION)	2PUFF FOUR	Night Sweats					
TIMES PER DAY							
RESPIRATORY				Beclomethasone	C	Glaxo Wellcome	
(INHALATION)	2PUFF TWICE						
PER DAY							

Date:11/13/00ISR Number: 3609993-3Report Type:Expedited (15-DaCompany Report #B0086888A  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Bupropion	PS	Glaxo Wellcome	ORAL
150MG SEE							
TEXT		Feeling Abnormal					
		Personality Change					

Date:11/13/00ISR Number: 3609995-7Report Type:Expedited (15-DaCompany Report #B0089436A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Zyban	PS	Glaxo Wellcome	ORAL
150MG TWICE							
PER DAY		Feeling Abnormal					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/13/00ISR Number: 3610000-7Report Type:Expedited (15-DaCompany Report #B0091168A  
Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	1 DAY	Acidosis		Bupropion	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged		Hyperhidrosis		Paracetamol	SS		
Other		Hypotension		Alcohol	SS		
		Intentional Misuse					
		Pyrexia					
		Tachycardia					

Date:11/13/00ISR Number: 3610826-XReport Type:Expedited (15-DaCompany Report #2000UW04157  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Amnesia	Consumer	Tenormin	PS	Astrazeneca Pharmaceuticals Lp	
150 MG BID PO		Hypotension		Zyban	SS		ORAL
		Pneumonia		Nifedipine	SS		
		Sedation		Gabapentin	SS		
		Speech Disorder					
		Toxicologic Test Abnormal					

Date:11/13/00ISR Number: 3609948-9Report Type:Expedited (15-DaCompany Report #A0128539A  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	3 WK	Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
		Blood Creatine Phosphokinase Increased		No Concurrent Medications	C		
		Dermatomyositis					
		Face Oedema					
		Myalgia					
		Oedema					

Date:11/14/00ISR Number: 3611401-3Report Type:Expedited (15-DaCompany Report #A0128539A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 150 MG / ORAL	3 WK	Arthralgia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
Intervention to Prevent Permanent Impairment/Damage		Blood Creatine Phosphokinase Increased Dermatomyositis Face Oedema Myalgia	Professional				

Date:11/14/00ISR Number: 3611431-1Report Type:Expedited (15-DaCompany Report #A0131125A  
Age: Gender:Male I/FU:I

Outcome	PT
Congenital Anomaly	Aortic Valve Disease Caesarean Section Complications Of Maternal Exposure To Therapeutic Drugs Congenital Aortic Valve Incompetence Heart Disease Congenital

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Twin Pregnancy

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE		Study	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ORAL		Health				
		Professional				

Date:11/14/00ISR Number: 3611630-9Report Type:Expedited (15-DaCompany Report #B0091014A  
Age:63 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Extrasystoles	Foreign	Zyban	PS	Pharmacia And Upjohn Co	ORAL
Intervention to		Palpitations	Other				
150MG (TWICE				Thyroxine Sodium	C		
Prevent Permanent							
PER DAY) ORAL							
Impairment/Damage							

Date:11/14/00ISR Number: 3611631-0Report Type:Expedited (15-DaCompany Report #B0090959A  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
			Health				
			Professional				
			Other				

Date:11/14/00ISR Number: 3611644-9Report Type:Expedited (15-DaCompany Report #B0086470A  
Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Atrial Fibrillation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/			Health				
Disability		Hyperhidrosis					
TWICE PER							

Required	Nausea	Professional				
DAY/ ORAL						
Intervention to	Night Sweats			Combivent	C	
Prevent Permanent				Beclomethasone		
Impairment/Damage				Dipropion	C	

Date:11/14/00ISR Number: 3611826-6Report Type:Expedited (15-DaCompany Report #B0089436A  
 Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL			Health				
			Professional				
			Company				
			Representative				

Date:11/14/00ISR Number: 3611827-8Report Type:Expedited (15-DaCompany Report #B0086888A  
 Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG SEE							
TEXT ORAL		Feeling Abnormal	Health				
			Professional				



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/14/00ISR Number: 3611921-1Report Type:Expedited (15-DaCompany Report #B0091168A  
Age:18 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 150 MG /ORAL Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage	Acidosis Blood Ph Decreased Coma Convulsion Hyperhidrosis Hypotension Irritability Medication Error Overdose Pyrexia Tachycardia	Foreign	Bupropion Hydrochloride  Paracetamol (Acetaminophen) Ethanol (Alcohol)	PS  SS SS	Glaxo Wellcome Inc	ORAL

Date:11/14/00ISR Number: 3611940-5Report Type:Expedited (15-DaCompany Report #B0091033A  
Age:41 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death  150 MG/TWICE  PER DAY/ ORAL  ORAL	Brain Stem Infarction Cerebellar Infarction  Drug Interaction  Headache Hypertension	Foreign	Bupropion Hydrochloride  Atenolol (Formulation Unknown) (Atenolol)  Catecholamine Reuptake Inhibitor (Formulation	PS  SS SS	Glaxo Wellcome Inc	ORAL  ORAL

Date:11/14/00ISR Number: 3611946-6Report Type:Expedited (15-DaCompany Report #B0091004A  
Age:50 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization -	Cerebrovascular Accident	Foreign	Bupropion			

Initial or Prolonged 150 MG/PER	Hemiparesis	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
DAY/ ORAL	Transient Ischaemic Attack	Aspirin Salbutamol Sulphate Ipratropium Bromide Salmeterol Xinafoate	C C C C		

Date:11/14/00ISR Number: 3611948-XReport Type:Expedited (15-DaCompany Report #B0091015A  
 Age:50 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL	Chest Pain Hypokalaemia Tachycardia	Foreign	Zyban  Aspirin Enalapril Zopiclone Ca Carbonate+Na Etidronat Atenolol Mebeverine Ranitidine Hydrochloride	PS  C C C C C C C	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/14/00ISR Number: 3611949-1Report Type:Expedited (15-DaCompany Report #B0091092A  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG/ TWICE	Vision Blurred	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL		Visual Field Defect		Ascorbic Acid	C		
				Multivitamin	C		

Date:11/14/00ISR Number: 3611950-8Report Type:Expedited (15-DaCompany Report #B0090989A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required	ORAL	Migraine With Aura	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Intervention to Prevent Permanent Impairment/Damage		Visual Disturbance					

Date:11/14/00ISR Number: 3612383-0Report Type:Expedited (15-DaCompany Report #A0125460A  
Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	ORAL	Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - Initial or Prolonged		Ecchymosis	Professional	Clozapine	C		
		Excoriation	Company	Lisinopril	C		
		Fall	Representative	Docusate Sodium	C		
		Gastrointestinal Disorder		Amlodipine	C		
		Hepatic Congestion					
		Hepatic Failure					
		Hepatic Steatosis					
		Intestinal Ischaemia					
		Laceration					
		Liver Disorder					
		Mucosal Haemorrhage					
		Pancreatic Disorder					
		Pulmonary Congestion					

Pulmonary Oedema  
Rib Fracture  
Splenic Injury  
Wrist Fracture

Date:11/14/00ISR Number: 3612564-6Report Type:Expedited (15-DaCompany Report #A130601A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL Disability		Arthralgia Blindness Unilateral Brain Damage Brain Hypoxia Cardiac Arrest Cerebrovascular Accident Coma Convulsion Dyspnoea Haemorrhage Headache Hemiplegia Rash Generalised	Literature Health Professional Company Representative	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/14/00ISR Number: 3612648-2Report Type:Expedited (15-DaCompany Report #A0130681A  
Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ PER Initial or Prolonged DAY/ ORAL		Fall	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Grand Mal Convulsion	Professional				
		Humerus Fracture	Company	Ethanol	C		
		Intentional Misuse	Representative	Cannabis	C		
		Suicide Attempt					

Date:11/15/00ISR Number: 3611257-9Report Type:Expedited (15-DaCompany Report #B0091314A  
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG PER DAY 1 DAY		Left Ventricular Failure		Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY (INHALATION)				Beclomethasone	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)				Salbutamol	C	Glaxo Wellcome	

Date:11/15/00ISR Number: 3611261-0Report Type:Expedited (15-DaCompany Report #A0070194A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 1 DAY		Adjustment Disorder		General Anesthesia	PS		
150MG PER DAY 24 DAY		Amnesia		Zyban	SS	Glaxo Wellcome	ORAL
		Anger		Motrin	C		
		Anxiety		Surgery	C		
		Anxiety Disorder		Imodium	C		
		Asthenia		D5 L.R	C		
INTRAVENOUS	1 DAY						

		Cognitive Disorder	Tylenol #3	C	ORAL
		Confusional State	Levsin	C	
		Convulsion	Marcaine	C	
1	DAY				
		Crying	Demerol	C	
		INTRA VENOUS			
		1 DAY			
		Decreased Activity	Hydrocodone +		
		Depressed Mood	Acetaminophen	C	
		Depression	Diprivan	C	
1	DAY				
		Diarrhoea	Lidocaine	C	
1	DAY				
		Disturbance In Attention	Fentanyl	C	
1	DAY				
		Drug Ineffective	Midazolam	C	
1	DAY				
		Early Morning Awakening	Succinylcholine	C	Glaxo Wellcome
1	DAY				
		Encephalopathy	Desflurane	C	
1	DAY				
		Fatigue	Zofran	C	Glaxo Wellcome
1	DAY				
		Fear	Oxygen	C	
1	DAY				
		Feeling Abnormal	Droperidol	C	
1	DAY				
		Headache	Nitrous Oxide	C	
1	DAY				
		Hypoxia			
		Injury			
		Insomnia			
		Mental Impairment			
		Mood Swings			
		Nausea			
		Nervous System Disorder			
		Thinking Abnormal			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/15/00ISR Number: 3611267-1Report Type:Expedited (15-DaCompany Report #B0089784A

Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG TWICE		Grand Mal Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged PER DAY	4 DAY	Hypersensitivity					
1TAB PER DAY		Loss Of Consciousness		Losartan	C		ORAL
1TAB PER DAY		Thrombocythaemia		Bromazepam	C		ORAL

Date:11/15/00ISR Number: 3611766-2Report Type:Direct

Company Report #

Age:12 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 3X DAILY,		Bipolar I Disorder Condition Aggravated		Wellbutrin 75mg Burroughs Wellcome	PS	Burroughs Wellcome	
Initial or Prolonged 75MG EACH Required Intervention to Prevent Permanent Impairment/Damage	1 YR	Suicide Attempt					

Date:11/15/00ISR Number: 3611241-5Report Type:Expedited (15-DaCompany Report #A0131385A

Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG TWICE		Confusional State		Wellbutrin	PS	Glaxo Wellcome	ORAL
Hospitalization - PER DAY		Dermatitis					
Initial or Prolonged Other		Liver Function Test Abnormal Multi-Organ Failure Pyrexia Stevens-Johnson Syndrome					

Toxic Epidermal  
Necrolysis

Date:11/15/00ISR Number: 3611247-6Report Type:Expedited (15-DaCompany Report #B0089979A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2 WK Initial or Prolonged		Erythema Multiforme Hyperkeratosis Rash Pustular Skin Disorder		Zyban Coproxamol	PS C	Glaxo Wellcome	ORAL

Date:11/15/00ISR Number: 3611248-8Report Type:Expedited (15-DaCompany Report #B0090811A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 5 DAY		Crohn'S Disease		Zyban Mesalazine	PS C	Glaxo Wellcome	ORAL

Date:11/16/00ISR Number: 3611880-1Report Type:Expedited (15-DaCompany Report #A0131900A  
Age:80 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Blood Creatine Phosphokinase Increased

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Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG TWICE PER DAY	65 DAY	Blood Creatine Phosphokinase Mb Increased Chest Pain	Bupropion	PS	Glaxo Wellcome	ORAL
1TAB TWICE PER DAY	65 DAY	Myocardial Infarction	Placebo	SS		ORAL

Date:11/16/00ISR Number: 3611883-7Report Type:Expedited (15-DaCompany Report #B0085539A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG TWICE PER DAY	15 DAY	Motion Sickness Vertigo		Zyban	PS	Glaxo Wellcome	ORAL
SUBCUTANEOUS RESPIRATORY (INHALATION)		Visual Acuity Reduced		Insulin	C		
				Flixotide	C	Glaxo Wellcome	

Date:11/16/00ISR Number: 3611886-2Report Type:Expedited (15-DaCompany Report #B0089069A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG PER DAY Hospitalization - 5MG THREE Initial or Prolonged TIMES PER DAY	11 DAY	Brain Stem Infarction		Bupropion	PS	Glaxo Wellcome	ORAL
		Circulatory Collapse		Diclofenac	C		ORAL
300MG PER DAY	185 DAY			Allopurinol	C	Glaxo Wellcome	ORAL

Date:11/16/00ISR Number: 3611887-4Report Type:Expedited (15-DaCompany Report #B0089166A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG TWICE		Aggression		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged PER DAY 2 MON Other		Agitation Emotional Disorder Emotional Distress Fatigue Hallucination Nightmare Paranoia Sleep Walking					

Date:11/16/00ISR Number: 3611890-4Report Type:Expedited (15-DaCompany Report #B0089994A  
Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG TWICE		Cerebellar Ataxia		Zyban	PS	Glaxo Wellcome	ORAL
PER DAY 11 DAY		Cerebral Ischaemia					
2.5MG PER DAY		Viral Infection		Atenolol + Chlorthalidone Lisinopril	C C		ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/16/00ISR Number: 3611894-1Report Type:Expedited (15-DaCompany Report #B0090960A  
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG TWICE		Glaucoma		Zyban	PS	Glaxo Wellcome	ORAL
PER DAY	10	DAY					

Date:11/16/00ISR Number: 3611899-0Report Type:Expedited (15-DaCompany Report #B0091448A  
Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG SINGLE		Bronchospasm Cardiovascular Disorder		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
DOSE	1	DAY					
UNKNOWN		Decreased Activity Dyspnoea		Corticosteroids	C		
UNKNOWN		Muscular Weakness		Antibiotics	C		
UNKNOWN				Unspecified Medication	C		

Date:11/16/00ISR Number: 3611901-6Report Type:Expedited (15-DaCompany Report #B0091583A  
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 150MG PER DAY		Sudden Death		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL

Date:11/17/00ISR Number: 3613270-4Report Type:Direct Company Report #  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG PO		Chills		Zyban (Bupropion)	PS		ORAL
Initial or Prolonged		Malaise Oral Pain Pyrexia Rash Erythematous Serum Sickness Vomiting					

Date:11/17/00ISR Number: 3613388-6Report Type:Expedited (15-DaCompany Report #A0131385A  
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG / Hospitalization - TWICE PER DAY		Confusional State	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged / ORAL		Dermatitis	Professional				
Required Intervention to Prevent Permanent Impairment/Damage		Liver Function Test  Abnormal Multi-Organ Failure Pyrexia Rash Generalised Stevens-Johnson Syndrome					

Date:11/17/00ISR Number: 3613411-9Report Type:Expedited (15-DaCompany Report #B0091314A  
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150 MG / PER  DAY / ORAL		Left Ventricular Failure	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Beclomethasone  
 Dipropion C  
 Salbutamol Sulphate C

Date:11/17/00ISR Number: 3613414-4Report Type:Expedited (15-DaCompany Report #B0089979A  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNK / UNK / Initial or Prolonged ORAL	2 WK	Erythema Multiforme	Foreign Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Hyperkeratosis	Professional Company Representative	Co-Proxamol	C		
		Psoriasis					
		Rash Pustular					

Date:11/17/00ISR Number: 3613415-6Report Type:Expedited (15-DaCompany Report #B0090811A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required UNK / UNK / Intervention to ORAL	5 DAY	Crohn'S Disease	Foreign Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
Prevent Permanent Impairment/Damage			Professional	Mesalazine	C		

Date:11/17/00ISR Number: 3613533-2Report Type:Expedited (15-DaCompany Report #1458727A  
 Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Other Required Intervention to Prevent Permanent		Coma Convulsion Hyperhidrosis Hypotension Irritability Metabolic Acidosis Nervousness	Foreign Other	Tylenol	PS	Mcneil Consumer Products Co Div Mcneilab Inc	
				Zyban (R) Prolonged Release Tablets 150 Mg	SS		

Impairment/Damage Oral Intake Reduced  
Overdose  
Pyrexia  
Tachycardia

Date:11/17/00ISR Number: 3612413-6Report Type:Expedited (15-DaCompany Report #A0131854A  
Age:70 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG PER DAY		Cerebrovascular Accident	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Fall Intracranial Aneurysm					

Date:11/17/00ISR Number: 3612414-8Report Type:Expedited (15-DaCompany Report #B0091238A  
Age:57 YR Gender:Male I/FU:F

Outcome	PT
Life-Threatening	Cardiac Failure Congestive Cardiomegaly Chest X-Ray Abnormal Dyspnoea

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Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Face Oedema Fatigue Fluid Retention				
150MG TWICE		Hyponatraemia Oedema Peripheral	Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
PER DAY	40 DAY	Urine Osmolarity Decreased	Lisinopril	C		ORAL
20MG PER DAY			Atorvastatin	C		ORAL
10MG PER DAY	62 DAY		Atorvastatin	C		ORAL
10MG PER DAY	62 DAY		Atorvastatin	C		ORAL
90MG PER DAY			Aspirin	C		ORAL
150MG PER DAY						

Date:11/17/00ISR Number: 3612415-XReport Type:Expedited (15-DaCompany Report #B0091443A  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged	10MG PER DAY	UNKNOWN	Asthma	Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
				Cetirizine	C	Glaxo Wellcome	ORAL
				Flixonase	C	Glaxo Wellcome	
				Beclomethasone	C	Glaxo Wellcome	
				Salbutamol	C	Glaxo Wellcome	

Date:11/17/00ISR Number: 3612416-1Report Type:Expedited (15-DaCompany Report #B0091452A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		UNKNOWN	Asthma	Bupropion			

150MG PER DAY	9	DAY	Pruritus	Hydrochloride	PS	Glaxo Wellcome	ORAL
			Rash Macular				
			Upper Respiratory Tract				
			Infection				

Date:11/17/00ISR Number: 3612417-3Report Type:Expedited (15-DaCompany Report #B0091581A  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged				Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG TWICE							
PER DAY	19	DAY	Amplitude Decreased				
			Face Oedema	Gaviscon	C		
			Oesophageal Disorder				
			Urticaria				

Date:11/20/00ISR Number: 3613233-9Report Type:Expedited (15-DaCompany Report #A0127813A  
 Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other				Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG PER DAY			Blood Potassium Decreased				
			Epilepsy	K-Dur	C		
			Fall	Vitamins	C		
			Fatigue	Alcohol	C		
			Headache				
			Muscle Rigidity				
			Myalgia				
			Pallor				



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Freedom Of Information (FOI) Report

Date:11/20/00ISR Number: 3613236-4Report Type:Expedited (15-DaCompany Report #A0129577A  
 Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia		Zyban	PS	Glaxo Wellcome	ORAL
150MG TWICE							
PER DAY		Erectile Dysfunction					
		Fatigue		Cardiac Medication	SS		
25MG TWICE		Hypoaesthesia		Metoprolol	C		ORAL
PER DAY		Paraesthesia					
		Sexual Dysfunction		Ramipril	C		ORAL
5MG PER DAY							
		Triple Vessel Bypass		Metformin	C		ORAL
1000MG TWICE							
PER DAY		Graft					
				Glyburide	C		ORAL
10MG TWICE							
PER DAY							
				Losec	C		ORAL
20MG PER DAY							
				Gemfibrozil	C		ORAL
600MG TWICE							
PER DAY							

Date:11/20/00ISR Number: 3614432-2Report Type:Expedited (15-DaCompany Report #B0085539A  
 Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Motion Sickness	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL		Vertigo	Health				
		Visual Acuity Reduced	Professional	Insulin	C		
			Company	Fluticasone			
			Representative	Propionate	C		

Date:11/20/00ISR Number: 3614435-8Report Type:Expedited (15-DaCompany Report #B0089069A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Brain Stem Infarction	Foreign	Bupropion			
Hospitalization - ORAL		Circulatory Collapse		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged				Diclofenac	C		
				Allopurinol	C		

Date:11/20/00ISR Number: 3614436-XReport Type:Expedited (15-DaCompany Report #B0091448A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Bronchospasm	Foreign	Bupropion			
Intervention to 150 MG/ Prevent Permanent SINGLE DOSE/ Impairment/Damage ORAL		Cardiovascular Disorder	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Decreased Activity	Professional				
		Dyspnoea					
		Muscular Weakness		Corticosteroid	C		
				Antibiotics	C		
				Unknown	C		

Date:11/20/00ISR Number: 3614437-1Report Type:Expedited (15-DaCompany Report #B0090960A  
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Glaucoma	Foreign	Bupropion			
150 MG/ TWICE			Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL			Professional				

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Freedom Of Information (FOI) Report

Date:11/20/00ISR Number: 3614439-5Report Type:Expedited (15-DaCompany Report #B0089994A  
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent PER DAY/ ORAL Impairment/Damage		Cerebellar Ataxia Cerebral Ischaemia	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Atenolol + Chlorthalidone Lisinopril	C C		

Date:11/20/00ISR Number: 3614444-9Report Type:Expedited (15-DaCompany Report #B0091583A  
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death  150 MG / PER DAY / ORAL		Sudden Death	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:11/20/00ISR Number: 3614490-5Report Type:Expedited (15-DaCompany Report #A0070194A  
Age:39 YR Gender:Female I/FU:F

Outcome	PT
Disability	Amnesia Anger Anhedonia Anxiety Anxiety Disorder Asthenia Brain Contusion Breast Hyperplasia Breast Neoplasm Cognitive Disorder Condition Aggravated Confusional State Convulsion Decreased Appetite

Depressed Mood  
Depression  
Diarrhoea  
Disturbance In Attention  
Drug Ineffective  
Dysphoria  
Early Morning Awakening  
Encephalopathy  
Fatigue  
Feeling Abnormal  
Fibroadenoma Of Breast  
Fibrocystic Breast  
Disease  
Headache  
Hodgkin'S Disease Nodular  
Sclerosis Stage  
Unspecified  
Hypochondriasis  
Hypoxia  
Initial Insomnia  
Loss Of Libido  
Memory Impairment  
Mood Swings

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Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Professional				
			General Anesthetic (Formulation Unknown) (General Anesthetic)	SS		
			Ibuprofen	C		
			Surgery	C		
			Loperamide			
			Hydrochloride	C		
			Dextrose + Lactated			
			Ringers	C		
			Tylenol No. 3	C		
			Hyoscyamine Sulphate	C		
			Bupivacaine			
			Hydrochloride	C		
			Pethidine			
			Hydrochloride	C		
			Hydrocodone +			
			Paracetamol	C		
			Propofol	C		
			Lignocaine	C		
			Fentanyl	C		
			Midazolam	C		
			Suxmethonium			
			Chloride	C		
			Desflurane	C		
			Ondansetron			
			Hydrochloride	C		
			Oxygen	C		
			Droperidol	C		
			Nitrous Oxide	C		

Date:11/20/00ISR Number: 3614600-XReport Type:Expedited (15-DaCompany Report #B0091238A  
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Life-Threatening	Cardiac Failure	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE						
	Congestive	Health				
PER DAY/ ORAL						
	Cardiomegaly	Professional	Aspirin	C		
	Chest X-Ray Abnormal		Lisinopril	C		
	Dyspnoea		Atorvastatin Calcium	C		
	Face Oedema					
	Fatigue					
	Fluid Retention					
	Hyponatraemia					
	Oedema Peripheral					
	Urine Osmolarity					
	Decreased					

Date:11/20/00ISR Number: 3614602-3Report Type:Expedited (15-DaCompany Report #B0091452A  
Age:35 YR Gender:Female I/FU:I

Outcome  
Required  
Intervention to

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Prevent Permanent  
Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/	ORAL	Asthma	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Pruritus Rash Macular Upper Respiratory Tract Infection					

Date:11/20/00ISR Number: 3614604-7Report Type:Expedited (15-DaCompany Report #B0091443A  
Age:53 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged	150 MG/	ORAL	Asthma	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
					Cetirizine Hydrochloride	C		
					Fluticasone Propionate	C		
					Beclomethasone Dipropion.	C		
					Salbutamol Sulphate	C		

Date:11/20/00ISR Number: 3614606-0Report Type:Expedited (15-DaCompany Report #B0091581A  
Age:40 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER DAY/	150 MG/	TWICE ORAL	Chest Pain	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Dermatitis Exfoliative					
			Electrocardiogram T Wave Amplitude Decreased Face Oedema Oesophageal Disorder Urticaria		Gaviscon	C		

Date:11/20/00ISR Number: 3614634-5Report Type:Expedited (15-DaCompany Report #B0089166A  
Age:36 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG (TWICE Required PER DAY) ORAL Intervention to Prevent Permanent Impairment/Damage	Aggression Emotional Disorder Emotional Distress Fatigue Hallucination Nightmare Paranoia Sleep Disorder	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:11/20/00ISR Number: 3614655-2Report Type:Expedited (15-DaCompany Report #B0089784A  
Age:52 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ TWICE Initial or Prolonged DAILY/ ORAL	Grand Mal Convulsion Hypersensitivity Loss Of Consciousness Thrombocythaemia	Foreign Health Professional	Zyban Losartan Potassium Bromazepam	PS C C	Glaxo Wellcome Inc	ORAL



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Freedom Of Information (FOI) Report

Date:11/20/00ISR Number: 3614849-6Report Type:Expedited (15-DaCompany Report #A0131854A  
Age:70 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / PER Initial or Prolonged DAY / ORAL	Cerebrovascular Accident	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
	Fall	Consumer				
	Intracranial Aneurysm					

Date:11/20/00ISR Number: 3615282-3Report Type:Expedited (15-DaCompany Report #A0131900A  
Age:80 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL	Cardiac Enzymes Increased	Study	Zyban	PS	Glaxo Wellcome Inc	ORAL
	Chest Pain	Health				
1 TABLET/TWICE PER DAY/ORAL	Myocardial Infarction	Professional	Placebo	SS		ORAL

Date:11/20/00ISR Number: 3618538-3Report Type:Periodic Company Report #HQ9690310AUG2000  
Age:35 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 75 MG 1 X PER 1 DAY, ORAL ORAL	Abortion Spontaneous Pregnancy	Health Professional	Effexor Xr	PS	Wyeth Ayerst Laboratories	ORAL
			Tegretol (Carbamazepine)	SS		ORAL
			Wellbutrin (Amfebutamone)			

ORAL				Hydrochloride)	SS		ORAL
				Zyrtec (Cetirizine Hydrochloride)	SS		ORAL

Date:11/20/00ISR Number: 3613246-7Report Type:Expedited (15-DaCompany Report #B0091703A  
Age:28 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hyperventilation Pyrexia		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL

Date:11/20/00ISR Number: 3613248-0Report Type:Expedited (15-DaCompany Report #D0010443A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1TAB PER DAY Initial or Prolonged 1TAB PER DAY 1TAB TWICE PER DAY		Angioneurotic Oedema Arthralgia C-Reactive Protein Increased Diarrhoea Joint Swelling Lymphocytosis Pruritus Urticaria		Zyban Thyronajod Orgametril	PS C C	Glaxo Wellcome	ORAL ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/21/00ISR Number: 3614109-3Report Type:Expedited (15-DaCompany Report #A0109223A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angiopathy		Zyban	PS	Glaxo Wellcome	ORAL
39 DAY		Arthralgia		Ibuprofen	C		
		Bone Density Decreased					
		Immobile					
		Inflammation					
		Intervertebral Disc					
		Degeneration					
		Intervertebral Disc					
		Disorder					
		Leukocytoclastic					
		Vasculitis					
		Osteoporosis					
		Pain In Extremity					
		Polyarthritits					
		Pruritus					
		Purpura					
		Rash Erythematous					
		Sneezing					
		Throat Irritation					
		Urticaria					
		White Blood Cell Count					
		Increased					

Date:11/21/00ISR Number: 3614111-1Report Type:Expedited (15-DaCompany Report #A0128188A  
Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Adrenal Disorder		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG TWICE							
Initial or Prolonged		Aggression					
PER DAY	1 YR						
		Atrial Fibrillation		Neurontin	C		
		Blood Disorder		Coumadin	C	Glaxo Wellcome	
		Constipation		Ativan	C		
		Coordination Abnormal		Ambien	C		
		Depressed Mood		Restoril	C		
		Difficulty In Walking		T3	C	Glaxo Wellcome	

UNKNOWN

Disorientation  
Fatigue  
Hyperthyroidism  
Libido Decreased  
Loss Of Consciousness  
Malaise  
Restlessness  
Sexual Dysfunction  
Sinus Congestion  
Sleep Disorder  
Speech Disorder  
Urinary Tract Infection

Date:11/21/00ISR Number: 3614124-XReport Type:Expedited (15-DaCompany Report #B0090537A  
Age:62 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG TWICE	Fall		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged PER DAY	Femoral Neck Fracture Loss Of Consciousness					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/21/00ISR Number: 3615555-4Report Type:Expedited (15-DaCompany Report #B0091703A

Age:28 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL Initial or Prolonged	Hyperventilation Pyrexia	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:11/21/00ISR Number: 3615559-1Report Type:Expedited (15-DaCompany Report #D0010443A

Age:37 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1 TABLET / Initial or Prolonged PER DAY/ ORAL	Angioneurotic Oedema Arthralgia C-Reactive Protein Increased Diarrhoea Joint Swelling Lymphocytosis Pruritus Urticaria	Foreign Health Professional	Zyban Thyronajod Lynoestrenol	PS C C	Glaxo Wellcome Inc	ORAL

Date:11/21/00ISR Number: 3615591-8Report Type:Expedited (15-DaCompany Report #A0129577A

Age:50 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG/ TWICE PER DAY/ ORAL	Asthenia Erectile Dysfunction Fatigue Hypoaesthesia Paraesthesia	Foreign Health Professional	Zyban Cardiac Medication (Formulation Unknown) (Cardiac Medication) Metoprolol Ramipril Metformin	PS SS C C C	Glaxo Wellcome Inc	ORAL

Glibenclamide C  
Omeprazole C  
Gemfibrozil C

Date:11/21/00ISR Number: 3615657-2Report Type:Expedited (15-DaCompany Report #A0127813A  
Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Potassium Decreased	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG, PER		Condition Aggravated	Professional				
DAY, ORAL		Convulsive Threshold Lowered		Potassium Chloride	C		
		Fall		Multivitamin	C		
		Fatigue		Ethanol	C		
		Grand Mal Convulsion					
		Headache					
		Muscle Rigidity					
		Pain					
		Skin Discolouration					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/21/00ISR Number: 3617568-5Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #S00-USA-01104-01

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Anxiety Asthenia Depersonalisation Dizziness Drug Withdrawal Syndrome	Health Professional	Celexa	PS	Forest Laboratories Inc	
150 MG QD PO	Hypertension Palpitations Tachycardia Thinking Abnormal		Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL

Date:11/21/00ISR Number: 3631290-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #S00-USA-01152-01

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose 20 MG QD PO	Tremor	Health Professional	Celexa	PS	Forest Laboratories Inc	ORAL
100 MG QD PO		Company Representative	Wellbutrin (Amfebumatone Hydrochloride)	SS		ORAL
200 MG QD PO			Wellbutrin (Amfebumatone Hydrochloride)	SS		ORAL

Date:11/21/00ISR Number: 3614117-2Report Type:Expedited (15-DaCompany Report #A0131760A  
Age:42 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 350MG PER DAY 8 MON	Alanine Aminotransferase Increased		Wellbutrin	PS	Glaxo Wellcome	ORAL
UNKNOWN	Blood Amylase Increased		Zoloft	C		

Blood Glucose Increased  
Chromaturia  
Hepatitis

Date:11/22/00ISR Number: 3615123-4Report Type:Expedited (15-DaCompany Report #A0132245A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG TWICE		Blindness Transient		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged PER DAY	1 MON	Communication Disorder					
150MG TWICE		Convulsion		Zyban	SS	Glaxo Wellcome	ORAL
PER DAY	3 MON	Dizziness					
		Feeling Hot		Vitamin E	C		
		Hearing Impaired		Aspirin	C		
		Hyperglycaemia		Vitamin C	C		
		Hyperhidrosis		Multivitamin	C		
		Hypoglycaemia					
		Nervous System Disorder					
		Tinnitus					
		Tremor					
		Tunnel Vision					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/22/00ISR Number: 3615125-8Report Type:Expedited (15-DaCompany Report #B0082488A  
 Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG TWICE Other PER DAY	31 DAY	Eye Irritation Maculopathy Post Procedural Complication Retinal Disorder Retinal Oedema Retinal Vasculitis Retinitis Visual Disturbance		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL

Date:11/22/00ISR Number: 3615139-8Report Type:Expedited (15-DaCompany Report #B0091958A  
 Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG TWICE Initial or Prolonged PER DAY	1 DAY	Unevaluable Event		Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY (INHALATION)	250MCG TWICE PER DAY			Beclomethasone	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)	40MG THREE TIMES PER DAY			Ipratropium	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)				Aminophylline Salbutamol	C C	Glaxo Wellcome	ORAL
40MG PER DAY				Prednisolone	C		ORAL

Eformoterol

C

RESPIRATORY

(INHALATION) 12MCG TWICE

PER DAY

Date:11/22/00ISR Number: 3615140-4Report Type:Expedited (15-DaCompany Report #B0091959A

Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG TWICE		Abdominal Pain		Bupropion	PS	Glaxo Wellcome	ORAL
Initial or Prolonged PER DAY		Constipation					
Other 10MG PER DAY	5 DAY	Diarrhoea		Lisinopril	SS		ORAL
		Dysphagia		Betahistine	C		ORAL
16MG THREE TIMES PER DAY		Rectal Haemorrhage					
10MG THREE TIMES PER DAY		Urticaria		Prochlorperazine	C		ORAL
10MG PER DAY				Clomipramine	C		ORAL
2.5MG PER DAY				Bendrofluazide	C	Glaxo Wellcome	ORAL

Date:11/22/00ISR Number: 3615366-XReport Type:Direct

Company Report #

Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG SR FOR 3DY THEN BID ORAL		Hyperhidrosis		Zyban	PS	Glaxo Wellcome	ORAL
		Sleep Disorder					
		Tremor		Xanax	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/22/00 ISR Number: 3615407-X Report Type:Direct  
 Age:42 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG PO ONE	Cough		Wellbutrin 150mg	PS		ORAL
DOSE		Movement Disorder					
		Urinary Incontinence					

Date:11/22/00 ISR Number: 3616584-7 Report Type:Expedited (15-DaCompany Report #B0091959A  
 Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG (TWICE)	Abdominal Pain Constipation	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Required	PER DAY) ORAL	Diarrhoea					
Intervention to Prevent Permanent Impairment/Damage	10 MG (PER DAY) ORAL	Dysphagia Rectal Haemorrhage		Lisinopril (Lisinopril)	C		ORAL
		Urticaria		Betahistine	C		
				Prochlorperazine	C		
				Clomipramine Hcl	C		
				Bendrofluazide	C		

Date:11/22/00 ISR Number: 3616585-9 Report Type:Expedited (15-DaCompany Report #B0091958A  
 Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG (TWICE)	Chronic Obstructive Airways Disease	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY) ORAL		Exacerbated					
				Beclomethasone Dipropion	C		

Ipratropium Bromide C  
 Aminophylline C  
 Salbutamol Sulphate C  
 Prednisolone C  
 Eformoterol C

Date:11/22/00ISR Number: 3616586-0Report Type:Expedited (15-DaCompany Report #B0091876A  
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Metrorrhagia	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Intervention to 150 MG (TWICE Prevent Permanent PER DAY) ORAL Impairment/Damage				Norimin	C		

Date:11/22/00ISR Number: 3616587-2Report Type:Expedited (15-DaCompany Report #B0091876A  
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Metrorrhagia	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Intervention to 150 MG (TWICE Prevent Permanent PER DAY) ORAL Impairment/Damage				Norimin	C		

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Freedom Of Information (FOI) Report

Date:11/22/00ISR Number: 3616588-4Report Type:Expedited (15-DaCompany Report #B0091290A  
Age:57 YR Gender:Female I/FU:I

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG (TWICE PER DAY) ORAL	Condition Aggravated Rheumatic Fever  Systemic Lupus Erythematosus	Foreign Study  Health  Professional	Bupropion Hydrochloride   Prednisone Azathioprine Flupirtine Maleate Atorvastatin Calcium Magnerot N Dorzolamide Hcl	PS    C C C C C C	Glaxo Wellcome Inc	ORAL

Date:11/22/00ISR Number: 3616815-3Report Type:Expedited (15-DaCompany Report #B0082488A  
Age:32 YR Gender:Male I/FU:F

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY Required / ORAL Intervention to Prevent Permanent Impairment/Damage	Maculopathy Retinal Deposits Retinal Disorder Retinal Oedema Retinitis Vasculitis Visual Disturbance	Foreign Study Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:11/22/00ISR Number: 3617771-4Report Type:Expedited (15-DaCompany Report #A036897  
Age:58 YR Gender:Female I/FU:I

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL  300.00 MG	Amnesia Confusional State  Hypotension	Health Professional	Procardia Xl  Bupropion	PS  SS	Pfizer Laboratories Div Pfizer Inc	ORAL  ORAL

TOTAL: BID: ORA

Pneumonia

Sedation

Speech Disorder  
Toxicologic Test Abnormal

Gabapentin SS  
Atenolol SS

Date: 11/22/00 ISR Number: 3617824-0 Report Type: Expedited (15-DaCompany Report #A0132245A

Age: 52 YR Gender: Male I/FU: I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY		Blindness Transient	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
/ ORAL		Convulsion					
		Deafness					
150 MG / TWICE PER DAY		Dizziness Feeling Hot Hyperglycaemia		Zyban Tablet - Zyban (Bupropion Hydrochloride)	SS		ORAL
/ ORAL		Hyperhidrosis					
		Hypoglycaemia					
		Nervous System Disorder		Multivitamin	C		
		Speech Disorder		Vitamin E	C		
		Tinnitus		Ascorbic Acid	C		
		Tremor		Aspirin	C		
		Tunnel Vision					

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Freedom Of Information (FOI) Report

Date:11/22/00ISR Number: 3615129-5Report Type:Expedited (15-DaCompany Report #B0091290A  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG TWICE		Condition Aggravated Rheumatic Fever		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
PER DAY		Systemic Lupus					
5MG PER DAY		Erythematosus		Decortin	C		
100MG PER DAY				Katadolon	C		
10MG PER DAY				Sortis	C		
20MG TWICE				Trusopt	C		
PER DAY							
48.6MG TWICE				Magnerot N	C		
PER DAY							
50MG PER DAY				Imurek	C	Glaxo Wellcome	

Date:11/22/00ISR Number: 3615138-6Report Type:Expedited (15-DaCompany Report #B0091876A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG TWICE		Metrorrhagia		Zyban	PS	Glaxo Wellcome	ORAL
PER DAY				Norimin	C		ORAL

Date:11/24/00ISR Number: 3616420-9Report Type:Expedited (15-DaCompany Report #A0128188A  
Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Adrenal Disorder Consumer Wellbutrin Sr PS Glaxo Wellcome Inc ORAL  
150 MG/ TWICE  
Initial or Prolonged Aggression  
PER DAY/ ORAL

Atrial Fibrillation	Warfarin Sodium	C
Blood Disorder	Lorazepam	C
Constipation	Zolpidem Tartrate	C
Coordination Abnormal	Temazepam	C
Difficulty In Walking		
Disorientation		
Dysarthria		
Fatigue		
Hyperthyroidism		
Insomnia		
Libido Decreased		
Loss Of Consciousness		
Malaise		
Restlessness		
Sexual Dysfunction		
Sinus Congestion		
Speech Disorder		
Urinary Tract Infection		

Date:11/24/00ISR Number: 3616450-7Report Type:Expedited (15-DaCompany Report #A0109223A  
Age:62 YR Gender:Male I/FU:F

Outcome	PT
Other	Arthralgia
	Arthritis
	Back Pain
	Bone Density Decreased
	Condition Aggravated
	Decreased Activity

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL		Dysphonia Intervertebral Disc Degeneration Leukocytoclastic Vasculitis	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Osteoporosis Pain In Extremity Polyarthritus Pruritus Purpura Rash Erythematous Rheumatoid Arthritis Sneezing Urticaria Vasculitis White Blood Cell Count Increased		Ibuprofen	C		

Date:11/24/00ISR Number: 3616474-XReport Type:Expedited (15-DaCompany Report #A0131760A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 350 MG PER DAY; ORAL	8 MON	Alanine Aminotransferase Increased Blood Amylase Increased Blood Glucose Increased Chromaturia Hepatitis	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
				Sertraline Hydrochloride	C		

Date:11/24/00ISR Number: 3616888-8Report Type:Expedited (15-DaCompany Report #B0090537A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG (TWICE		Fall Feeling Abnormal	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

PER DAY) ORAL  
Femoral Neck Fracture Professional  
Loss Of Consciousness

Date:11/27/00ISR Number: 3616080-7Report Type:Expedited (15-DaCompany Report #A0131639A  
Age:73 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG PER DAY 2 YR	Hyperkalaemia		Zantac	PS	Glaxo Wellcome	ORAL
Initial or Prolonged 150MG PER DAY	Intestinal Ischaemia		Zyban	SS	Glaxo Wellcome	ORAL
	Rectal Haemorrhage		Theophylline Multivitamin	C C		

Date:11/27/00ISR Number: 3616091-1Report Type:Expedited (15-DaCompany Report #B0086711A  
Age:38 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 6 DAY	Asthenia		Bupropion	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged	Fatigue Muscle Spasms Paraesthesia Respiratory Alkalosis Tetany		Atenolol Benzodiazepines	C C		ORAL

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Freedom Of Information (FOI) Report

Date:11/27/00ISR Number: 3616092-3Report Type:Expedited (15-DaCompany Report #B0088230A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coma Convulsion Hypotension Metabolic Acidosis Overdose		Zyban	PS	Glaxo Wellcome	

Date:11/27/00ISR Number: 3616094-7Report Type:Expedited (15-DaCompany Report #B0089068A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Brain Oedema		Bupropion	PS	Glaxo Wellcome	ORAL
150MG PER DAY	57 DAY	Coma Haematoma Haemorrhagic Stroke Headache Hemiparesis Ruptured Cerebral Aneurysm Sedation Subarachnoid Haemorrhage					

Date:11/27/00ISR Number: 3616097-2Report Type:Expedited (15-DaCompany Report #B0090063A

Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Aphasia		Zyban	PS	Glaxo Wellcome	ORAL
2TAB PER DAY	14 DAY	Cerebral Artery Embolism Cerebral Infarction Hemiplegia					
Initial or Prolonged							
Other							

Date:11/27/00ISR Number: 3616109-6Report Type:Expedited (15-DaCompany Report #B0092034A

Age:52 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Joint Swelling		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG TWICE							
PER DAY	9	DAY					
3UNIT PER DAY				Sulphasalazine	SS		ORAL
200UG THREE				Metronidazole	C		ORAL
TIMES PER DAY	6	DAY					
150MG PER DAY				Indomethacin	C		ORAL
10MG PER DAY				Prednisolone	C		ORAL

Date:11/27/00ISR Number: 3616667-1Report Type:Expedited (15-DaCompany Report #D0010841A  
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Amnesia	Health	Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown	9	DAY					
Hospitalization -		Cardiac Arrest	Professional				
Initial or Prolonged		Electroencephalogram					
Other		Abnormal Epilepsy					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/27/00ISR Number: 3617031-1Report Type:Expedited (15-DaCompany Report #A0131639A  
Age:73 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / PER	Colitis Ischaemic	Health	Zantac 150	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged DAY / ORAL	Hyperkalaemia	Professional				
	Rectal Haemorrhage		Zyban Tablet - Zyban (Bupropion Hydrochloride)	SS		ORAL
150 MG / PER						
DAY / ORAL			Theophylline	C		
			Multivitamin	C		

Date:11/27/00ISR Number: 3617253-XReport Type:Expedited (15-DaCompany Report #B0091999A  
Age:60 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG ORAL	Aspiration Cardio-Respiratory Arrest	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
	Depressed Level Of Consciousness		Aspirin	C		
	Salivary Hypersecretion		Ranitidine Hydrochloride	C		
	Skin Discolouration		Co-Dydramol	C		
	Ventricular Fibrillation		Nitrazepam	C		

Date:11/27/00ISR Number: 3617316-9Report Type:Expedited (15-DaCompany Report #2000035041GB  
Age:52 YR Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Rheumatoid Arthritis Skin Nodule	Foreign Health	Azulfidine	PS	Pharmacia And Upjohn Co	ORAL
TID, ORAL		Professional	Zyban(Amfebutamone Hydrochloride)	SS		ORAL
150		Other				

MG,BID,ORAL

Metronidazole C  
Indocid(Indometacin) C  
Delta-Cortril C

Date:11/27/00ISR Number: 3617400-XReport Type:Expedited (15-DaCompany Report #B0092002A

Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Arthralgia	Foreign	Bupropion			
150 MG ORAL		Difficulty In Walking	Other	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Erythema		Thyroxine Sodium	C		
		Oedema		Evening Primrose Oil	C		
		Pruritus					
		Synovitis					
		Viral Infection					

Date:11/27/00ISR Number: 3617401-1Report Type:Expedited (15-DaCompany Report #B0092034A

Age:52 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Arthropathy	Foreign	Bupropion			
Intervention to		Rheumatoid Arthritis	Other	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE							
Prevent Permanent							
PER DAY) ORAL							
Impairment/Damage				Sulphasalazine			

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ORAL (Sulfasalazine) SS ORAL  
 Metronidazole C  
 Indomethacin C  
 Prednisolone C

Date:11/27/00ISR Number: 3617403-5Report Type:Expedited (15-DaCompany Report #B0092029A  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG ORAL	Atrial Fibrillation	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Kapake	C		

Date:11/27/00ISR Number: 3617404-7Report Type:Expedited (15-DaCompany Report #D0011104A  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1 TABLET (PER DAY) ORAL	Arthralgia Dyspnoea Face Oedema Myalgia Pharyngolaryngeal Pain Pruritus Urine Analysis Abnormal Urticaria	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:11/27/00ISR Number: 3617406-0Report Type:Expedited (15-DaCompany Report #B0088740A  
 Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG / TWICE PER DAY	Diabetes Mellitus Inadequate Control Vision Blurred	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

/ ORAL

Insulin Lispro C  
Human Insulin C

Date:11/27/00ISR Number: 3617571-5Report Type:Expedited (15-DaCompany Report #B0086711A  
Age:38 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150 MG/ORAL	Asthenia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - Initial or Prolonged	Fatigue Muscle Spasms Paraesthesia Respiratory Alkalosis Tetany	Health Professional	Atenolol Benzodiazepines	C C		

Date:11/27/00ISR Number: 3617573-9Report Type:Expedited (15-DaCompany Report #B0088230A  
Age: Gender:Male I/FU:F

Outcome	PT
Other	Acidosis Coma Convulsion Hypotension Metabolic Acidosis



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Overdose

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
50 TABLET		Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	

Date:11/27/00ISR Number: 3617574-0Report Type:Expedited (15-DaCompany Report #B0089068A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death ORAL		Brain Oedema	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Cerebral Infarction Coma Haematoma Headache Hemiparesis Ruptured Cerebral Aneurysm Sedation Subarachnoid Haemorrhage					

Date:11/27/00ISR Number: 3617575-2Report Type:Expedited (15-DaCompany Report #B0090063A  
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL Initial or Prolonged Other		Aphasia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Cerebral Artery Embolism Cerebral Infarction Hemiplegia	Health Professional				

Date:11/27/00ISR Number: 3616093-5Report Type:Expedited (15-DaCompany Report #B0088740A  
Age:24 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Vision Blurred		Zyban	PS	Glaxo Wellcome	ORAL
150MG TWICE							
PER DAY	9	DAY		Humalog	C		
				Humulin	C		

Date:11/27/00ISR Number: 3616106-0Report Type:Expedited (15-DaCompany Report #B0091999A  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Cardio-Respiratory Arrest		Zyban	PS	Glaxo Wellcome	ORAL
39 DAY							
Initial or Prolonged		Cyanosis		Aspirin	C		ORAL
		Depressed Level Of		Ranitidine	C	Glaxo Wellcome	ORAL
		Consciousness		Co-Dydramol	C	Glaxo Wellcome	ORAL
		Ventricular Fibrillation		Nitrazepam	C		ORAL

Date:11/27/00ISR Number: 3616107-2Report Type:Expedited (15-DaCompany Report #B0092002A  
Age:28 YR Gender:Female I/FU:I

Outcome	PT
Disability	Arthralgia
	Autoimmune Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG PER DAY	14 DAY	Difficulty In Walking Erythema Oedema	Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG PER DAY		Pruritus Synovitis	Thyroxine	C	Glaxo Wellcome	ORAL
			Evening Primrose Oil	C		ORAL

Date:11/27/00ISR Number: 3616108-4Report Type:Expedited (15-DaCompany Report #B0092029A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150MG PER DAY	9 DAY	Atrial Fibrillation	Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
2TAB PER DAY				Kapake	C		ORAL

Date:11/27/00ISR Number: 3616113-8Report Type:Expedited (15-DaCompany Report #D0011104A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1TAB PER DAY		Arthralgia	Zyban	PS	Glaxo Wellcome	ORAL
			Dyspnoea Face Oedema Haematuria Myalgia Pharyngolaryngeal Pain Pruritus Urticaria White Blood Cells Urine Positive				

Date:11/28/00ISR Number: 3617660-5Report Type:Expedited (15-DaCompany Report #A0130408A  
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Dermatitis		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	5 DAY	Dyspnoea Malaise Nervousness Vision Blurred					

Date:11/28/00ISR Number: 3617667-8Report Type:Expedited (15-DaCompany Report #B0088745A  
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 20 DAY		Chest Pain		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - RESPIRATORY		Electrocardiogram T Wave		Ipratropium Bromide	C	Glaxo Wellcome	
Initial or Prolonged (INHALATION)	500MCG Six	Inversion					
Disability times per day		Myocardial Infarction		Salbutamol	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)	5MG Six times			Aspirin	C		ORAL
per day				Bendrofluazide	C	Glaxo Wellcome	ORAL
75MG Per day				Prednisolone	C		ORAL
2.5MG Per day				Beclomethasone	C	Glaxo Wellcome	
30MG Per day							
RESPIRATORY (INHALATION)	2PUFF Twice						

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Freedom Of Information (FOI) Report

per day

Date:11/28/00ISR Number: 3617672-1Report Type:Expedited (15-DaCompany Report #B0091225A

Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1200MG				Zyban	PS	Glaxo Wellcome	ORAL
cumulative dose	6	DAY					
			Coordination Abnormal				
			Delusion				
			Diarrhoea				
			Dry Mouth				
			Dysarthria				
			Extrapyramidal Disorder				
			Hypotension				
			Insomnia				
			Muscle Rigidity				
			Pneumonia Aspiration				
			Sedation				

Date:11/28/00ISR Number: 3617676-9Report Type:Expedited (15-DaCompany Report #B0092066A

Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG As Initial or Prolonged directed				Zyban	PS	Glaxo Wellcome	
			Hyperhidrosis				
			Hypertension				
			Ventricular Hypertrophy				
			Visual Disturbance				

Date:11/28/00ISR Number: 3617681-2Report Type:Expedited (15-DaCompany Report #B0092404A

Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other				Zyban	PS	Glaxo Wellcome	ORAL
			Asthma				

10MG Per day	Peak Expiratory Flow Rate	Prednisolone	C		ORAL
	Decreased	Fluticasone	C	Glaxo Wellcome	
RESPIRATORY					
(INHALATION)	1000MCG Twice				
per day		Salmeterol	C	Glaxo Wellcome	
RESPIRATORY					
(INHALATION)	50MCG Twice				
per day		Salbutamol	C	Glaxo Wellcome	
RESPIRATORY					
(INHALATION)	200MCG As				
required		Quinine	C		
		Nasonex	C		

Date:11/28/00ISR Number: 3617684-8Report Type:Expedited (15-DaCompany Report #B0092571A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Suicide Attempt		Zyban	PS	Glaxo Wellcome	

Date:11/28/00ISR Number: 3617686-1Report Type:Expedited (15-DaCompany Report #D0010583A

Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	7 DAY	Visual Disturbance Vitreous Disorder		Zyban	PS	Glaxo Wellcome	ORAL

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Freedom Of Information (FOI) Report

1TAB Twice  
 per day 4 DAY  
 Bactrim C Glaxo Wellcome ORAL

Date:11/29/00ISR Number: 3618203-2Report Type:Expedited (15-DaCompany Report #A0117940A  
 Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Acne Cystic		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day 3 DAY		Arthropod Bite					
		Asthenia		Soma	C		
		Bronchospasm		Valium	C		
		Burning Sensation					
		Chest Pain					
		Dermatitis					
		Dizziness					
		Drug Hypersensitivity					
		Dyspnoea Exertional					
		Ear Haemorrhage					
		Eye Disorder					
		Eye Irritation					
		Eyelid Oedema					
		Folliculitis					
		Haemorrhage					
		Hypoventilation					
		Infection					
		Irritability					
		Oedema					
		Pain					
		Palpitations					
		Pruritus					
		Rash Erythematous					
		Rash Papular					
		Rash Pustular					
		Respiratory Disorder					
		Scratch					
		Skin Lesion					
		Sunburn					
		Toxic Shock Syndrome					
		Urticaria					

Date:11/29/00ISR Number: 3618215-9Report Type:Expedited (15-DaCompany Report #A0132668A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Bupropion	PS	Glaxo Wellcome	ORAL
150MG Twice		Vaginal Haemorrhage					
per day							

Date:11/29/00ISR Number: 3618224-XReport Type:Expedited (15-DaCompany Report #B0091997A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Angioneurotic Oedema		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day 18 DAY		Bronchial Obstruction					
Initial or Prolonged		Eyelid Oedema					
		Pruritus					
		Rash Generalised					
		Urticaria					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/29/00ISR Number: 3618230-5Report Type:Expedited (15-DaCompany Report #B0092355A  
Age:60 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Toxic Epidermal Necrolysis		Zyban Statines	PS C	Glaxo Wellcome	ORAL

Date:11/29/00ISR Number: 3618231-7Report Type:Expedited (15-DaCompany Report #B0092499A  
Age:35 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Blindness Blister Rash Pruritic		Zyban	PS	Glaxo Wellcome	

Date:11/29/00ISR Number: 3619214-3Report Type:Expedited (15-DaCompany Report #2000AP05073  
Age:61 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 10 MG DAILY Required PO Intervention to 150 MG BID PO Prevent Permanent Impairment/Damage	Abdominal Pain Constipation Diarrhoea Dysphagia Rectal Haemorrhage Urticaria	Foreign Health Professional Other	Zestril  Bupropion Betahistine Prochlorperazine Anafranil Bendrofluazide	PS  SS C C C C	Astrazeneca Pharmaceuticals Lp	ORAL  ORAL

Date:11/30/00ISR Number: 3618945-9Report Type:Expedited (15-DaCompany Report #A0131760A  
Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 350MG Per day 8 MON	Alanine Aminotransferase		Wellbutrin	PS	Glaxo Wellcome	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/00ISR Number: 3625779-8Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #A0114567A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150 MG /TWICE Other PER DAY / ORAL	Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:11/30/00ISR Number: 3625785-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0114618A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG/ TWICE PER DAY / ORAL	Convulsion Hypoglycaemia	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:11/30/00ISR Number: 3625793-2Report Type:Periodic  
Age:17 YR Gender:Male I/FU:I

Company Report #A0114245A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG /THREE TIMES PER DA /ORAL 150 MG/ ORAL	Convulsion Drug Interaction Medication Error	Health Professional Company Representative	Wellbutrin Sr   Nortriptyline (Nortriptyline)	PS   SS	Pharmacia And Upjohn Co	ORAL   ORAL

Date:11/30/00ISR Number: 3625795-6Report Type:Periodic Company Report #A0114247A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company	Wellbutrin Sr Furosemide Tablet (Furosemide)	PS	Glaxo Wellcome Inc	ORAL
ORAL			Representative		SS		

Date:11/30/00ISR Number: 3625800-7Report Type:Periodic Company Report #A0114119A

Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG/		Head Injury	Professional Company				
VARIABLE			Representative	Paroxetine Hydrochloride	C		
DOSE/ ORAL							

Date:11/30/00ISR Number: 3625804-4Report Type:Periodic Company Report #A0113009A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion	Health Professional Company	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
4500 MG /		Overdose	Professional Company				
Initial or Prolonged			Representative				
SINGLE DOSE/							
Other							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/00ISR Number: 3625807-XReport Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0112414A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 200 MG /TWICE Initial or Prolonged PER DAY /		Grand Mal Convulsion	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL	6 MON	Company Representative	Sertraline Hydrochloride (Sertraline Hydrochloride)	SS		ORAL
50 MG / PER DAY/ ORAL	3 DAY					

Date:11/30/00ISR Number: 3625808-1Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #A0112470A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200 MG /TWICE PER DAY/ ORAL		Grand Mal Convulsion	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
		Health Professional Company Representative	Quetiapine Fumarate	C		

Date:11/30/00ISR Number: 3625810-XReport Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0112385A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG/TWICE PER DAY/ ORAL		Convulsion	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
	DAY	Company Representative				

Date:11/30/00ISR Number: 3625812-3Report Type:Periodic Company Report #A0112386A

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG /			Professional				
TWICE PER DAY			Company				
/ ORAL	DAY		Representative				

Date:11/30/00ISR Number: 3625815-9Report Type:Periodic Company Report #A0111400A

Age:11 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
100 MG / PER			Professional				
DAY / ORAL			Company	Carbamazepine	C		
			Representative				

Date:11/30/00ISR Number: 3625819-6Report Type:Periodic Company Report #A0111423A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
300 MG / PER			Professional				
DAY / ORAL			Company				
			Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/00ISR Number: 3625822-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0111466A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Grand Mal Convulsion	Professional				
150 M G /			Company				
THREE TIMES			Representative				
PER DAY /							
ORAL							

Date:11/30/00ISR Number: 3625823-8Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:I

Company Report #A0110050A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Hospitalization -		Abdominal Pain	Professional				
150 MG /		Difficulty In Walking					
Initial or Prolonged							
TWICE PER DAY		Erythema Multiforme					
Disability		Face Oedema					
/ ORAL		Oedema					
Required		Pain In Extremity					
Intervention to		Pharyngeal Oedema					
Prevent Permanent		Tenderness					
Impairment/Damage		Urinary Tract Infection					
		Urticaria					

Date:11/30/00ISR Number: 3625824-XReport Type:Periodic  
 Age: Gender:Unknown I/FU:I

Company Report #A0109827A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Convulsion	Professional				
150 MG /							
TWICE PER DAY							

/ ORAL

Company  
Representative

Date:11/30/00ISR Number: 3625825-1Report Type:Periodic Company Report #A0109828A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG /			Professional				
TWICE PER DAY			Company				
/ ORAL	2	MON	Representative				

Date:11/30/00ISR Number: 3625826-3Report Type:Periodic Company Report #A0109899A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Dyskinesia	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
100 MG /			Professional				
SINGLE DOSE /		Epilepsy					
ORAL				Cannabis	C		
				Echinacea	C		
				Semisodium Valproate	C		
				Fluoxetine			
				Hydrochloride	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/00ISR Number: 3625828-7Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0103570A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG /			Professional				
Required			Company				
TWICE PER DAY							
Intervention to			Representative	Alprazolam	C		
/ ORAL				Fluoxetine	C		
Prevent Permanent				Hydrochloride	C		
Impairment/Damage				Zolpidem Tartrate	C		

Date:11/30/00ISR Number: 3625829-9Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0105852A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG /			Professional				
TWICE PER DAY		Epileptic Aura					
/ ORAL		Syncope					
		Urinary Incontinence		Nortriptyline	C		
				Citalopram	C		
				Hydrobromide	C		
				Lithium Salt	C		

Date:11/30/00ISR Number: 3625831-7Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #A0108726A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Grand Mal Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG /			Professional				
Other			Company				
TWICE PER DAY							
Required							
/ ORAL							

Intervention to Prevent Permanent Impairment/Damage  
Representative  
Prednisone  
Hydrochloroquine  
C  
C

Date:11/30/00ISR Number: 3627138-0Report Type:Periodic Company Report #A0121035A  
Age:49 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100 MG / Initial or Prolonged TWICE PER DAY  / ORAL	Pancreatitis	Health  Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
3 MON						

Date:11/30/00ISR Number: 3627139-2Report Type:Periodic Company Report #A0120808A  
Age:32 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY Other / ORAL	Convulsion  Delirium  Intentional Misuse  Suicide Attempt	Health  Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/00ISR Number: 3627142-2Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0120790A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - VARIABLE DOSE	Confusional State	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged / ORAL	Convulsion	Professional				
	Delirium		Baclofen Oxycodone Hydrochloride	C C		

Date:11/30/00ISR Number: 3627144-6Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0120622A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL	Alanine Aminotransferase	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged	Increased Blood Alkaline Phosphatase Increased Grand Mal Convulsion Intentional Misuse	Professional Company Representative				

Date:11/30/00ISR Number: 3627146-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0119914A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other	Convulsion Hallucination	Health Professional Company Representative	Wellbutrin Sr Venlafaxine Hydrochloride	PS C	Glaxo Wellcome Inc	

Date:11/30/00ISR Number: 3627147-1Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0118761A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Other 150 MG / SEE		Abnormal Faeces	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
TEXT / ORAL	5	DAY	Defaecation Urgency	Professional			
ORAL			Feeling Hot	Company	Levofloxacin Tablet	SS	ORAL
			Grand Mal Convulsion	Representative	Oral Contraceptive	C	
			Hyperhidrosis				
			Nausea				
			Tremor				
			Urinary Tract Infection				

Date:11/30/00ISR Number: 3627149-5Report Type:Periodic Company Report #A0118454A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / PER		Anxiety	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged DAY / ORAL	4 WK	Asthenia					
		Blood Pressure Increased		Citalopram			
		Dissociation		Hydrobromide			
		Disturbance In Attention		(Formulation			
		Dizziness		Unknown)	SS		
		Drug Withdrawal Syndrome					
		Palpitations					
		Simple Partial Seizures					
		Syncope					
		Tachycardia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/00ISR Number: 3627152-5Report Type:Periodic  
Age:21 YR Gender:Female I/FU:I

Company Report #A0117091A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150 MG / Hospitalization - TWICE PER DAY Initial or Prolonged / ORAL 3 YR Required Intervention to Prevent Permanent Impairment/Damage	Drug Effect Decreased Grand Mal Convulsion Pyrexia	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
			Cetirizine Hydrochloride Salbutamol Sulphate Fluticasone Propionate	C C C		

Date:11/30/00ISR Number: 3627153-7Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0116851A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL Initial or Prolonged	Convulsion	Health Professional Company Representative	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:11/30/00ISR Number: 3627155-0Report Type:Periodic  
Age:54 YR Gender:Male I/FU:I

Company Report #A0116644A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability 400 MG / PER DAY / ORAL 6 WK 250 MG / FOUR TIMES PER DAY	Tic Tremor	Health Professional	Wellbutrin Sr Lithium Salt Methadone Hydrochloride	PS SS SS	Glaxo Wellcome Inc	ORAL ORAL

ORAL

Tamsulosin Hcl C  
Psychiatric  
Consultation C

Date:11/30/00ISR Number: 3627161-6Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #A0116263A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
ORAL		Grand Mal Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
			Professional Company Representative	Antidepressant	C		

Date:11/30/00ISR Number: 3627162-8Report Type:Periodic  
Age:29 YR Gender:Male I/FU:I

Company Report #A0116264A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
ORAL		Grand Mal Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
			Professional Company Representative	Antidepressant	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/00ISR Number: 3627163-XReport Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0116265A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Grand Mal Convulsion	Professional Company Representative	Antidepressant	C		
ORAL							

Date:11/30/00ISR Number: 3627164-1Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0116266A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Grand Mal Convulsion	Professional Company Representative	Antidepressant	C		
ORAL							

Date:11/30/00ISR Number: 3627167-7Report Type:Periodic  
Age: Gender:Not SpecifiedI/FU:I

Company Report #A0116329A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Dermatitis	Professional				
ORAL		Hospitalization - Initial or Prolonged					

Date:11/30/00ISR Number: 3627170-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0116346A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Convulsion	Professional Company Representative				
ORAL							

Date:11/30/00ISR Number: 3627173-2Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0116040A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/PER Initial or Prolonged DAY/ORAL, 225 MG / PER DAY/ ORAL	2 WK	Back Injury Convulsion Drug Level Changed Vomiting	Consumer	Wellbutrin Sr  Carbamazepine Primidone Topiramate	PS  C C C	Glaxo Wellcome Inc	ORAL

Date:11/30/00ISR Number: 3627176-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0114700A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ORAL Initial or Prolonged		Depression Drug Effect Decreased Weight Increased	Consumer	Wellbutrin Sr  Semisodium Valproate Citalopram Hydrobromide	PS  C C C	Glaxo Wellcome Inc	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/00ISR Number: 3627178-1Report Type:Periodic  
Age:7 MON Gender:Female I/FU:I

Company Report #A0114779A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / PER Initial or Prolonged DAY / ORAL Other	Convulsion	Health  Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:11/30/00ISR Number: 3627181-1Report Type:Periodic  
Age:6 YR Gender:Male I/FU:I

Company Report #A0114780A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 100 MG / Required TWICE PER Intervention to DAY/ ORAL 6 MON Prevent Permanent Impairment/Damage	Drug Ineffective  Dyskinesia  Grunting Tic	Health  Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:11/30/00ISR Number: 3627373-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0103624A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY Other / ORAL	Convulsion  Nuclear Magnetic  Resonance Imaging  Abnormal	Health  Professional  Company  Representative	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:11/30/00ISR Number: 3627377-9Report Type:Periodic  
Age:75 YR Gender:Male I/FU:I

Company Report #A0105553A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL	6 MON	Confusional State Convulsion Epileptic Aura Syncope	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:11/30/00ISR Number: 3627379-2Report Type:Periodic Company Report #A0105559A  
 Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / ORAL Initial or Prolonged		Hyponatraemia Inappropriate Antidiuretic Hormone Secretion	Health Professional Company Representative	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:11/30/00ISR Number: 3627381-0Report Type:Periodic Company Report #A0107151A  
 Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG / ORAL Initial or Prolonged		Anaphylactic Reaction Choking Dizziness Dyspnoea Food Allergy Hypersensitivity Retching	Health Professional Company Representative	Wellbutrin Sr Methylphenidate Chl	PS C	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/00ISR Number: 3627383-4Report Type:Periodic  
Age:12 MON Gender:Female I/FU:I

Company Report #A0107580A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
75 MG / TWICE							
PER DAY /		Maternal Drugs Affecting					
ORAL		Foetus					
				Prenatal	C		

Date:11/30/00ISR Number: 3627385-8Report Type:Periodic  
Age:17 YR Gender: I/FU:I

Company Report #A0124973A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY			Professional				
/ ORAL			Company				
			Representative	Wellbutrin Tablet (Bupropion Hydrochloride)	SS		ORAL
75 MG / TWICE							
PER DAY /							
ORAL							

Date:11/30/00ISR Number: 3627389-5Report Type:Periodic  
Age:46 YR Gender:Female I/FU:F

Company Report #A0108085A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Grand Mal Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
450 MG / PER							
Initial or Prolonged			Professional				
DAY / ORAL			Company	Quetiapine Fumarate	C		
Disability							

Required  
Intervention to  
Prevent Permanent  
Impairment/Damage

Representative

Temazepam

C

Date:11/30/00ISR Number: 3627393-7Report Type:Periodic  
Age:39 YR Gender:Female I/FU:F

Company Report #A0105971A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other / ORAL		Cyanosis	Professional				
		Drooling					
		Tremor		Ortho Tri-Cyclen	C		

Date:11/30/00ISR Number: 3627396-2Report Type:Periodic  
Age:15 YR Gender:Female I/FU:F

Company Report #A0104843A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG / PER DAY / ORAL		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
			Professional				
			Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/00ISR Number: 3627399-8Report Type:Periodic  
Age:13 YR Gender:Male I/FU:F

Company Report #A0101377A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
75 SEE TEXT /		Memory Impairment	Professional				
TWICE PER DAY		Tongue Disorder	Company				
/ ORAL			Representative	Desipramine	C		
				Methylphenidate Hcl	C		

Date:11/30/00ISR Number: 3627472-4Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #PRIUSA2000008297

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Defaecation Urgency	Health	Levaquin	PS	Rw Johnson	
		Grand Mal Convulsion	Professional			Pharmaceutical	
		Hyperhidrosis				Research Institute	
		Nausea				Div Ortho Pharm	ORAL
ORAL		Paraesthesia		Wellbutrin			
		Tremor		(Amfebutamone			
150 MG,				Hydrochloride)	SS		ORAL
DIALY, ORAL	5	DAY		Oral Contraceptive			
				(Oral Contraceptive			
				Nos)	C		

Date:11/30/00ISR Number: 3627524-9Report Type:Periodic  
Age:19 YR Gender:Male I/FU:I

Company Report #A0123848A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL			Professional	Methylphenidate Hcl	C		
			Company				

Representative

Date:11/30/00ISR Number: 3627528-6Report Type:Periodic Company Report #A0123716A  
 Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / ORAL		Dermatitis	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Urticaria	Professional				

Date:11/30/00ISR Number: 3627532-8Report Type:Periodic Company Report #A0123213A  
 Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG /		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
TWICE PER			Professional				
DAY/ ORAL				Golden Seal (Formulation Unknown) (Golden Seal)	SS		ORAL
ORAL				Cannabis	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/00ISR Number: 3627536-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0123216A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Convulsion					
ORAL		Drug Interaction	Professional Company Representative	Cannabis (Formulation Unknown) (Cannabis)	SS		

Date:11/30/00ISR Number: 3627538-9Report Type:Periodic  
Age:7 YR Gender:Male I/FU:I

Company Report #A0123110A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Life-Threatening		Grand Mal Convulsion					
200 MG / PER			Professional				
Hospitalization -			Company Representative				
DAY/ ORAL							
Initial or Prolonged							
Required							
Intervention to							
Prevent Permanent							
Impairment/Damage							

Date:11/30/00ISR Number: 3627541-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0123131A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Convulsion					
150 MG /			Professional				
TWICE PER DAY			Company				
/ ORAL			Representative	Venlafaxine Hydrochloride	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY/ ORAL	Agitation Anxiety Confusional State Disturbance In Attention Insomnia Irritability Memory Impairment Paranoia Thinking Abnormal	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG/ ORAL	Convulsion Insomnia Tooth Disorder	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/00ISR Number: 3627548-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0121951A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Accidental Overdose	Health Professional Company Representative	Wellbutrin	PS	Glaxo Wellcome Inc	

Date:11/30/00ISR Number: 3627552-3Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0121920A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Anxiety	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL							
Hospitalization - Initial or Prolonged Required		Back Pain Chest Pain Convulsion	Professional Company Representative	Clonazepam (Formulation Unknown)			
Intervention to Prevent Permanent Impairment/Damage		Electroencephalogram Abnormal Headache		(Clonazepam) Zolpidem Tartrate (Formulation Unknown) (Zolpidem Tartrate)	SS SS		ORAL

10 MG / AT

NIGHT / ORAL

Thyroxine Sodium	C
Loestrin	C
Quinapril Hydrochloride	C

Date:11/30/00ISR Number: 3627737-6Report Type:Periodic  
Age:21 YR Gender:Female I/FU:F

Company Report #A0102154A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
300 MG/ TWICE							
PER DAY/ ORAL		Overdose	Professional				

Date:11/30/00ISR Number: 3627739-XReport Type:Periodic Company Report #A0128927A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL			Professional Company Representative				

Date:11/30/00ISR Number: 3627740-6Report Type:Periodic Company Report #A0128474A  
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
1 TABLET/ TWICE PER DAY/ ORAL	1 YR		Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/00ISR Number: 3627743-1Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0128492A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Convulsion	Professional				
ORAL							

Date:11/30/00ISR Number: 3627746-7Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #A0128275A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Disability		Anaphylactic Reaction	Professional				
150 MG/ PER							
Required		Dermatitis					
DAY/ ORAL							
Intervention to		Urticaria		Oral Contraceptive	C		
Prevent Permanent							
Impairment/Damage							

Date:11/30/00ISR Number: 3627748-0Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #A0128201A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Convulsion	Professional	Ssri	C		
ORAL			Company	Cannabis	C		
			Representative				

Date:11/30/00ISR Number: 3627749-2Report Type:Periodic  
Age:33 YR Gender:Male I/FU:I

Company Report #A0127739A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Fall					
150 MG/ TWICE							

PER DAY/ ORAL	Grand Mal Convulsion	Professional				
	Headache		Amoxicillin			C
	Memory Impairment		Carisoprodol			C
	Motor Dysfunction		Cyclobenzaprine Hcl			C
	Movement Disorder					
	Pain In Extremity					
	Tremor					

Date:11/30/00ISR Number: 3627752-2Report Type:Periodic Company Report #A0127242A  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abdominal Pain Upper	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Fatigue					
PER DAY/ ORAL		Pruritus		Prednisone			
		Tenderness		(Formulation			
		Urticaria		Unknown)	SS		
				Progesterone	C		

Date:11/30/00ISR Number: 3627753-4Report Type:Periodic Company Report #A0127247A  
 Age:31 YR Gender:Male I/FU:I

Outcome	PT	Report Source
Hospitalization - Initial or Prolonged	Dermatitis	Health Professional Company

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Freedom Of Information (FOI) Report

Representative

Dose	Duration	Product	Role	Manufacturer	Route
ORAL		Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:11/30/00ISR Number: 3627777-7Report Type:Periodic Company Report #A0126856A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Convulsion	Health  Professional Company Representative	Wellbutrin Sr  Fluoxetine Hydrochloride (Formulation Unknown) (Fluoxetine Hydrochloride)	PS  SS	Glaxo Wellcome Inc	ORAL

Date:11/30/00ISR Number: 3627780-7Report Type:Periodic Company Report #A0126468A  
 Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG / TWICE PER DAY/ ORAL	3 WK	Convulsion	Health  Professional	Wellbutrin Sr  Beta-Blocker	PS  C	Glaxo Wellcome Inc	ORAL

Date:11/30/00ISR Number: 3627782-0Report Type:Periodic Company Report #A0126488A  
 Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG / PER		Convulsion	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:11/30/00ISR Number: 3627785-6Report Type:Periodic Company Report #A0125786A  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL		Productive Cough	Professional Company Representative	Olanzapine (Formulation Unknown) (Olanzapine)	SS		
5 MG				Theophylline Inhaler Oral Contraceptive	C C C		

Date:11/30/00ISR Number: 3627787-XReport Type:Periodic Company Report #A0125239A  
 Age:43 YR Gender:Female I/FU:I

Outcome	PT
Other	Convulsion Electroencephalogram Abnormal Epilepsy Hypertension Intentional Misuse

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Freedom Of Information (FOI) Report

Dose	Duration	Loss Of Consciousness Memory Impairment Oedema Peripheral	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL			Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
SUBCUTANEOUS DAY /	20 MG / PER			Glatiramer Acetate Injection (Glatiramer Acetate)	SS		
SUBCUTANEOUS				Oxcarbazepine (Formulation Unknown) (Oxcarbazepine)	SS		ORAL
150 MG/ SEE TEXT/ ORAL				Rantidine Hydrochloride Amantadine Hydrochloride Estroven Chromium Picolinate	C C C C		

Date:11/30/00ISR Number: 3627789-3Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #A0124316A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG / ORAL	3 MON	Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Sr Alprazolam (Formulation Unknown) (Alprazolam)	PS SS	Glaxo Wellcome Inc	ORAL

Date:11/30/00ISR Number: 3627790-XReport Type:Periodic Company Report #A0124317A  
Age:10 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
3000 MG / SEE		Overdose	Professional				
TEXT/ ORAL			Company Representative				

Date:11/30/00ISR Number: 3627792-3Report Type:Periodic Company Report #A0124152A  
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Burning Sensation	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL	6 DAY						
Initial or Prolonged		Condition Aggravated		Atenolol	C		
		Gastrooesophageal Reflux		Lisinopril	C		
		Disease		Atorvastatin Calcium	C		
		Insomnia					

Date:11/30/00ISR Number: 3627794-7Report Type:Periodic Company Report #A0124006A  
Age:18 YR Gender:Female I/FU:I

Outcome  
Other  
Required



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Intervention to Prevent Permanent Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Decreased Appetite	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
		Dermatitis	Professional				
		Dizziness		Fiorinal	C		
		Hypersensitivity					
		Oedema Peripheral					
		Orthostatic Hypotension					
		Pruritus					
		Rash Erythematous					
		Urticaria					

Date:11/30/00ISR Number: 3627797-2Report Type:Periodic Company Report #A0123987A  
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Cardiac Failure	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Chest Pain	Professional	Propranolol			
		Myocardial Infarction	Company Representative	Hydrochloride Hydroxyzine	C C		

Date:11/30/00ISR Number: 3627799-6Report Type:Periodic Company Report #A0123847A  
 Age:10 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
			Professional	Methylphenidate Hcl	C		
			Company Representative	Imipramine	C		

Date:12/01/00ISR Number: 3619959-5Report Type:Expedited (15-DaCompany Report #A0132705A  
 Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day		Arthralgia  Cellulitis  Dyspnoea Erythema Feeling Hot Hypersensitivity Pain In Extremity Swelling Urticaria		Zyban	PS	Glaxo Wellcome	ORAL

Date:12/01/00ISR Number: 3619961-3Report Type:Expedited (15-DaCompany Report #B0086031A  
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1TAB Twice per day	12 DAY	Depression  Headache  Oedema Peripheral Physical Assault Suicide Attempt		Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/01/00ISR Number: 3619964-9Report Type:Expedited (15-DaCompany Report #B0092315A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death				Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG Twice		Haemorrhagic Stroke Headache					
per day	48 DAY	Nausea					

Date:12/01/00ISR Number: 3619967-4Report Type:Expedited (15-DaCompany Report #B0092451A  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged				Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG Twice		Hyperreflexia Paraesthesia					
per day	5 DAY	Tachycardia Tremor		Aciclovir	C	Glaxo Wellcome	

Date:12/01/00ISR Number: 3619970-4Report Type:Expedited (15-DaCompany Report #B0092811A  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other				Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Amenorrhoea					
per day	46 DAY	Dysgeusia		Loestrin	C		ORAL
30MCG Per day							

Date:12/01/00ISR Number: 3619971-6Report Type:Expedited (15-DaCompany Report #B0092839A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day	17 DAY	Dermatitis Psoriasis		Zyban	PS	Glaxo Wellcome	ORAL
TOPICAL				Calcipotriol	C		
TOPICAL				Betamethasone	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)				Salbutamol	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)				Beclomethasone	C	Glaxo Wellcome	
15MG Per day				Meloxicam	C		ORAL

Date:12/01/00ISR Number: 3619972-8Report Type:Expedited (15-DaCompany Report #B0092845A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day	6 DAY	Periorbital Oedema		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged RESPIRATORY (INHALATION)	100MCG Per day	Urticaria		Beclomethasone	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)				Salbutamol	C	Glaxo Wellcome	

Date:12/01/00ISR Number: 3619974-1Report Type:Expedited (15-DaCompany Report #D0011022A  
Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 4 WK		Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged 1 YR		Hypertension		Oral Contraceptives	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/01/00ISR Number: 3620886-8Report Type:Expedited (15-DaCompany Report #A0130408A  
Age:46 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG TWICE	Duration Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged PER DAY ORAL	Dyspnoea Malaise Nervousness Vision Blurred					

Date:12/01/00ISR Number: 3621136-9Report Type:Expedited (15-DaCompany Report #B0091225A  
Age:34 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to 150 MG / ORAL	Duration Coordination Abnormal Delusion	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Prevent Permanent Impairment/Damage	Diarrhoea Dry Mouth Dysarthria Extrapyramidal Disorder Feeling Abnormal Hypotension Insomnia Muscle Rigidity Pneumonia Aspiration Sedation	Professional				

Date:12/01/00ISR Number: 3621137-0Report Type:Expedited (15-DaCompany Report #B0092404A  
Age:48 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to ORAL	Duration Asthma Peak Expiratory Flow Rate	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Prevent Permanent Impairment/Damage	Decreased	Professional	Prednisolone Fluticasone Propionate	C C		

Salmeterol Xinafoate C  
 Salbutamol Sulphate C  
 Quinine C  
 Nasonex C

Date:12/01/00ISR Number: 3621138-2Report Type:Expedited (15-DaCompany Report #B0092066A  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/AS		Hyperhidrosis Hypertension	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
DIRECTED		Ventricular Hypertrophy Visual Disturbance	Professional				

Date:12/01/00ISR Number: 3621139-4Report Type:Expedited (15-DaCompany Report #B0092571A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Suicide Attempt	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/01/00ISR Number: 3621140-0Report Type:Expedited (15-DaCompany Report #B0088745A  
Age:49 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - SEE TEXT / Initial or Prolonged ORAL Disability	Chest Pain Electrocardiogram T Wave Inversion Myocardial Infarction	Foreign Health Professional	Bupropion Hydrochloride  Ipratropium Bromide Salbutamol Sulphate Aspirin Bendrofluazide Prednisolone Beclomethasone Dipropion	 PS  C C C C C C	 Glaxo Wellcome Inc	 ORAL

Date:12/01/00ISR Number: 3621277-6Report Type:Expedited (15-DaCompany Report #D0010841A  
Age:24 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 150 MG/ORAL Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage	Amnesia Cardiac Arrest Electroencephalogram Abnormal Grand Mal Convulsion	Foreign Health Professional	Bupropion Hydrochloride	 PS	 Glaxo Wellcome Inc	 ORAL

Date:12/01/00ISR Number: 3621279-XReport Type:Expedited (15-DaCompany Report #D0010583A  
Age:23 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG/TWICE PER DAY/ORAL	Photopsia Visual Disturbance Vitreous Opacities	Foreign Health Professional	Bupropion Hydrochloride  Co-Trimoxazole	 PS  C	 Glaxo Wellcome Inc	 ORAL



Date:12/04/00ISR Number: 3620761-9Report Type:Expedited (15-DaCompany Report #B0093052A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -		Eye Allergy		Fluoxetine	C		
Initial or Prolonged		Rash Pruritic		Chlorhexidine	C		

Date:12/04/00ISR Number: 3620762-0Report Type:Expedited (15-DaCompany Report #B0093061A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxo Wellcome	
Disability		Diplopia					
		Insomnia					
		Tinnitus					
		Tongue Disorder					
		Vision Blurred					
		Visual Disturbance					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/04/00ISR Number: 3620763-2Report Type:Expedited (15-DaCompany Report #B0093064A  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Vision Blurred		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	9	DAY					
				Humalog	C		
				Humulin I	C		

Date:12/04/00ISR Number: 3620764-4Report Type:Expedited (15-DaCompany Report #B0093067A  
Age:67 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rectal Haemorrhage		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	6	DAY		Co-Codamol	C		ORAL
2U Four times							
per day							
400MG Per day				Ibuprofen	C		ORAL

Date:12/04/00ISR Number: 3620810-8Report Type:Expedited (15-DaCompany Report #B0092443A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Chemical Poisoning	Health	Zyban	PS	Glaxo Wellcome	
Initial or Prolonged		Drug Interaction	Professional	Fluoxetine	I		
		Intentional Self-Injury					

Date:12/04/00ISR Number: 3621668-3Report Type:Expedited (15-DaCompany Report #A0131760A  
Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Alanine Aminotransferase Health Wellbutrin Sr PS Glaxo Wellcome Inc ORAL  
350 MG / PER  
Increased Professional  
DAY / ORAL 8 MON  
Blood Amylase Increased Sertraline  
Blood Glucose Increased Hydrochloride C  
Chromaturia  
Hepatitis C

Date:12/04/00ISR Number: 3621683-XReport Type:Expedited (15-DaCompany Report #A0117940A  
Age:47 YR Gender:Male I/FU:F

Outcome PT  
Hospitalization - Acne Cystic  
Initial or Prolonged Asthenia  
Bronchospasm  
Burning Sensation  
Chest Pain  
Dermatitis  
Dizziness  
Dyspnoea Exertional  
Ear Haemorrhage  
Erythema  
Eyelid Oedema  
Folliculitis  
Haemorrhage  
Hypersensitivity  
Hypoventilation  
Irritability  
Oedema

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Pain Palpitations Pruritus	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Purulence Rash Erythematous Rash Papular					
/ ORAL		Respiratory Disorder					
		Scratch		Carisoprodol	C		
		Shock		Diazepam	C		
		Skin Infection					
		Skin Lesion					
		Urticaria					
		Visual Disturbance					

Date:12/04/00ISR Number: 3621705-6Report Type:Expedited (15-DaCompany Report #A0132668A  
Age:34 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG /		Abortion Spontaneous	Study	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY			Haemorrhage	Health Professional				
/ ORAL								

Date:12/04/00ISR Number: 3622221-8Report Type:Expedited (15-DaCompany Report #A0132722A  
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG /		Myocarditis	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE ER DAY								
/ ORAL								

Medroxyprogesterone

INTRAMUSCULAR 150 MG / SEE

Ace. Injection SS

TEXT /

INTRAMUSCULAR

Alfalfa	C
Calcium Salt	C
Activated Charcoal	C
Kelp	C
Potassium Salt	C
Rose Hips	C
Ascorbic Acid	C

Date:12/04/00ISR Number: 3622293-0Report Type:Expedited (15-DaCompany Report #B0092499A  
Age:35 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Blister Eye Disorder Rash Pruritic Visual Acuity Reduced	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:12/04/00ISR Number: 3622294-2Report Type:Expedited (15-DaCompany Report #B0092355A  
Age:60 YR Gender:Male I/FU:I

Outcome	PT	Report Source
Hospitalization - Initial or Prolonged	Toxic Epidermal Necrolysis	Foreign Health

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Professional Company Representative	Product	Role	Manufacturer	Route
ORAL			Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
			Statines	C		

Date:12/04/00ISR Number: 3622295-4Report Type:Expedited (15-DaCompany Report #B0091997A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG / PER DAY / ORAL		Angioneurotic Oedema Bronchial Obstruction Dermatitis Eyelid Oedema Pruritus Rash Generalised Urticaria	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:12/04/00ISR Number: 3622389-3Report Type:Expedited (15-DaCompany Report #WAES 00111289  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other 10 MG	1832 DAY	Abdominal Pain Constipation Diarrhoea Dysphagia Rectal Haemorrhage Urticaria	Foreign Other	Prinivil  Zyban (Bupropion Hcl)  Anafranil Bendroflumethiazide Betahistine Prochlorperazine	PS  SS  C C C C	Merck Research Laboratories Div Merck Co Inc	ORAL  ORAL
150 MG							

Date:12/05/00ISR Number: 3621719-6Report Type:Expedited (15-DaCompany Report #A0086224A  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day				No Concurrent Medications	C		

Date:12/05/00ISR Number: 3621720-2Report Type:Expedited (15-DaCompany Report #A0105888A  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG Per day	2 DAY						

Date:12/05/00ISR Number: 3621725-1Report Type:Expedited (15-DaCompany Report #A0131022A  
Age:30 YR Gender:Female I/FU:F

Outcome PT  
Other Bleeding Time Prolonged  
Cholelithiasis  
Platelet Count Decreased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Post Procedural Haemorrhage	Report Source	Product	Role	Manufacturer	Route
150MG Three times per day				Wellbutrin	PS	Glaxo Wellcome	ORAL
25MG As directed				Lamictal	SS	Glaxo Wellcome	ORAL
				Risperdal	C		
				Paxil	C		
				Ambien	C		
				Celexa	C		
				Sonata	C		

Date:12/05/00ISR Number: 3622806-9Report Type:Direct Company Report #  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eye Pain Headache		Zyban (Bupropion Sr) (150 Mg)	PS		ORAL
150 MG PO QD		Mental Impairment Neck Pain Tremor					

Date:12/06/00ISR Number: 3622603-4Report Type:Expedited (15-DaCompany Report #A0133264A  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Deafness Tinnitus		Wellbutrin Wellbutrin Sr	PS SS	Glaxo Wellcome Glaxo Wellcome	ORAL
Other							

Date:12/06/00ISR Number: 3622626-5Report Type:Expedited (15-DaCompany Report #B0093039A  
 Age:68 YR Gender:Male I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice		Myocardial Infarction	Zyban	PS	Glaxo Wellcome	ORAL
	per day		Sudden Death				
	75MG Per day			Aspirin	C		ORAL
	150MG Per day			Ranitidine	C	Glaxo Wellcome	ORAL

Date:12/06/00ISR Number: 3622627-7Report Type:Expedited (15-DaCompany Report #B0093101A  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	7 DAY	Death	Zyban	PS	Glaxo Wellcome	ORAL
				Clopidogrel	C		ORAL
				Amitriptyline	C		ORAL
				Ipratropium	C	Glaxo Wellcome	ORAL

Date:12/06/00ISR Number: 3622628-9Report Type:Expedited (15-DaCompany Report #B0093206A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide	Zyban	PS	Glaxo Wellcome	ORAL



Other	Depression	Foreign	Bupropion	PS	Glaxo Wellcome Inc	ORAL
1 TABLET	Headache	Health	Hydrochloride			
(TWICE PER	Oedema Peripheral	Professional				
DAY) ORAL	Physical Assault					
	Suicide Attempt					

Date:12/06/00ISR Number: 3623680-7Report Type:Expedited (15-DaCompany Report #A0132705A  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Foreign	Bupropion			
Other		Cellulitis	Consumer	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE		Dyspnoea					
PER DAY) ORAL		Erythema					
		Feeling Hot					
		Hypersensitivity					
		Joint Swelling					
		Oedema Peripheral					
		Pain In Extremity					
		Swelling					
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/06/00ISR Number: 3623682-0Report Type:Expedited (15-DaCompany Report #B0092315A

Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Haemorrhagic Stroke	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL		Nausea	Professional				

Date:12/06/00ISR Number: 3623715-1Report Type:Expedited (15-DaCompany Report #B0092839A

Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to prevent Permanent Impairment/Damage		Psoriasis	Foreign Other	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Calcipotriene	C		
				Betamethasone	C		
				Salbutamol Sulphate	C		
				Beclomethasone			
				Dipropion	C		
				Meloxicam	C		

Date:12/06/00ISR Number: 3623716-3Report Type:Expedited (15-DaCompany Report #B0092451A

Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hyperreflexia Paraesthesia	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL		Tachycardia	Professional				
		Tremor		Acyclovir	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day							
		International Normalised		Warfarin	I	Glaxo Wellcome	ORAL
5MG Per day		Ratio Increased					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Wellbutrin Sr	PS		ORAL
100 MG BID PO							
		Anger		Tenex	SS		ORAL
0.5 MG Q AM							
		Epilepsy					
PO							
		Headache		Risperdal	SS		ORAL
1 MG Q HS PO							
		Loss Of Consciousness		Risperdal	SS		ORAL
1 MG BID PO							
		Memory Impairment		Adderal	C		
		Vision Blurred		Flonase	C		
				Haldol Im	C		
				Haldol	C		
				Tylenol	C		
				Vistaril Im	C		
				Vistaril	C		
				Maalox	C		
				Mom	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/08/00ISR Number: 3624600-1Report Type:Expedited (15-DaCompany Report #A0111797A  
 Age:67 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Per day	Confusional State		Bupropion	PS	Glaxo Wellcome	ORAL
Initial or Prolonged 1TAB Per day	Dizziness		Placebo	SS		ORAL
75MG Per day	Dysarthria		Plavix	C		ORAL
25MG Per day	Hypotension		Atenolol	C		ORAL
2.5MG Per day	Syncope		Zestril	C		ORAL
20MG Twice per day	Ventricular Tachycardia		Prilosec	C		ORAL
			Enteric-Coated Aspirin	C		ORAL
			Lipitor	C		ORAL
SUBLINGUAL			Nitroglycerin 1/150	C	Glaxo Wellcome	

Date:12/08/00ISR Number: 3624618-9Report Type:Expedited (15-DaCompany Report #B0089165A  
 Age:26 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150MG Twice per day	Abortion Spontaneous Pregnancy	Health Professional	Zyban	PS	Glaxo Wellcome	ORAL
	9 DAY					

Date:12/08/00ISR Number: 3624620-7Report Type:Expedited (15-DaCompany Report #B0090959A  
 Age:26 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other	Abortion Spontaneous	Health Professional	Zyban	PS	Glaxo Wellcome	

Date:12/08/00ISR Number: 3624621-9Report Type:Expedited (15-DaCompany Report #B0092066A  
Age:38 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Dizziness		Zyban	PS	Glaxo Wellcome	ORAL
150MG As						
Hospitalization -	Hyperhidrosis					
directed 7 DAY						
Initial or Prolonged	Malignant Hypertension					
	Ventricular Hypertrophy					
	Visual Disturbance					

Date:12/08/00ISR Number: 3624629-3Report Type:Expedited (15-DaCompany Report #B0093404A  
Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Haemoptysis		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged	International Normalised Ratio Increased					

Date:12/08/00ISR Number: 3625343-0Report Type:Expedited (15-DaCompany Report #HQ2762525OCT2000  
Age:67 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Agitation
Initial or Prolonged	Asthenia
	Disorientation
	Hallucination
	Headache

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
175 MG, 450 MG, 75 MG, DAILY, ORAL		Hyperhidrosis Hypoaesthesia Lethargy Paraesthesia Thinking Abnormal	Consumer	Effexor Xr	PS	Wyeth Ayerst Laboratories	ORAL
ORAL				Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL
				Synthroid (Levothyroxine Sodium)	C		

Date:12/08/00ISR Number: 3625451-4Report Type:Expedited (15-DaCompany Report #A0131022A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG / THREE TIMES PER DAY / ORAL		Bleeding Time Prolonged Cholelithiasis Platelet Count Decreased Post Procedural Haemorrhage	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
25 MG / ORAL/ AS DIRECTED				Lamictal Tablet (Lamotrigine)	SS		ORAL
				Risperidone	C		
				Paroxetine Hydrochloride	C		
				Zolpidem Tartrate	C		
				Citalopram Hydrobromide	C		
				Zaleplon	C		



Date:12/08/00ISR Number: 3625616-1Report Type:Expedited (15-DaCompany Report #A0086224A  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Overdose	Literature				
PER DAY/ ORAL			Health Professional				

Date:12/08/00ISR Number: 3625617-3Report Type:Expedited (15-DaCompany Report #A0105888A  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							
DAY/ ORAL	2	DAY					

Date:12/08/00ISR Number: 3625620-3Report Type:Expedited (15-DaCompany Report #B0093064A  
Age:24 YR Gender:Female I/FU:I

Outcome  
Required  
Intervention to  
Prevent Permanent

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Vision Blurred	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Insulin Lispro Human Int/Long Insulin	C C		

Date:12/08/00ISR Number: 3625621-5Report Type:Expedited (15-DaCompany Report #B0093067A  
Age:67 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 150 MG/ PER Intervention to DAY/ ORAL Prevent Permanent Impairment/Damage		Rectal Haemorrhage	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Co-Codamol Ibuprofen	C C		

Date:12/08/00ISR Number: 3625622-7Report Type:Expedited (15-DaCompany Report #B0093052A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG / UNK, ORAL		Hypersensitivity	Foreign Other	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Fluoxetine Chlorhexidine	C C		

Date:12/08/00ISR Number: 3625623-9Report Type:Expedited (15-DaCompany Report #B0093061A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Diplopia	Foreign	Bupropion			
		Dry Mouth	Consumer	Hydrochloride	PS	Glaxo Wellcome Inc	
UNKNOWN	150 MG/	UNK/ Insomnia					
UNKNOWN		Tinnitus Vision Blurred Visual Disturbance					

Date:12/08/00ISR Number: 3625624-0Report Type:Expedited (15-DaCompany Report #B0092443A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abnormal Behaviour	Foreign	Zyban	PS	Glaxo Wellcome Inc	
UNKNOWN	UNK/UNK/						
Initial or Prolonged		Disinhibition	Health				
UNKNOWN		Drug Interaction Intentional Self-Injury	Professional Other	Fluoxetine (Formulation Unknown) (Fluoxetine)	SS		
UNKNOWN	UNK/ UNK/						
UNKNOWN							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/11/00ISR Number: 3625422-8Report Type:Expedited (15-DaCompany Report #B0093505A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability			Blood Creatine	Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice Other per day	63	DAY	Phosphokinase Increased				
150MG Per day			Chest Pain	Cetirizine	C	Glaxo Wellcome	
200MG Twice per day			Convulsion	Omeprazole	C		
			Gamma-Glutamyltransferase				
			Increased Insomnia	Fluticasone Salbutamol	C C	Glaxo Wellcome Glaxo Wellcome	NASAL
RESPIRATORY (INHALATION)			Lethargy				
RESPIRATORY (INHALATION)			Muscle Spasms	Beclomethasone	C	Glaxo Wellcome	

Date:12/11/00ISR Number: 3625423-XReport Type:Expedited (15-DaCompany Report #B0093507A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Initial or Prolonged Variable dose	12	DAY	Psoriasis	Zyban	PS	Glaxo Wellcome	ORAL

Date:12/11/00ISR Number: 3625424-1Report Type:Expedited (15-DaCompany Report #B0093510A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Per day	17	DAY	Periorbital Oedema Stridor	Zyban	PS	Glaxo Wellcome	ORAL

Urticaria

Date:12/11/00ISR Number: 3626713-7Report Type:Expedited (15-DaCompany Report #A0133264A  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness Tinnitus	Consumer	Wellbutrin Wellbutrin	PS SS	Glaxo Wellcome Inc	ORAL
ORAL							

Date:12/11/00ISR Number: 3627087-8Report Type:Expedited (15-DaCompany Report #B0093206A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL							
			Professional Company Representative				

Date:12/11/00ISR Number: 3627089-1Report Type:Expedited (15-DaCompany Report #B0093033A  
 Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction International Normalised Ratio Increased	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG PER DAY, ORAL							
				Warfarin Sodium	SS		ORAL
5 MG (PER DAY) ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/11/00ISR Number: 3627108-2Report Type:Expedited (15-DaCompany Report #B0093101A  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER DAY/ORAL				Clopidogrel	C		
				Amitriptyline	C		
				Ipratropium Bromide	C		

Date:12/11/00ISR Number: 3627343-3Report Type:Expedited (15-DaCompany Report #B0093039A  
Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Foreign Other	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY / ORAL				Aspirin	C		
				Ranitidine Hydrochloride	C		

Date:12/12/00ISR Number: 3626592-8Report Type:Expedited (15-DaCompany Report #A0109223A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 39 DAY		Arthralgia	Health	Zyban	PS	Glaxo Wellcome	ORAL
		Back Pain	Professional	Ibuprofen	C		
		Bone Density Decreased					
		Intervertebral Disc Degeneration					
		Leukocytoclastic Vasculitis					

Osteoporosis  
Pain  
Pain In Extremity  
Pruritus  
Purpura  
Rash Erythematous  
Rheumatoid Arthritis  
Sneezing  
Throat Irritation  
Urticaria  
White Blood Cell Count  
Increased

Date:12/12/00ISR Number: 3627261-0Report Type:Direct  
Age:7 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Grand Mal Convulsion		Bupropion	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/12/00ISR Number: 3627615-2Report Type:Expedited (15-DaCompany Report #A0111797A  
Age:67 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG (PER DAY) ORAL	Dehydration Myocardial Ischaemia Syncope Ventricular Tachycardia	Foreign Health Professional	Bupropion Hydrochloride  Placebo Tablet (Placebo)	PS  SS	Glaxo Wellcome Inc	ORAL  ORAL
1 TABLET (PER DAY) ORAL			Clopidogrel Bisulphate Atenolol Lisinopril Omeprazole Aspirin Atorvastatin Calcium Nitroglycerin	C C C C C C C		

Date:12/12/00ISR Number: 3627687-5Report Type:Expedited (15-DaCompany Report #B0093404A  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG	Haemoptysis International Normalised Ratio Increased	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:12/12/00ISR Number: 3627806-0Report Type:Expedited (15-DaCompany Report #B0092066A  
Age:38 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 150 MG/ AS	Malignant Hypertension Ventricular Hypertrophy	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL



Initial or Prolonged  
DIRECTED/

Professional

ORAL

Date:12/12/00ISR Number: 3627808-4Report Type:Expedited (15-DaCompany Report #B0090959A  
Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:12/12/00ISR Number: 3627809-6Report Type:Expedited (15-DaCompany Report #B0089165A  
Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/13/00ISR Number: 3627847-3Report Type:Expedited (15-DaCompany Report #B0091833A  
Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Twice	Angioneurotic Oedema	Health	Zyban	PS	Glaxo Wellcome	ORAL
per day	16 DAY	Asthma	Professional				
		Urticaria					

Date:12/13/00ISR Number: 3627861-8Report Type:Expedited (15-DaCompany Report #D0010662A  
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	450MG per day 10 DAY	Dyspnoea	Health	Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -	Initial or Prolonged	Grand Mal Convulsion	Professional				
		Pulse Absent					

Date:12/13/00ISR Number: 3627862-XReport Type:Expedited (15-DaCompany Report #D0011022A  
Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	4 WK	Angina Pectoris	Health	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	1 YR	Catecholamines Urine	Professional	Oral Contraceptives	C		
		Increased					
		Epistaxis					
		Headache					
		Hypertension					

Date:12/13/00ISR Number: 3628961-9Report Type:Expedited (15-DaCompany Report #B0093510A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Periorbital Oedema	Foreign	Bupropion			

150 MG /PER Stridor Other Hydrochloride PS Glaxo Wellcome Inc ORAL  
 Urticaria  
 DAY/ORAL

Date:12/13/00ISR Number: 3628962-0Report Type:Expedited (15-DaCompany Report #B0093507A  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/ VARIABLE		Psoriasis	Foreign Health Professional Other	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
DOSE/ORAL							

Date:12/13/00ISR Number: 3628986-3Report Type:Expedited (15-DaCompany Report #B0093505A  
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required 150 MG/ TWICE Intervention to PER DAY/ Prevent Permanent ORAL Impairment/Damage		Blood Creatine Phosphokinase Increased Chest Pain Convulsion Gamma-Glutamyltransferase Increased Insomnia Lethargy Muscle Spasms	Foreign	Bupropion Hydrochloride Cetirizine Hydrochloride Omeprazole Fluticasone Propionate	PS C C C	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Salbutamol Sulphate C  
 Beclomethasone  
 Dipropion C

Date:12/14/00ISR Number: 3628923-1Report Type:Direct  
 Age:66 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG BID Initial or Prolonged ORAL		Urticaria		Bupropion 150mg Sa	PS		ORAL
				Asa	C		
				Atenolol	C		
				Ipratropium	C		
				Isdn	C		
				Niacin	C		
				Nicotine Patch	C		
				Simvastatin	C		

Date:12/14/00ISR Number: 3628924-3Report Type:Expedited (15-DaCompany Report #A0133478A  
 Age:17 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 38TAB Single Initial or Prolonged dose		Coma		Wellbutrin	PS	Glaxo Wellcome	ORAL
54TAB Single dose		Ileus Paralytic		Tegretol	SS		ORAL
12TAB Single dose		Overdose					
38TAB Single dose		Oxygen Saturation Decreased		Trazodone	SS		ORAL
				Seroquel	SS		ORAL

dose

Date:12/14/00ISR Number: 3628931-0Report Type:Expedited (15-DaCompany Report #B0089929A  
 Age:67 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Circulatory Collapse	Health	Zyban	PS	Glaxo Wellcome	
150MG Twice		Depressed Level Of	Professional				
per day	51	DAY		Cimetidine	C		ORAL
400MG Twice		Dyspepsia					
per day	6	DAY		Temazepam	C		ORAL
		Dyspnoea					
		Dysuria					
		Eye Irritation					
		Grand Mal Convulsion					
		Keratoconjunctivitis					
		Sicca					
		Myocardial Infarction					

Date:12/14/00ISR Number: 3628977-2Report Type:Direct Company Report #  
 Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Diarrhoea		Bupropion	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/15/00ISR Number: 3630326-0Report Type:Expedited (15-DaCompany Report #A0134365A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphasia		Zyban	PS	Glaxo Wellcome	ORAL
1 MON		Cystitis Dysarthria		No Concurrent Medication	C		

Date:12/15/00ISR Number: 3630335-1Report Type:Expedited (15-DaCompany Report #B0092251A

Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eyelid Disorder		Zyban	PS	Glaxo Wellcome	ORAL
30 DAY		Nausea Pain Retinal Disorder Retinal Vascular Disorder Vision Blurred Visual Acuity Reduced		Oestradiol	C		

Date:12/15/00ISR Number: 3630342-9Report Type:Expedited (15-DaCompany Report #B0093973A

Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Arthropathy		Zyban	PS	Glaxo Wellcome	
Other		Stevens-Johnson Syndrome					

Date:12/15/00ISR Number: 3630343-0Report Type:Expedited (15-DaCompany Report #B0093986A

Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Burning Sensation		Zyban	PS	Glaxo Wellcome	
150MG		Pruritus					
Variable dose	6 DAY						

Date:12/18/00ISR Number: 3631624-7Report Type:Expedited (15-DaCompany Report #A0132705A  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability							
150MG Twice				Zyban	PS	Glaxo Wellcome	ORAL
Other							
per day	24 DAY	Cellulitis					
		Dyspnoea					
		Feeling Hot					
		Hypersensitivity					
		Pain In Extremity					
		Rash Erythematous					
		Stevens-Johnson Syndrome					
		Swelling					
		Urticaria					

Date:12/18/00ISR Number: 3631633-8Report Type:Expedited (15-DaCompany Report #B0093969A  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening							
Other				Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY				Salbutamol	C	Glaxo Wellcome	
(INHALATION)	2PUFF As	Urticaria					

required

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Freedom Of Information (FOI) Report

Antihistamine C

Date:12/18/00ISR Number: 3631634-XReport Type:Expedited (15-DaCompany Report #B0093975A  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	23 DAY	Abortion Spontaneous Vaginal Haemorrhage		Zyban	PS	Glaxo Wellcome	ORAL
1TAB Per day				Microgynon	C		ORAL
VAGINAL				Sultrin	C		

Date:12/18/00ISR Number: 3631635-1Report Type:Expedited (15-DaCompany Report #B0093979A  
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 6 DAY Hospitalization - Initial or Prolonged		Akinesia Dyspnoea		Zyban Atenolol + Chlorthalidone	PS C	Glaxo Wellcome	ORAL ORAL

Date:12/18/00ISR Number: 3633253-8Report Type:Expedited (15-DaCompany Report #B0089929A  
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG /TWICE PER DAY		Circulatory Collapse Dyspepsia Dyspnoea	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
		Grand Mal Convulsion Keratoconjunctivitis Sicca Loss Of Consciousness Myocardial Infarction		Cimetidine Temazepam	C C		



Urinary Tract Disorder

Date:12/18/00ISR Number: 3633257-5Report Type:Expedited (15-DaCompany Report #B0091833A  
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Angioneurotic Oedema Asthma	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE PER DAY/ ORAL		Urticaria	Professional				

Date:12/18/00ISR Number: 3633289-7Report Type:Expedited (15-DaCompany Report #A0133478A  
 Age:17 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 38 TABLET/ Initial or Prolonged SINGLE DOSE/ ORAL		Coma Dialysis Ileus Paralytic	Health Professional	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
54 TABLET/ SINGLE DOSE/ ORAL		Overdose Oxygen Saturation Decreased Sedation		Carbamazepine Tablet (Carbamazepine)	SS		ORAL
12 TABLET/ SINGLE DOSE/ ORAL				Trazodone Tablet (Trazodone)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

38	Quetiapine Fumarate Tablet (Quetiapine Fumarate)	SS	ORAL
TABLET/SINGLE			
DOSE/ ORAL			
1600 MG/	Ibuprofen Tablet (Ibuprofen)	SS	ORAL
SINGLE DOSE/			
ORAL			

Date:12/18/00 ISR Number: 3633358-1 Report Type:Expedited (15-DaCompany Report #D0011022A  
 Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Angina Pectoris Blood Catecholamines	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Abnormal Epistaxis Headache Pulmonary Arterial Wedge Pressure Increased	Professional	Oral Contraceptive	C		

Date:12/18/00 ISR Number: 3633359-3 Report Type:Expedited (15-DaCompany Report #D0010662A  
 Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - ORAL	10 DAY	Convulsion Dyspnoea	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Pulse Absent	Professional				

Date:12/18/00ISR Number: 3633571-3Report Type:Direct  
Age:32 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tremor		Wellbutrin Sr 150mg	PS		ORAL
150MG ONCE A							
DAY ORAL							

Date:12/19/00ISR Number: 3632879-5Report Type:Expedited (15-DaCompany Report #A0055937A  
Age:55 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Accelerated Hypertension
Initial or Prolonged	Alopecia
Disability	Anaphylactic Shock
	Anuria
	Anxiety
	Asthenia
	Autonomic Nervous System
	Imbalance
	Back Pain
	Bipolar Disorder
	Bradycardia
	Chills
	Cognitive Disorder
	Convulsion
	Cystocele
	Depersonalisation

Freedom Of Information (FOI) Report

Depression  
Disorientation  
Dizziness  
Drug Hypersensitivity  
Dysarthria  
Dyspepsia  
Dyspnoea  
Dysthymic Disorder  
Fatigue  
Fear  
Feeling Abnormal  
Haematuria  
Headache  
Heart Rate Abnormal  
Hyperglycaemia  
Hyperhidrosis  
Hypertension  
Hyperventilation  
Hypoaesthesia  
Hypomania  
Intervertebral Disc  
Degeneration  
Labile Blood Pressure  
Memory Impairment  
Mental Impairment  
Micturition Urgency  
Mitral Valve Incompetence  
Movement Disorder  
Muscle Disorder  
Muscle Rigidity  
Muscle Spasms  
Muscle Twitching  
Muscular Weakness  
Musculoskeletal Stiffness  
Neck Pain  
Nervousness  
Neurosis  
Obsessive-Compulsive  
Disorder  
Oedema  
Palpitations  
Panic Disorder  
Paraesthesia  
Renal Disorder  
Renal Failure Chronic  
Respiratory Disorder  
Schizophrenia

Sciatica  
Stupor  
Swelling  
Tachycardia  
Tangentiality  
Throat Tightness  
Tinnitus  
Tongue Oedema  
Transient Ischaemic  
Attack  
Tremor  
Urethral Stricture  
Urinary Incontinence  
Ventricular Extrasystoles

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Weight Increased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	12 DAY		Zyban	PS	Glaxo Wellcome	ORAL
1.25MG Per day			Premarin	C		
			Vitamin	C		
			Asa	C		
			Melatonin	C		

Date:12/19/00ISR Number: 3632904-1Report Type:Expedited (15-DaCompany Report #B0082488A  
Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Macular Degeneration Retinal Exudates		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG Twice Other per day	31 DAY	Retinal Oedema Retinal Vasculitis Retinitis					

Date:12/19/00ISR Number: 3632907-7Report Type:Expedited (15-DaCompany Report #B0092236A  
Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Twice per day	18 DAY	Astigmatism Eye Movement Disorder		Zyban	PS	Glaxo Wellcome	ORAL
4PUFF Twice per day		Hypermetropia Hypertension		Salbutamol	C	Glaxo Wellcome	
4PUFF Twice		Vision Blurred Visual Disturbance		Beclomethasone Dipropionate	C	Glaxo Wellcome	

per day

Frusemide C

30MG Per day

Spirolactone C

100MG Per day

Digoxin C Glaxo Wellcome

.25MG Per day

Date:12/19/00ISR Number: 3632909-0Report Type:Expedited (15-DaCompany Report #B0092251A

Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	30 DAY	Eye Pain	Health	Zyban	PS	Glaxo Wellcome	ORAL
Other		Nausea Retinoschisis	Professional	Oestradiol	C		

Date:12/19/00ISR Number: 3632911-9Report Type:Expedited (15-DaCompany Report #B0093187A

Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day 3 DAY	Agitation		Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY (INHALATION)		Delirium Tremens Hallucination Suicidal Ideation Trismus Visual Disturbance		Salmeterol Alprazolam	C	Glaxo Wellcome	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/19/00ISR Number: 3632925-9Report Type:Expedited (15-DaCompany Report #B0093977A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day	16 DAY	Asthma	Zyban	PS	Glaxo Wellcome	ORAL
Other	1PUFF Three		Dyspnoea	Salbutamol	C	Glaxo Wellcome	
	times per day			Eformoterol	C		
	1PUFF Three						
	times per day			Steroids	C		

Date:12/19/00ISR Number: 3632926-0Report Type:Expedited (15-DaCompany Report #B0094080A  
Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Twice		Atrial Fibrillation	Zyban	PS	Glaxo Wellcome	ORAL
	per day	14 DAY	Gait Disturbance				
	50MG per day		Hypoaesthesia Oral	Atenolol	C		ORAL
	2.5MG per day			Bendrofluazide	C	Glaxo Wellcome	ORAL
				Nicotine	C		

Date:12/19/00ISR Number: 3632927-2Report Type:Expedited (15-DaCompany Report #B0094092A  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice		Coronary Artery Disease	Zyban	PS	Glaxo Wellcome	ORAL
	Hospitalization -						
	per day	51 DAY		Co-Dydramol	C	Glaxo Wellcome	ORAL
	Initial or Prolonged						



Date:12/19/00ISR Number: 3632929-6Report Type:Expedited (15-DaCompany Report #D0011104A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 1TAB Twice		Arthralgia	Health	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	2 WK	Dyspnoea	Professional				
		Haematuria					
		Hypersensitivity					
		Myalgia					
		Urticaria					
		White Blood Cells Urine Positive					

Date:12/19/00ISR Number: 3633679-2Report Type:Expedited (15-DaCompany Report #A0134365A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphasia	Foreign	Bupropion			
		Cystitis	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL		Dysarthria	Professional Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/19/00ISR Number: 3633681-0Report Type:Expedited (15-DaCompany Report #B0093973A  
Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required 150 MG Intervention to Prevent Permanent Impairment/Damage		Arthropathy Stevens-Johnson Syndrome	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:12/19/00ISR Number: 3633688-3Report Type:Expedited (15-DaCompany Report #B0092251A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG ORAL Prevent Permanent Impairment/Damage		Eye Pain Nausea Retinal Disorder Visual Disturbance	Foreign Health Professional	Bupropion Hydrochloride Oestradiol	PS  C	Glaxo Wellcome Inc	ORAL

Date:12/19/00ISR Number: 3633690-1Report Type:Expedited (15-DaCompany Report #B0093986A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG Prevent Permanent VARIABLE DOSE Impairment/Damage		Burning Sensation Pruritus	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:12/19/00ISR Number: 3633960-7Report Type:Periodic Company Report #0323283A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Effect Decreased	Consumer	Habitrol	PS	Novartis Consumer	

1

PATCH/QD/TTS

Wellbutrin-Bupropion  
Hcl

SS

Glaxo Wellcome

1

PATCH/QD/TTS

Date:12/19/00ISR Number: 3633996-6Report Type:Expedited (15-DaCompany Report #HQ2762525OCT2000

Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 175 MG DAILY, ORAL		Agitation Asthenia Disorientation	Health Professional	Effexor Xr	PS	Wyeth Ayerst Laboratories	ORAL
ORAL; SEE IMAGE		Hallucination Headache Hyperhidrosis Hypoaesthesia Lethargy Panic Attack		Wellbutrin (Amfebutamone Hydrochloride, Capsule, Extended Release)	SS		ORAL
		Paraesthesia Syncope Thinking Abnormal		Synthroid (Levothyroxine Sodium)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/19/00ISR Number: 3634102-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #0303996A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Application Site Reaction	Consumer	Habitrol	PS	Novartis Consumer Health Inc	
1		Pruritus					
		Sedation					
PATCH/QD/TTS							
				Zyban - Bupropion Hcl 150mg - Glaxo	SS	Glaxo	ORAL
150 MG/PO							

Date:12/19/00ISR Number: 3634166-8Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #0269684A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams	Health Professional	Habitrol	PS	Novartis Consumer Health Inc	
		Insomnia					
TRANSDERMAL	1						
PATCH/QD/TTS							
				Zyban-Bupropion Hcl 150 Mg-Glaxo	SS		ORAL
PO QD							
				Nasonex	C		
				Zyrtec	C		
				Depo Provera	C		

Date:12/19/00ISR Number: 3634175-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #0272578A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea	Consumer	Habitrol	PS	Novartis Consumer Health Inc	
		Dizziness					
1 PATCH /TTS							
		Nausea					
				Zyban-Bupropion Hcl-Glaxo	SS		ORAL
PO							
				Zestoretic	C		

Date:12/19/00ISR Number: 3634235-2Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #0272281A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pruritus Rash Erythematous	Consumer	Habitrol	PS	Novartis Consumer Health Inc	
1							
PATCH/QD/TTS		Urticaria					
				Wellbutrin-Bupropion Hcl Tab-Glaxo Wellcome	SS	Glaxo Wellcome	ORAL
PO							

Date:12/19/00ISR Number: 3634266-2Report Type:Periodic  
Age:64 YR Gender:Male I/FU:I

Company Report #0309310A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Oedema Pruritus	Consumer	Habitrol	PS	Novartis Consumer Health Inc	
TRANSDERMAL	2						
PATCHES/QD/TT							
S							
150 MG OD PO,				Zyban-Bupropion-Hcl 150 Mg	SS	Glaxo	ORAL
THEN 300				Synthroid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/19/00ISR Number: 3634309-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #0315170A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State	Consumer	Habitrol	PS	Novartis Consumer Health Inc	
TRANSDERMAL	1 PATCH/						
QD/TTS							
				Wellbutrin - Bupropion Hcl - Glaxo Wellcome	SS	Glaxo Wellcome	

Date:12/19/00ISR Number: 3634744-6Report Type:Periodic  
 Age:50 YR Gender:Female I/FU:I

Company Report #0266794A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Oedema Mouth Pharyngeal Oedema	Consumer	Habitrol	PS	Novartis Consumer Health Inc	
TRANSDERMAL	1 PATCH/ QD/						
TTS		Urticaria					
QD PO				Zyban (Glaxo Wellcome )	SS	Glaxo Wellcome	ORAL
				Zyban Synthroid	C		

Date:12/19/00ISR Number: 3634804-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #0268640A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache Nervousness	Consumer	Habitrol	PS	Novartis Consumer Health Inc	
TRANSDERMAL	TTS	15 DAY					
PO				Zyban - Bupropion Hcl 150 Mg - Glaxo	SS	Glaxo	ORAL

Date:12/19/00ISR Number: 3634812-9Report Type:Periodic Company Report #0268835A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Application Site Reaction	Consumer	Habitrol	PS	Novartis Consumer Health Inc	
1		Oedema					
		Paraesthesia					
PATCH/QD/TTS		Weight Increased		Zyban-Bupropion Hcl 150 Mg-Glaxo	SS	Glaxo	ORAL
PO							

Date:12/19/00ISR Number: 3634886-5Report Type:Periodic Company Report #0296321A  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Consumer	Habitrol	PS	Novartis Consumer Health Inc	
1		Headache					
		Hyperhidrosis					
PATCH/QD/TTS				Zyban-Bupropion Hcl 150 Mg - Glaxo	SS		ORAL
150 MG BID PO							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/20/00ISR Number: 3633819-5Report Type:Direct  
Age:45 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1 TAB ORAL Initial or Prolonged		Urticaria		Wellbutrin Sr 150 Mg	PS		ORAL
				Paxil	C		
				Lithium	C		
				Zyprexa	C		

Date:12/20/00ISR Number: 3633822-5Report Type:Expedited (15-DaCompany Report #A0121104A  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 1 WK		Myocardial Infarction		Zyban	PS	Glaxo Wellcome	ORAL
				Bactrim	C	Glaxo Wellcome	
				Viagra	I		ORAL
1 DAY							

Date:12/20/00ISR Number: 3633824-9Report Type:Expedited (15-DaCompany Report #A0128274A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day	18 DAY	Anxiety		Zyban	PS	Glaxo Wellcome	ORAL
		Feeling Abnormal					
11 DAY		Feeling Hot		Nicoderm	C		
		Hyperaesthesia					
		Irritability					
		Nervousness					
		Nightmare					
		Oedema					
		Pruritus					
		Skin Discolouration					
		Urticaria					



Date:12/20/00ISR Number: 3633839-0Report Type:Expedited (15-DaCompany Report #B0093439A  
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 7 DAY	Bronchospasm		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged	Dizziness Dyspnoea Tremor		Ipratropium	C	Glaxo Wellcome	

Date:12/20/00ISR Number: 3633845-6Report Type:Expedited (15-DaCompany Report #B0094062A  
Age:40 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability 300MG Per day 20 DAY	Decreased Activity		Zyban	PS	Glaxo Wellcome	ORAL
	Urticaria					

Date:12/20/00ISR Number: 3633846-8Report Type:Expedited (15-DaCompany Report #B0094077A  
Age:55 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice Initial or Prolonged per day	Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
75MG Per day			Aspirin	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/20/00ISR Number: 3633847-XReport Type:Expedited (15-DaCompany Report #B0094079A

Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Confusional State		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Other		Dry Mouth					
per day	14 DAY	Malaise		Propranolol	C		
		Migraine					
		Nausea					
		Pain					

Date:12/20/00ISR Number: 3633849-3Report Type:Expedited (15-DaCompany Report #B0094291A

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Psoriasis		Zyban	PS	Glaxo Wellcome	ORAL
3 WK							
Initial or Prolonged							

Date:12/20/00ISR Number: 3635022-1Report Type:Expedited (15-DaCompany Report #B0093187A

Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation	Foreign	Bupropion			
150 MG/PER		Delirium Tremens	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Hallucination	Professional				
DAY/ORAL		Suicidal Ideation	Company	Salmeterol	C		
		Trismus	Representative	Alprazolam	C		
		Visual Disturbance					

Date:12/20/00ISR Number: 3635023-3Report Type:Expedited (15-DaCompany Report #B0094080A

Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Atrial Fibrillation	Foreign	Bupropion			
Intervention to		Gait Disturbance		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
Prevent Permanent		Hypoaesthesia Oral					
PER DAY/'							
Impairment/Damage							
ORAL							
				Atenolol	C		
				Bendrofluazide	C		
				Nicotine	C		

Date:12/20/00ISR Number: 3635024-5Report Type:Expedited (15-DaCompany Report #B0093977A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Asthma	Foreign	Bupropion			
Required				Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER							
Intervention to							
DAY/ ORAL							
Prevent Permanent				Salbutamol Sulphate	C		
Impairment/Damage				Eformoterol	C		
				Steroid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/20/00ISR Number: 3635025-7Report Type:Expedited (15-DaCompany Report #B0094092A  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coronary Artery Disease	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL				Co-Dydramol	C		

Date:12/20/00ISR Number: 3635026-9Report Type:Expedited (15-DaCompany Report #B0092236A  
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Astigmatism Eye Movement Disorder	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE PER DAY/ORAL		Hypermetropia	Professional	Salbutamol Sulphate	C		
		Hypertension Vision Blurred		Beclomethasone	C		
				Dipropion	C		
				Frusemide	C		
				Spirolactone	C		
				Digoxin	C		

Date:12/20/00ISR Number: 3635033-6Report Type:Expedited (15-DaCompany Report #B0092251A  
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required 150 MG / ORAL		Nausea Retinal Detachment	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Intervention to Prevent Permanent Impairment/Damage			Professional	Oestradiol	C		

Date:12/20/00ISR Number: 3635034-8Report Type:Expedited (15-DaCompany Report #D0011104A  
Age:38 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1	Arthralgia Haematuria Hypersensitivity Myalgia White Blood Cells Urine Positive	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TABLET/TWICE PER DAY/ORAL						

Date:12/20/00ISR Number: 3635035-XReport Type:Expedited (15-DaCompany Report #B0082488A  
Age:32 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG / Required TWICE PER DAY Intervention to / ORAL Prevent Permanent Impairment/Damage	Eye Irritation Maculopathy Retinal Oedema Retinal Tear Retinitis	Foreign Study Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Freedom Of Information (FOI) Report

Date:12/20/00  
 ISR Number: 3635126-3  
 Report Type:Expedited (15-DaCompany Report #A0055937A  
 Age:55 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG / PER Disability DAY / ORAL	Alopecia Anxiety  Blood Glucose Increased  Blood Triglycerides Increased Bradycardia Cognitive Disorder Confusional State Cystocele Diplopia Dizziness Drug Hypersensitivity Dry Mouth Dry Throat Dysarthria Dyspepsia Dysthymic Disorder Dysuria Enuresis Faeces Pale Feeling Abnormal Feeling Hot And Cold Fluid Retention Headache Hypertension Hypoaesthesia Insomnia Intervertebral Disc Protrusion Memory Impairment Mental Impairment Movement Disorder Muscle Spasms Muscular Weakness Musculoskeletal Stiffness Nausea Nervousness Oedema Palpitations Panic Disorder	Health Professional	Bupropion Hydrochloride   Conjugated Estrogens Vitamin Aspirin Melatonin		Glaxo Wellcome Inc	ORAL
				PS		
				C		
				C		
				C		
				C		

Paraesthesia  
Sciatica  
Spinal Osteoarthritis  
Syncope  
Throat Tightness  
Tinnitus  
Tremor

Date:12/21/00ISR Number: 3634583-6Report Type:Expedited (15-DaCompany Report #A0134853A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cervix Carcinoma		Wellbutrin	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged		Drug Toxicity Overdose					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/21/00ISR Number: 3634599-XReport Type:Expedited (15-DaCompany Report #B0092443A

Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	47 DAY	Amputation		Zyban	PS	Glaxo Wellcome	
75MG Per day		Disinhibition		Aspirin	C		ORAL
2.5MG Per day		Intentional Self-Injury		Bendrofluazide	C	Glaxo Wellcome	ORAL
10MG Per day				Fosinopril	C		ORAL
				Fluoxetine	I		

Date:12/21/00ISR Number: 3634623-4Report Type:Expedited (15-DaCompany Report #B0094547A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Twice per day		Completed Suicide		Zyban	PS	Glaxo Wellcome	ORAL

Date:12/21/00ISR Number: 3634624-6Report Type:Expedited (15-DaCompany Report #B0094550A

Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Single dose	1 DAY	Decreased Appetite		Zyban	PS	Glaxo Wellcome	ORAL
1U Per day		Panic Attack		Cilest	C		ORAL
		Psychomotor Hyperactivity					
		Thirst					
		Vision Blurred					



Date:12/21/00ISR Number: 3634625-8Report Type:Expedited (15-DaCompany Report #B0094552A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain		Zyban	PS	Glaxo Wellcome	ORAL
150MG		Dry Mouth					
Variable dose	3	DAY					
		Insomnia					
		Vision Blurred					

Date:12/21/00ISR Number: 3634626-XReport Type:Expedited (15-DaCompany Report #B0094555A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Bipolar Disorder		Zyban	PS	Glaxo Wellcome	ORAL
150MG As		Mania					
directed							
				Lithium	C		ORAL
600MG Per day				Paracetamol	C		ORAL

Date:12/21/00ISR Number: 3634627-1Report Type:Expedited (15-DaCompany Report #B0094565A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Disorientation		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Fear					
Hospitalization -							
per day	3	DAY					
Initial or Prolonged		Feeling Abnormal		Salbutamol	C	Glaxo Wellcome	
RESPIRATORY							
Other		Malaise					
(INHALATION)							
				Beclomethasone	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/21/00ISR Number: 3634628-3Report Type:Expedited (15-DaCompany Report #B0094566A  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
				Amlodipine	C		ORAL
				Atenolol	C		ORAL
50MG Twice							
per day							
75MG Per day				Aspirin	C		ORAL
7.5MG Per day				Meloxicam	C		ORAL

Date:12/21/00ISR Number: 3634630-1Report Type:Expedited (15-DaCompany Report #B0094567A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Psoriasis		Zyban	PS	Glaxo Wellcome	ORAL
28 DAY							

Date:12/21/00ISR Number: 3634631-3Report Type:Expedited (15-DaCompany Report #B0094571A  
Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day	36 DAY						
Hospitalization -		Myocardial Infarction		Aspirin	C		
75MG Twice							
Initial or Prolonged							
per day							
50MG Per day				Atenolol	C		
1PUFF As				Nitroglycerin	C	Glaxo Wellcome	
required							
TOPICAL				Fucibet	C		

5MG Per day	Finasteride	C	
10MG Per day	Lisinopril	C	
TOPICAL	Betamethasone	C	Glaxo Wellcome
300MG Per day	Quinine	C	

Date:12/21/00ISR Number: 3635694-1Report Type:Expedited (15-DaCompany Report #B0093969A  
 Age:37 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Asthma	Foreign	Bupropion			
Required	Erythema Multiforme		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ORAL						
Intervention to	Urticaria		Salbutamol Sulphate	C		
Prevent Permanent			Antihistamine	C		
Impairment/Damage						

Date:12/21/00ISR Number: 3635696-5Report Type:Expedited (15-DaCompany Report #B0093979A  
 Age:63 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Dyspnoea	Foreign	Bupropion			
Hospitalization -	Paralysis		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ORAL						
Initial or Prolonged			Atenolol +			
			Chlorothalidone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/21/00ISR Number: 3635697-7Report Type:Expedited (15-DaCompany Report #B0093975A  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Abortion Spontaneous	Foreign	Bupropion			
Intervention to		Menstruation Irregular		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
Prevent Permanent		Vaginal Haemorrhage					
PER DAY/ ORAL							
Impairment/Damage				Microgynon	C		
				Sultrin	C		

Date:12/21/00ISR Number: 3635700-4Report Type:Expedited (15-DaCompany Report #A0132705A  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Cellulitis	Foreign	Bupropion			
Other		Hypersensitivity	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
Required		Stevens-Johnson Syndrome	Professional				
PER DAY/ ORAL							
Intervention to							
Prevent Permanent							
Impairment/Damage							

Date:12/22/00ISR Number: 3635299-2Report Type:Expedited (15-DaCompany Report #A0135070A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Hypersensitivity		Zyban	PS	Glaxo Wellcome	

Date:12/22/00ISR Number: 3635320-1Report Type:Expedited (15-DaCompany Report #B0089482A  
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Left Ventricular Failure	Consumer	Bupropion			

150MG Twice	Oedema	Hydrochloride	PS	Glaxo Wellcome	ORAL
per day	20 DAY	Oedema Peripheral			
2PUFF At		Beclomethasone	C	Glaxo Wellcome	NASAL
night					

Date:12/22/00ISR Number: 3635322-5Report Type:Expedited (15-DaCompany Report #B0091290A  
 Age:57 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Condition Aggravated	Health	Bupropion			
Initial or Prolonged	Rheumatic Fever	Professional	Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG Twice	Systemic Lupus					
per day	Erythematosus		Decortin	C		
5MG Per day			Azathioprine	C	Glaxo Wellcome	
50MG Per day			Katadolon	C		
100MG Per day			Sortis	C		
10MG Per day			Magnerot N	C		
48.6MG Twice						
per day			Trusopt	C		
20MG Twice						
per day						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/22/00ISR Number: 3635324-9Report Type:Expedited (15-DaCompany Report #B0092236A  
 Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Twice per day	18 DAY	Astigmatism	Health	Zyban	PS	Glaxo Wellcome	ORAL
		Blood Pressure Increased	Professional				
4PUFF Twice per day		Eye Movement Disorder		Salbutamol	C	Glaxo Wellcome	
		Hypermetropia					
4PUFF Twice per day		Ophthalmoplegia		Beclomethasone Dipropionate	C	Glaxo Wellcome	
30MG Per day				Frusemide	C		
100MG Per day				Spiroinolactone	C		
.25MG Per day				Digoxin	C	Glaxo Wellcome	

Date:12/22/00ISR Number: 3635354-7Report Type:Expedited (15-DaCompany Report #B0094589A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Pain Upper		Bupropion			
		Arthralgia		Hydrochloride	PS	Glaxo Wellcome	ORAL
		Chest Pain		Losec	C		ORAL
		Cold Sweat		Vioxx	C		ORAL
		Condition Aggravated					
		Dizziness					
		Dry Mouth					
		Dry Throat					
		Feeling Hot					
		Hypertension					
		Ill-Defined Disorder					
		Myalgia					
		Ulcer					
		Vision Blurred					

Date:12/22/00ISR Number: 3635361-4Report Type:Expedited (15-DaCompany Report #B0094740A  
Age:54 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150MG Twice per day	Duration 7 DAY	Claustrophobia Depressed Mood Suicidal Ideation	Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY (INHALATION) day	10PUFF per		Salbutamol	C	Glaxo Wellcome	
RESPIRATORY (INHALATION) per day	500MCG Twice		Beclomethasone	C	Glaxo Wellcome	

Date:12/22/00ISR Number: 3635362-6Report Type:Expedited (15-DaCompany Report #B0094743A  
Age:51 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice per day	Duration 15 DAY	Dermatitis Exfoliative Oedema Peripheral Periorbital Oedema Urticaria	Zyban Hrt	PS C	Glaxo Wellcome	ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/22/00ISR Number: 3635363-8Report Type:Expedited (15-DaCompany Report #B0094744A  
Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	1 DAY	Acidosis	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -	1 DAY	Convulsion		Anadin	SS		ORAL
Initial or Prolonged		Hypotension					
Other		Intentional Misuse					
		Tachycardia					

Date:12/22/00ISR Number: 3635364-XReport Type:Expedited (15-DaCompany Report #B0094745A  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day 11 DAY	Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
	400MG per day	Confusional State		Etidronic Acid	C		ORAL
		Fatigue		Sulphasalazine	C		ORAL

Date:12/22/00ISR Number: 3635365-1Report Type:Expedited (15-DaCompany Report #B0094748A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	32 DAY	Erythema		Zyban	PS	Glaxo Wellcome	ORAL
				Co-Codamol	C		

Date:12/22/00ISR Number: 3636019-8Report Type:Expedited (15-DaCompany Report #A0134853A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cervix Carcinoma	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
SEE TEXT /							



Hospitalization - Drug Toxicity  
ORAL  
Initial or Prolonged Overdose

Date:12/22/00ISR Number: 3636319-1Report Type:Expedited (15-DaCompany Report #B0094552A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Chest Pain	Foreign	Bupropion			
Intervention to		Dry Mouth		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
Prevent Permanent		Insomnia					
VARIABLE DOSE							
Impairment/Damage		Vision Blurred					
/ ORAL							

Date:12/22/00ISR Number: 3636325-7Report Type:Expedited (15-DaCompany Report #B0094565A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Disorientation	Foreign	Bupropion			
Hospitalization -		Malaise		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
Initial or Prolonged							
TWICE PER DAY							
Required							
/ ORAL							
Intervention to				Salbutamol Sulphate	C		
Prevent Permanent				Beclomethasone			
Impairment/Damage				Dipropion	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/22/00ISR Number: 3636328-2Report Type:Expedited (15-DaCompany Report #B0093439A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG		Bronchospasm Dizziness	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
		Dyspnoea Tremor		Ipratropium Bromide	C		

Date:12/22/00ISR Number: 3636355-5Report Type:Expedited (15-DaCompany Report #B0094571A

Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 300 MG / PER Initial or Prolonged DAY/ ORAL		Angina Pectoris Myocardial Infarction	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Aspirin	C		
				Atenolol	C		
				Nitroglycerin	C		
				Fucibet	C		
				Finasteride	C		
				Lisinopril	C		
				Betamethasone	C		
				Quinine	C		

Date:12/22/00ISR Number: 3636357-9Report Type:Expedited (15-DaCompany Report #B0094567A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG / ORAL Prevent Permanent Impairment/Damage		Psoriasis	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:12/22/00ISR Number: 3636359-2Report Type:Expedited (15-DaCompany Report #B0094566A  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG/ ORAL Prevent Permanent Impairment/Damage		Angina Pectoris	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Amlodipine	C		
				Atenolol	C		
				Aspirin	C		
				Meloxicam	C		

Date:12/22/00ISR Number: 3639382-7Report Type:Expedited (15-DaCompany Report #B0094744A  
Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 7.5G/ORAL Initial or Prolonged Required 7.5G/ORAL Intervention to Prevent Permanent Impairment/Damage		Acidosis Convulsion	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Hypotension Intentional Misuse		Anadin (Formulation Unknown) (Anadin)	SS		ORAL
		Tachycardia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/26/00ISR Number: 3636409-3Report Type:Expedited (15-DaCompany Report #B0092443A  
 Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG / TWICE PER DAY		Disinhibition Medication Error	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Fluoxetine (Formulation Unknown) (Fluoxetine)	SS
Aspirin	C
Bendrofluazide	C
Fosinopril	C

Date:12/26/00ISR Number: 3638607-1Report Type:Expedited (15-DaCompany Report #A0135070A  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Hypersensitivity	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:12/26/00ISR Number: 3638608-3Report Type:Expedited (15-DaCompany Report #A0135193A  
 Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 3 MON		Completed Suicide	Literature	Wellbutrin	PS	Glaxo Wellcome Inc	
		Euphoric Mood Murder	Health Professional Company Representative	Pain Medication (Formulation Unknown) (Pain Medication)	SS		

Date:12/26/00ISR Number: 3638775-1Report Type:Expedited (15-DaCompany Report #A0128274A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG TWICE Prevent Permanent PER DAY ORAL Impairment/Damage		Anxiety Feeling Abnormal Feeling Hot Hyperaesthesia Irritability Nervousness Nightmare Pallor Pruritus Swelling Urticaria	Health Professional	Bupropion Hydrochloride  Nicotine	PS  C	Glaxo Wellcome Inc	ORAL

Date:12/26/00ISR Number: 3638777-5Report Type:Expedited (15-DaCompany Report #A0121104A  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG ORAL		Drug Interaction Myocardial Infarction	Health Professional	Bupropion Hydrochloride  Sildenafil Citrate Tablet (Sildenafil Citrate)	PS  SS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Co-Trimoxazole C

Date:12/26/00ISR Number: 3639159-2Report Type:Expedited (15-DaCompany Report #B0092236A  
Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG / TWICE PER DAY / ORAL	Astigmatism Blood Pressure Increased Hypermetropia Ophthalmoplegia Visual Disturbance	Foreign Health Professional	Bupropion Hydrochloride Salbutamol Sulphate Beclomethasone Dipropion Frusemide Spironolactone Digoxin	PS  C C C C	Glaxo Wellcome Inc	ORAL

Date:12/26/00ISR Number: 3639160-9Report Type:Expedited (15-DaCompany Report #B0091290A  
Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG / TWICE PER DAY / ORAL		Rheumatic Fever Systemic Lupus Erythematosis	Foreign Study Health Professional	Bupropion Hydrochloride Prednisone Azathioprine Flupirtine Maleate Atorvastatin Calcium Magnerot N Dorzolamide Hcl	PS  C C C C C C	Glaxo Wellcome Inc	ORAL

Date:12/26/00ISR Number: 3639161-0Report Type:Expedited (15-DaCompany Report #B0089482A  
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG / Prevent Permanent TWICE PER DAY Impairment/Damage / ORAL		Left Ventricular Failure Oedema  Oedema Peripheral	Foreign	Bupropion Hydrochloride  Beclomethasone Dipropion	PS  C	Glaxo Wellcome Inc	ORAL

Date:12/26/00ISR Number: 3639375-XReport Type:Expedited (15-DaCompany Report #B0094740A  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability  150 MG/TWICE  PER DAY/ORAL		Claustrophobia Depressed Mood  Suicidal Ideation	Foreign	Bupropion Hydrochloride  Salbutamol Sulphate Beclomethasone Dipropion	PS  C C	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/26/00ISR Number: 3639377-3Report Type:Expedited (15-DaCompany Report #B0094589A

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Abdominal Pain Upper Arthralgia	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
	Blood Pressure Increased		Omeprazole	C		
	Chest Pain		Rofecoxib	C		
	Cold Sweat					
	Condition Aggravated					
	Dizziness					
	Dry Mouth					
	Dry Throat					
	Feeling Hot					
	Heart Rate Increased					
	Malaise					
	Myalgia					
	Ulcer					
	Vision Blurred					

Date:12/26/00ISR Number: 3639378-5Report Type:Expedited (15-DaCompany Report #B0094748A

Age:52 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150 MG/ORAL	Rash Erythematous	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
			Co-Codamol	C		

Date:12/26/00ISR Number: 3639380-3Report Type:Expedited (15-DaCompany Report #B0094745A

Age:62 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to 150 MG/PER Prevent Permanent DAY/ORAL Impairment/Damage	Arthralgia Confusional State Fatigue	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
			Etidronic Acid	C		



Date:12/26/00ISR Number: 3639381-5Report Type:Expedited (15-DaCompany Report #B0094743A  
 Age:51 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Dermatitis Exfoliative	Foreign	Bupropion			
Initial or Prolonged	Oedema Peripheral		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE						
PER DAY/ORAL	Periorbital Oedema					
	Urticaria		Hrt	C		

Date:12/26/00ISR Number: 3639965-4Report Type:Expedited (15-DaCompany Report #B0094291A  
 Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Psoriasis	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL 3 WK						
Initial or Prolonged		Health Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/26/00ISR Number: 3639966-6Report Type:Expedited (15-DaCompany Report #B0094077A  
Age:55 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY	Angina Pectoris	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
/ ORAL			Aspirin	C		

Date:12/26/00ISR Number: 3639967-8Report Type:Expedited (15-DaCompany Report #B0094062A  
Age:40 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 300 MG / PER DAY / ORAL	Urticaria	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:12/26/00ISR Number: 3639969-1Report Type:Expedited (15-DaCompany Report #B0094079A  
Age:40 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150 MG / Required TWICE PER DAY Intervention to / ORAL Prevent Permanent Impairment/Damage	Dry Mouth  Feeling Abnormal  Headache  Malaise Pain Parosmia Syncope	Foreign	Zyban    Propranolol Hydrochloride	PS    C	Glaxo Wellcome Inc	ORAL

Date:12/28/00ISR Number: 3637903-1Report Type:Expedited (15-DaCompany Report #A0135312A  
Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	3	YR	Atrioventricular Block	Wellbutrin	PS	Glaxo Wellcome	ORAL
900MG Per day				Lithium	C		
10MG As				Largactil	C		

required

Date:12/28/00ISR Number: 3638077-3Report Type:Direct  
Age:44 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 AM + HS			Grand Mal Convulsion	Wellbutrin Sr Glaxo/Welcome	PS	Glaxo/Welcome	ORAL
PO				Olanzapine	C		

Date:12/29/00ISR Number: 3638853-7Report Type:Direct  
Age:49 YR Gender:Male I/FU:I

Company Report #

Outcome	PT
Hospitalization - Initial or Prolonged	Dermatitis Dysphagia Face Oedema Oedema Peripheral Pruritus Skin Lesion

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Urticaria

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
			Zyban Nicotine Inhaler	PS C		

Date:12/29/00ISR Number: 3640478-4Report Type:Periodic Company Report #A001253  
Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Eructation Nausea Sedation	Health Professional	Vistaril Wellbutrin Diphenhydramine/Pseu doephedrine Tagamet Hb Hydrocotone Valerate Cream	PS SS SS SS C	Pfizer Laboratories Div Pfizer Inc	

Date:01/02/01ISR Number: 3640473-5Report Type:Expedited (15-DaCompany Report #2000AU03968  
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 20 MG QD PO Initial or Prolonged		Blood Potassium Decreased Drug Interaction Drug Toxicity Heart Rate Increased	Health Professional	Prilosec Lithium Trazodone Wellbutrin	PS SS SS SS	Astrazeneca Lp	ORAL

Date:01/02/01ISR Number: 3640862-9Report Type:Expedited (15-DaCompany Report #A0135312A  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL		Atrioventricular Block	Foreign Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Professional

Lithium Salt

C

Largactil

C

Date:01/02/01ISR Number: 3640920-9Report Type:Expedited (15-DaCompany Report #HQ5327428DEC2000

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Craniopharyngioma	Health	Effexor Xr	PS	Wyeth Ayerst	
		Mouth Ulceration	Professional			Laboratories	ORAL
112.5 MG		Oral Pain					
DAILY, ORAL		Tardive Dyskinesia		Wellbutrin			
				(Amfebutamone			
				Hydrochloride, )	SS		ORAL
ORAL							

Date:01/03/01ISR Number: 3640294-3Report Type:Expedited (15-DaCompany Report #B0094866A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Asthenia		Zyban	PS	Glaxo Wellcome	
UNKNOWN							
Initial or Prolonged		Blood Potassium Decreased		Unspecified			
		Blood Sodium Decreased		Medications	C		
		Malaise					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/03/01  
 Age:           Gender:Female       I/FU:I  
 Report Type:Direct

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Wellbutrin Sr 100 Mg	PS		
100 MG BID		Vomiting					
ORAL							

Date:01/04/01  
 Age:           Gender:Female       I/FU:F  
 Report Type:Expedited (15-Da  
 Company Report #B0093061A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Diplopia	Consumer	Zyban	PS	Glaxo Wellcome	
		Insomnia					
		Tinnitus					
		Tongue Disorder					
		Vision Blurred					

Date:01/04/01  
 Age:           Gender:Female       I/FU:F  
 Report Type:Expedited (15-Da  
 Company Report #B0094941A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Gastritis		Zyban	PS	Glaxo Wellcome	
1 MON							
Initial or Prolonged		Rectal Haemorrhage					

Date:01/04/01  
 Age:42 YR   Gender:Male       I/FU:I  
 Report Type:Direct

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Convulsion		Wellbutrin Sr (Glaxo Wellcome)	PS	Glaxo Wellcome	
Hospitalization -		Diabetes Mellitus					
100MG 2 TABS							
Initial or Prolonged		Drug Interaction					
AM /PM							
		Ecchymosis		Clozaril (Novartis			

100MG 3 BID	Fall Haemorrhage	Pharmaceutical)	SS	Novartis Pharmaceutical
	Head Injury	Norvasc	C	
	Hepatic Failure	Zestril	C	
	Hypertension	Colace	C	
	Intestinal Ischaemia			
	Loss Of Consciousness			
	Medication Error			
	Mental Retardation			
	Severity Unspecified			
	Obesity			
	Schizoaffective Disorder			
	Sudden Death			

Date:01/05/01ISR Number: 3641975-8Report Type:Direct Company Report #  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tinnitus		Zyban 150 Mg	PS		ORAL
1 TABLET PER							

DAY=ROUTE=  
 ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/05/01ISR Number: 3642326-5Report Type:Expedited (15-DaCompany Report #B0091930A  
Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Acidosis		Zyban	PS	Glaxo Wellcome	ORAL
50TAB Single							
Hospitalization -		Coma					
dose	1 DAY						
Initial or Prolonged		Hypotension		Anadin Extra	SS		ORAL
24TAB Single							
		Intentional Misuse					
dose	1 DAY	Vomiting					

Date:01/05/01ISR Number: 3642335-6Report Type:Expedited (15-DaCompany Report #B0095016A  
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Arrhythmia		Zyban	PS	Glaxo Wellcome	ORAL
150MG See							
Hospitalization -		Cardiac Arrest					
text							
Initial or Prolonged		Coma					
Other		Convulsion					
		Intentional Misuse					
		Suicide Attempt					

Date:01/05/01ISR Number: 3642336-8Report Type:Expedited (15-DaCompany Report #B0095023A  
Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Renal Colic		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Initial or Prolonged							
per day				Analgesics	C		



Date:01/05/01ISR Number: 3642337-XReport Type:Expedited (15-DaCompany Report #B0095024A  
Age:73 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Petechiae Vasculitis		Zyban	PS	Glaxo Wellcome	

Date:01/05/01ISR Number: 3642341-1Report Type:Expedited (15-DaCompany Report #B0095172A  
Age:73 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150MG Twice Initial or Prolonged per day	Cerebrovascular Accident Urinary Tract Infection		Zyban	PS	Glaxo Wellcome	ORAL

Date:01/05/01ISR Number: 3642343-5Report Type:Expedited (15-DaCompany Report #D0013022A  
Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other 150MG Twice per day	Lenticular Opacities Vision Blurred		Zyban	PS	Glaxo Wellcome	ORAL
	13 DAY					

Date:01/05/01ISR Number: 3642344-7Report Type:Expedited (15-DaCompany Report #D0013064A  
Age:35 YR Gender:Male I/FU:I

Outcome	PT
Other	Dermatitis Atopic Skin Disorder

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Freedom Of Information (FOI) Report

Skin Exfoliation

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1TAB			Zyban	PS	Glaxo Wellcome	
cumulative dose	1 DAY					

Date:01/08/01ISR Number: 3642639-7Report Type:Direct  
 Age: Gender:Male I/FU:I Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Fatigue		Sporanox 100 Mg	PS		ORAL
Intervention to		Myalgia					
Prevent Permanent		Pyrexia					
ORAL Impairment/Damage		Rash Erythematous Rash Pruritic		Wellbutrin	SS		

Date:01/08/01ISR Number: 3643524-7Report Type:Expedited (15-DaCompany Report #B0095016A  
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization -		Arrhythmia Cardiac Arrest	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ SEE							
Initial or Prolonged		Coma	Professional				
TEXT/ ORAL							
Required		Convulsion					
Intervention to		Overdose					
Prevent Permanent		Suicide Attempt					
Impairment/Damage							

Date:01/08/01ISR Number: 3643532-6Report Type:Expedited (15-DaCompany Report #B0091930A  
Age:18 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 50 TABLET/ Initial or Prolonged SINGLE	Acidosis Coma Hypotension Intentional Misuse Vomiting	Foreign Health Professional	Bupropion Hydrochloride  Anadin Extra Tablet (Anadin Extra)	PS  SS	Glaxo Wellcome Inc	ORAL  ORAL
DOSE/ORAL						
24						
TABLET/SINGLE						
DOSE/ORAL						

Date:01/08/01ISR Number: 3643552-1Report Type:Expedited (15-DaCompany Report #D0013064A  
Age:35 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other	Hypersensitivity Rash Generalised Skin Disorder	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:01/08/01ISR Number: 3643555-7Report Type:Expedited (15-DaCompany Report #B0095172A  
Age:73 YR Gender:Female I/FU:I

Outcome	PT	Report Source
Hospitalization - Initial or Prolonged	Cerebrovascular Accident Urinary Tract Infection	Foreign Health

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Professional

Dose	Duration	Product	Role	Manufacturer	Route
150 MG (TWICE PER DAY) ORAL		Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:01/08/01ISR Number: 3643556-9Report Type:Expedited (15-DaCompany Report #D0013022A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Lenticular Opacities Vision Blurred	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL		Visual Disturbance	Professional				

Date:01/08/01ISR Number: 3643558-2Report Type:Expedited (15-DaCompany Report #B0095024A  
Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG	Petechiae Vasculitis	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
			Professional				

Date:01/08/01ISR Number: 3643559-4Report Type:Expedited (15-DaCompany Report #B0095023A  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG (TWICE PER DAY) ORAL	Renal Colic	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
			Professional				

Date:01/08/01ISR Number: 3643693-9Report Type:Expedited (15-DaCompany Report #B0094941A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1 MON		Gastritis Haematemesis  Rectal Haemorrhage Vomiting	Foreign Health  Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:01/08/01ISR Number: 3643694-0Report Type:Expedited (15-DaCompany Report #B0093061A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability  150 MG		Diplopia Insomnia  Tinnitus Tongue Disorder Vision Blurred Visual Disturbance	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

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Freedom Of Information (FOI) Report

Date:01/08/01ISR Number: 3644405-5Report Type:Expedited (15-DaCompany Report #B0094866A

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Asthenia	Foreign	Bupropion			
Initial or Prolonged	Blood Potassium Decreased	Health	Hydrochloride	PS	Glaxo Wellcome Inc	
	Blood Sodium Decreased	Professional				
	Malaise					

Date:01/08/01ISR Number: 3653655-3Report Type:Periodic Company Report #2000029474US

Age:78 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Dry Mouth	Consumer	Detrol	PS	Glaxo Wellcome Inc	ORAL
2 DF QD ORAL	Movement Disorder		Cefalexin (Cefalexin)	SS		
			Estraderm	C		
			Hydrodiuril	C		
			Slow-K	C		
			Elavil (Amitriptyline Hydrochloride)	C		
			Synthroid (Levothyroxine Sodium)	C		
			Urised (Metylionium Chloride, Benzoic Acid, Methenamine, Hyoscyamine, Phenylsalycate, Entex La	C C		

Date:01/08/01ISR Number: 3653875-8Report Type:Periodic Company Report #2000026940US

Age:68 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Dry Mouth	Consumer	Detrol	PS	Pharmacia & Upjohn	ORAL
2 MG BID ORAL	Dry Skin		Plendil	C		

Skin Exfoliation

Premarin (Estrogens  
Conjugated)

C

Date:01/08/01ISR Number: 3655507-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #2000033755US

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Detrol	PS	Pharmacia And Upjohn Co	ORAL
ORAL		Dry Mouth		Zyban/Amfebutamone	SS		

Date:01/09/01ISR Number: 3643529-6Report Type:Expedited (15-DaCompany Report #A0136085A  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Arthritis		Wellbutrin	PS	Glaxo Wellcome	ORAL
300MG Per day		Gastrointestinal Disorder		Adalat	C		
		Pain In Extremity		Synthroid	C	Glaxo Wellcome	
				Lipitor	C		
				Hytrin	C		

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Freedom Of Information (FOI) Report

Date:01/09/01ISR Number: 3643533-8Report Type:Expedited (15-DaCompany Report #B0092403A  
Age:63 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150MG Twice	Angina Unstable		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - per day 7 WK Initial or Prolonged	Atrial Fibrillation					

Date:01/09/01ISR Number: 3643537-5Report Type:Expedited (15-DaCompany Report #B0095014A  
Age:42 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 3 WK Initial or Prolonged	Amnesia		Zyban	PS	Glaxo Wellcome	
	Diabetes Mellitus Disturbance In Attention Haemorrhage Menstruation Irregular Paraesthesia Polydipsia Visual Disturbance					

Date:01/09/01ISR Number: 3643538-7Report Type:Expedited (15-DaCompany Report #B0095288A  
Age:55 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - Initial or Prolonged	Angina Pectoris Dyspnoea		Zyban	PS	Glaxo Wellcome	ORAL
60MG Per day			Amiodarone	C		ORAL
			Glycerol Trinitrate	C	Glaxo Wellcome	
			Isosorbide	C		ORAL
50MG As			Paracetamol	C		ORAL
required			Tramadol	C		ORAL
20MG Per day			Simvastatin	C		ORAL



12.5MG Per		Bisoprolol	C	ORAL
day		Pantoprazole	C	ORAL
10MG Twice		Clopidogrel	C	ORAL
per day		Rofecoxib	C	ORAL
		Nicorandil	C	ORAL

Date:01/09/01ISR Number: 3643540-5Report Type:Expedited (15-DaCompany Report #B0095293A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		International Normalised		Zyban	PS	Glaxo Wellcome	ORAL
Other		Ratio Increased					
150MG Twice	8 DAY			Warfarin	C	Glaxo Wellcome	
per day				Methotrexate	C		ORAL
10MG Weekly				Folic Acid	C		ORAL
5MG Two times							
per week				Co-Amilofruse	C		ORAL

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Freedom Of Information (FOI) Report

Date:01/09/01ISR Number: 3643541-7Report Type:Expedited (15-DaCompany Report #B0095307A  
Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Atrial Fibrillation		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	29 DAY			Pravastatin	C		ORAL
20MG Per day				Lisinopril	C		ORAL
10MG Per day				Gliclazide	C		ORAL
80MG Twice							
per day							

Date:01/09/01ISR Number: 3643542-9Report Type:Expedited (15-DaCompany Report #B0095308A  
Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	8 DAY			Gliclazide	C		ORAL
80MG Per day		Dyspnoea		Diltiazem	C		ORAL
300MG Per day				Nizatidine	C		ORAL
100MG Per day				Isosorbide Mononitrate	C		ORAL

Date:01/10/01ISR Number: 3644729-1Report Type:Expedited (15-DaCompany Report #A0130183A  
Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Toxicologic Test Abnormal	Consumer	Bupropion Atropine	PS SS	Glaxo Wellcome	
UNKNOWN				Chloral Hydrate	SS		
UNKNOWN							

UNKNOWN

Diazepam	SS	
Venlafaxine	SS	
Methadone	SS	Glaxo Wellcome
Sertraline	SS	
Lidocaine	SS	

Date:01/10/01ISR Number: 3644731-XReport Type:Expedited (15-DaCompany Report #A0131760A  
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	350MG Per day 8 MON	Blood Glucose Increased	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
UNKNOWN	100MG Per day 1 YR	Hepatitis C Hepatitis Infectious	Professional	Zoloft	C		

Date:01/11/01ISR Number: 3645633-5Report Type:Expedited (15-DaCompany Report #B0095597A  
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Twice	Aphasia		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -	per day 5 DAY	Hemiparesis					
Initial or Prolonged		Hemiplegia					

Date:01/12/01ISR Number: 3647307-3Report Type:Expedited (15-DaCompany Report #B0095288A  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG ORAL	Angina Pectoris	Foreign Other	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Hospitalization -	Initial or Prolonged			Amiodarone	C		

Freedom Of Information (FOI) Report

Nitroglycerin	C
Isosorbide	C
Paracetamol	C
Tramadol	
Hydrochloride	C
Simvastatin	C
Bisoprolol	C
Pantoprazole	C
Clopidogrel	C
Rofecoxib	C
Nicorandil	C

Date:01/12/01ISR Number: 3647308-5Report Type:Expedited (15-DaCompany Report #B0095307A  
Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG (PER Prevent Permanent DAY) ORAL Impairment/Damage		Atrial Fibrillation	Foreign Other	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Pravastatin	C		
				Lisinopril	C		
				Gliclazide	C		

Date:01/12/01ISR Number: 3647311-5Report Type:Expedited (15-DaCompany Report #B0095293A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG / Prevent Permanent TWICE PER DAY Impairment/Damage / ORAL		International Normalised Ratio Increased	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Warfarin	C		
				Methotrexate	C		
				Folic Acid	C		
				C0-Amilofruse	C		

Date:01/12/01ISR Number: 3647317-6Report Type:Expedited (15-DaCompany Report #B0092403A  
Age:63 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Angina Unstable	Foreign	Bupropion			
Hospitalization -	Atrial Fibrillation	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE						
Initial or Prolonged	Chest Pain	Professional				
PER DAY/ ORAL						

Date:01/12/01ISR Number: 3647319-XReport Type:Expedited (15-DaCompany Report #B0095308A  
Age:69 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Required	Angina Pectoris	Foreign	Bupropion			
Intervention to	Dyspnoea		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER						
Prevent Permanent						
DAY/ ORAL						
Impairment/Damage			Gliclazide	C		
			Diltiazem	C		
			Nizatidine	C		
			Isosorbide			
			Mononitrate	C		

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Freedom Of Information (FOI) Report

Date:01/12/01ISR Number: 3647320-6Report Type:Expedited (15-DaCompany Report #B0095014A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Amnesia Blindness Diabetes Mellitus Disturbance In Attention Haemorrhage Menstruation Irregular Mood Altered Paraesthesia Visual Disturbance	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:01/12/01ISR Number: 3647321-8Report Type:Expedited (15-DaCompany Report #A0136085A  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 300 MG/ PER DAY / ORAL		Arthritis Gastrointestinal Disorder Pain In Extremity	Foreign Health Professional	Wellbutrin Sr Nifedipine Thyroxine Sodium Atorvastatin Calcium Terazosin Hydrochloride	PS C C C C	Glaxo Wellcome Inc	ORAL

Date:01/15/01ISR Number: 3646711-7Report Type:Expedited (15-DaCompany Report #A0135962A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	2 WK	Chest Pain Dyspnoea Haemorrhage Lung Infiltration Nervous System Disorder Respiratory Arrest		Zyban	PS	Glaxo Wellcome	ORAL

Sarcoidosis

Date:01/15/01ISR Number: 3646713-0Report Type:Expedited (15-DaCompany Report #A0136308A  
Age:27 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly 400MG per day	Abortion Spontaneous		Wellbutrin	PS	Glaxo Wellcome	
	Complications Of Maternal Exposure To Therapeutic Drugs Congenital Anomaly Pregnancy					

Date:01/15/01ISR Number: 3646718-XReport Type:Expedited (15-DaCompany Report #B0093439A  
Age:61 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Disability Other	Bronchospasm Chest Discomfort Disorientation Dizziness Dyspnoea

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Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
7 DAY			Zyban	PS	Glaxo Wellcome	
RESPIRATORY (INHALATION)	120MCG per day		Combivent	C		

Date:01/15/01ISR Number: 3646719-1Report Type:Expedited (15-DaCompany Report #B0094653A  
Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day	Chest Pain		Zyban	PS	Glaxo Wellcome	ORAL
	4 DAY	Difficulty In Walking		Oestrogen	C		ORAL
625MCG per day				Fluvastatin	C		ORAL
40MG per day				Aspirin	C		ORAL
250MG per day							

Date:01/15/01ISR Number: 3646722-1Report Type:Expedited (15-DaCompany Report #B0095614A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	11 DAY	Chest Pain		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	150MG Per day			Aspirin	C		ORAL
				Alcohol	C		



Date:01/16/01ISR Number: 3647439-XReport Type:Expedited (15-DaCompany Report #A0131854A  
Age:70 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 8 DAY Initial or Prolonged Disability		Amnesia Cerebrovascular Accident Fall		Zyban	PS	Glaxo Wellcome	ORAL

Date:01/16/01ISR Number: 3647443-1Report Type:Expedited (15-DaCompany Report #A0136085A  
Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Arthralgia Arthritis		Wellbutrin	PS	Glaxo Wellcome	ORAL
60MG Per day		Gastrointestinal Disorder		Adalat Xl	C		ORAL
300MG Per day				Synthroid	C	Glaxo Wellcome	ORAL
40MG Per day				Lipitor	C		ORAL
				Hytrin	C		

Date:01/16/01ISR Number: 3647458-3Report Type:Expedited (15-DaCompany Report #B0095936A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2 WK Initial or Prolonged		Bronchospasm Lower Respiratory Tract Infection Respiratory Depression		Zyban	PS	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/16/01ISR Number: 3648779-0Report Type:Expedited (15-DaCompany Report #A0131760A  
Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hepatitis C	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
350 MG / PER							
DAY / ORAL	8	MON	Professional				
		Hepatitis Infectious		Sertraline Hydrochloride	C		

Date:01/16/01ISR Number: 3648799-6Report Type:Expedited (15-DaCompany Report #A0130183A  
Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Toxicologic Test Abnormal	Foreign	Wellbutrin	PS	Glaxo Wellcome Inc	
				Atropine	SS		
				Chloral Hydrate	SS		
				Diazepam	SS		
				Venlafaxine			
				Hydrochloride	SS		
				Methadone			
				Hydrochloride	SS		
				Sertraline	SS		
				Lignocaine	SS		

Date:01/16/01ISR Number: 3648872-2Report Type:Expedited (15-DaCompany Report #B0095597A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Aphasia	Foreign	Bupropion			
Hospitalization -		Hemiplegia	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
Initial or Prolonged			Professional				
PER DAY/ORAL							

Date:01/17/01ISR Number: 3648118-5Report Type:Direct Company Report #  
Age:6.5 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia		Wellbutrin			
		Dysuria		-Bupropion	PS		
50 MG QOD X 2							
WKS THEN 50		Erythema Multiforme					
MG QD		Joint Swelling					
		Pharyngolaryngeal Pain					
		Pyrexia					

Date:01/17/01ISR Number: 3648137-9Report Type:Expedited (15-DaCompany Report #A0131639A  
Age:73 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Colitis Ischaemic		Zantac	PS	Glaxo Wellcome	ORAL
150MG Per day 2 YR							
Initial or Prolonged		Hyperkalaemia		Zyban	SS	Glaxo Wellcome	ORAL
150MG Per day							
		Megaloblasts Increased		Theophylline	C		
		Rectal Haemorrhage		Multivitamin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/17/01ISR Number: 3648144-6Report Type:Expedited (15-DaCompany Report #A0136900A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG per day	46 DAY	Abortion Spontaneous	Bupropion	PS	Glaxo Wellcome	ORAL
			Haemorrhage				

Date:01/17/01ISR Number: 3648147-1Report Type:Expedited (15-DaCompany Report #B0088136A  
Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150MG Twice		Confusional State Headache	Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
per day			Loss Of Consciousness				
			Pallor Post-Traumatic Amnestic Disorder Tonic Convulsion	Contraceptive	C		ORAL

Date:01/17/01ISR Number: 3648154-9Report Type:Expedited (15-DaCompany Report #B0095172A  
Age:74 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150MG Twice		Cerebral Infarction	Zyban	PS	Glaxo Wellcome	ORAL
per day	7 DAY		Cerebrovascular Accident				
Other RESPIRATORY (INHALATION)			Trismus	Beclomethasone	C	Glaxo Wellcome	
			Urinary Tract Infection				
1PUFF Twice				Beconase Nasal	C	Glaxo Wellcome	NASAL
per day				Aspirin	C		ORAL
75MG per day							

60MG Twice  
 per day  
 5G per day

Theophylline C ORAL

Amlodipine Besylate C ORAL

Date:01/17/01ISR Number: 3648160-4Report Type:Expedited (15-DaCompany Report #B0095928A  
 Age:36 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 300MG Per day 39 DAY	Hydrocephalus		Zyban	PS	Glaxo Wellcome	ORAL

Date:01/18/01ISR Number: 3649141-7Report Type:Direct Company Report #  
 Age:35 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - P.O. 1 TAB Initial or Prolonged EACH AM	Grand Mal Convulsion		Wellbutrin 150 Sr	PS		ORAL

INCREASED TO  
 BID X 1 1/2  
 WK

Prozac C  
 Oral Contraceptive C  
 Proventil C  
 Entex C  
 Flonase C  
 Allegra C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/18/01ISR Number: 3649162-4Report Type:Direct  
 Age:54 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG BID PO	Bacterial Infection		Zyban (Bupropion 300mg) Glaxo Wellcome	PS	Glaxo Wellcome	ORAL
				Hydrochlorthiazide Ultram	C C		

Date:01/18/01ISR Number: 3649205-8Report Type:Expedited (15-DaCompany Report #A0126045A  
 Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG At night		Sudden Death		Wellbutrin	PS	Glaxo Wellcome	ORAL
				Tegretol	C		
				Zoloft	C		
				Ritalin	C		

Date:01/18/01ISR Number: 3649206-XReport Type:Expedited (15-DaCompany Report #A0134853A  
 Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Three Hospitalization - times per day 6 DAY Initial or Prolonged 1TAB Per day		Abdominal Adhesions Anaemia Anorexia		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Chest Pain Fistula Hypoaesthesia Lung Disorder Metastases To Lymph Nodes Neutropenia Night Sweats		Triphasil	C		ORAL
				St. John'S Wort	C		
				Vitamins	C		

Orthostatic Hypotension  
 Paraesthesia  
 Post Coital Bleeding  
 Pyrexia  
 Rhinitis  
 Sinus Tachycardia  
 Squamous Cell Carcinoma  
 Of The Cervix  
 Thyroid Function Test  
 Abnormal  
 Vaginal Haemorrhage

Date:01/18/01ISR Number: 3649212-5Report Type:Expedited (15-DaCompany Report #A0137061A

Age:17 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
4500MG per		Sinus Tachycardia					
day		Suicide Attempt		Celexa	SS		ORAL
400MG per day							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/18/01ISR Number: 3649214-9Report Type:Expedited (15-DaCompany Report #A0137073A

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged		Angina Pectoris	Zyban	PS	Glaxo Wellcome	ORAL

Date:01/18/01ISR Number: 3649215-0Report Type:Expedited (15-DaCompany Report #B0093922A

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged		Intervertebral Disc Protrusion Sciatica	Zyban	PS	Glaxo Wellcome	ORAL

Date:01/18/01ISR Number: 3649217-4Report Type:Expedited (15-DaCompany Report #B0095016A

Age:25 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150MG See Hospitalization - text Initial or Prolonged Other		Arrhythmia Brain Oedema Cardiac Arrest Coma Convulsion Intentional Misuse Suicide Attempt	Zyban	PS	Glaxo Wellcome	ORAL

Date:01/18/01ISR Number: 3649232-0Report Type:Expedited (15-DaCompany Report #B0096058A

Age:53 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice Initial or Prolonged per day		Psoriasis	Zyban	PS	Glaxo Wellcome	ORAL

24 DAY



Date:01/18/01ISR Number: 3649233-2Report Type:Expedited (15-DaCompany Report #B0096062A  
Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
15 DAY		Constipation		Zyban	PS	Glaxo Wellcome	ORAL
10MG Twice		Dermatitis Exfoliative		Nifedipine	C		ORAL
per day		Pruritus					
2.5MG In the				Bendrofluazide	C	Glaxo Wellcome	
morning							
5MG At night				Nitrazepam	C		
				Co-Codamol	C		

Date:01/18/01ISR Number: 3649234-4Report Type:Expedited (15-DaCompany Report #B0096063A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death							
150MG Twice		Cerebrovascular Accident		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -							
per day							
Initial or Prolonged							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/18/01ISR Number: 3649235-6Report Type:Expedited (15-DaCompany Report #D0012242A

Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day 2 WK	Hemiparesis		Zyban	PS	Glaxo Wellcome	ORAL
				Iodid 200	C		ORAL
				Bikalm	C		ORAL
				Diclofenac	C		
				Contraceptives	C		

Date:01/19/01ISR Number: 3650387-2Report Type:Direct

Company Report #

Age:82 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - SR 150MG / Initial or Prolonged DAY		Grand Mal Convulsion		Wellbutrin Sr	PS		
				Coumadin	C		
				Paxil	C		
				Bumex	C		
				Tegretol	C		
				Prinivil	C		
				Premarin	C		
				Synthroid	C		
				Verapamil	C		
				Ativan	C		

Date:01/19/01ISR Number: 3651566-0Report Type:Expedited (15-DaCompany Report #A0136085A

Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG/ TWICE	Osteoarthritis	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

PER DAY /

Health

ORAL

Professional

Nifedipine	C
Thyroxine Sodium	C
Atorvastatin Calcium	C
Terazosin	
Hydrochloride	C

Date:01/19/01ISR Number: 3651567-2Report Type:Expedited (15-DaCompany Report #A0131854A

Age:70 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG/ PER	Amnesia Aneurysm	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Disability DAY / ORAL		Cerebrovascular Accident	Professional				
		Fall Paraesthesia					

Date:01/19/01ISR Number: 3651568-4Report Type:Expedited (15-DaCompany Report #B0095936A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	2 WK	Lower Respiratory Tract Infection	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
			Professional				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/19/01ISR Number: 3651594-5Report Type:Expedited (15-DaCompany Report #B0093439A  
Age:61 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG	Duration Bronchospasm	Foreign	Zyban	PS	Glaxo Wellcome Inc	
Initial or Prolonged Disability Required Intervention to Prevent Permanent Impairment/Damage	Chest Discomfort Disorientation Dizziness Dyspnoea Tremor	Health Professional	Combivent	C		

Date:01/19/01ISR Number: 3651600-8Report Type:Expedited (15-DaCompany Report #A0136308A  
Age:27 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly	Duration Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs Foetal Disorder Pregnancy	Foreign Study Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	

Date:01/19/01ISR Number: 3651601-XReport Type:Expedited (15-DaCompany Report #B0095614A  
Age:42 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG/ ORAL	Duration Chest Pain	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged	Coronary Artery Disease		Aspirin Ethanol	C C		

Date:01/19/01ISR Number: 3651602-1Report Type:Expedited (15-DaCompany Report #B0094653A  
Age:56 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Disability	Chest Pain	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER						
	Myalgia	Health				
DAY/ ORAL		Professional	Estrogen	C		
			Fluvastatin Sodium	C		
			Aspirin	C		

Date:01/19/01ISR Number: 3652486-8Report Type:Expedited (15-DaCompany Report #A0135962A  
Age:36 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Chest Pain	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE						
Initial or Prolonged	Dyspnoea	Professional				
PER DAY/ ORAL						
	Haemorrhage					
	Lung Infiltration					
	Nervous System Disorder					
	Respiratory Disorder					
	Sarcoidosis					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/22/01ISR Number: 3651242-4Report Type:Expedited (15-DaCompany Report #A0133472A

Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Anxiety		Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY (INHALATION) per day	500MCG	Asthma Convulsion Decreased Appetite Twice Disturbance In Attention		Advair	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)		Fatigue Fear Feeling Abnormal Insomnia Nausea Nervousness Oedema Peripheral Speech Disorder Tremor		Ventolin	C	Glaxo Wellcome	

Date:01/22/01ISR Number: 3651243-6Report Type:Expedited (15-DaCompany Report #A0136520A

Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 60TABS cumulative dose		Haemorrhage Intentional Misuse Suicide Attempt Thyroid Gland Cancer		Zyban	PS	Glaxo Wellcome	
				Quinidine	SS		
				Benadryl	SS		
				Alcohol	SS		

Date:01/22/01ISR Number: 3651259-XReport Type:Expedited (15-DaCompany Report #B0095557A  
Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermographism		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Urticaria Papular					
per day	18	DAY					

Date:01/22/01ISR Number: 3651262-XReport Type:Expedited (15-DaCompany Report #B0096064A  
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Anxiety					
per day	20	DAY					
		Chest Pain		Alcohol	C		
		Confusional State		Imovane	C		
		Depression		Sobril	C		
		Dermatitis		Naproxen	C		
		Disturbance In Attention					
		Dysuria					
		Hallucination					
		Hyperhidrosis					
		Insomnia					
		Malaise					
		Nausea					
		Pyrexia					
		Tremor					
		Vertigo					
		Visual Disturbance					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/22/01ISR Number: 3651263-1Report Type:Expedited (15-DaCompany Report #B0096199A  
 Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
23 DAY		Bronchospasm		Simvastatin	C		ORAL
20MG per day				Isosorbide	C		ORAL
25MG per day				Atenolol	C		ORAL
50MG per day				Aspirin	C		ORAL
75MG per day							

Date:01/22/01ISR Number: 3651264-3Report Type:Expedited (15-DaCompany Report #B0096202A  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day 1 DAY		Difficulty In Walking					

Date:01/22/01ISR Number: 3651265-5Report Type:Expedited (15-DaCompany Report #B0096210A  
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Glossodynia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day 61 DAY		Pharyngitis		Allopurinol	C	Glaxo Wellcome	ORAL
100MG per day							

Date:01/22/01ISR Number: 3651266-7Report Type:Expedited (15-DaCompany Report #B0096213A  
 Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Amnesia		Zyban	PS	Glaxo Wellcome	ORAL



Hospitalization - 20MG per day	Fall	Paroxetine	C	ORAL
Initial or Prolonged 50MG per day	Gamma-Glutamyltransferase	Thioridazine	C	ORAL
Disability 1.25MG per day	Increased  Macrocytosis  Spinal Fracture	Oestrone	C	ORAL

Date:01/22/01ISR Number: 3651469-1Report Type:Expedited (15-DaCompany Report #B0094941A  
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - UNKNOWN 1 MON	Gastritis		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged	Haematemesis Rectal Haemorrhage					

Date:01/22/01ISR Number: 3651476-9Report Type:Expedited (15-DaCompany Report #D0013322A  
Age:47 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice	Dry Mouth		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day 1 DAY	Obstructive Airways					
Other	Disorder Panic Disorder Tachycardia Tongue Oedema					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/22/01ISR Number: 3652089-5Report Type:Direct  
Age:24 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG B.I.D.	Hepatitis Toxic Epidermal Necrolysis		Wellbutrin Sr 150mg Glaxowellcome	PS	Glaxowellcome	ORAL
ORAL							

Date:01/22/01ISR Number: 3652380-2Report Type:Expedited (15-DaCompany Report #A0137073A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/ ORAL		Angina Pectoris	Foreign Health  Professional Company Representative	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:01/22/01ISR Number: 3652383-8Report Type:Expedited (15-DaCompany Report #B0093922A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/ORAL		Intervertebral Disc Protrusion  Sciatica	Foreign Health  Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:01/22/01ISR Number: 3652385-1Report Type:Expedited (15-DaCompany Report #B0096058A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/ TWICE		Psoriasis	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

PER DAY/ ORAL

Date:01/22/01ISR Number: 3652387-5Report Type:Expedited (15-DaCompany Report #B0095016A  
Age:25 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - 150 MG/ SEE Initial or Prolonged TEXT/ ORAL Required Intervention to Prevent Permanent Impairment/Damage	Arrhythmia Brain Oedema Cardiac Arrest Coma Convulsion Intentional Misuse Suicide Attempt	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:01/22/01ISR Number: 3652411-XReport Type:Expedited (15-DaCompany Report #B0095928A  
Age:36 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other Required 300 MG (PER Intervention to DAY) ORAL Prevent Permanent Impairment/Damage	Hydrocephalus	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/22/01ISR Number: 3652413-3Report Type:Expedited (15-DaCompany Report #A0136900A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Foreign	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
ORAL		Occupational Exposure To Toxic Agent	Study Health Professional				

Date:01/22/01ISR Number: 3652417-0Report Type:Expedited (15-DaCompany Report #B0095172A  
Age:74 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Cerebral Infarction Cerebrovascular Accident	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE Required PER DAY), Intervention to ORAL Prevent Permanent Impairment/Damage		Urinary Tract Infection	Professional				
				Beclomethasone Dipropion Beclomethasone Dipropion Aspirin Theophylline Amlodipine Besylate	C C C C C		

Date:01/22/01ISR Number: 3652419-4Report Type:Expedited (15-DaCompany Report #B0088136A  
Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Amnesia Asthenia	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL		Circulatory Collapse	Professional				
		Eye Rolling Headache		Birth Control	C		

Loss Of Consciousness  
Pallor  
Salivary Hypersecretion  
Tonic Convulsion

Date:01/22/01ISR Number: 3652746-0Report Type:Expedited (15-DaCompany Report #B0096062A  
Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG ORAL Prevent Permanent Impairment/Damage		Constipation Dermatitis Dermatitis Exfoliative Pruritus	Foreign	Bupropion Hydrochloride Nifedipine Bendrofluazide Nitrazepam Co-Codamol	PS C C C	Glaxo Wellcome Inc	ORAL

Date:01/22/01ISR Number: 3652747-2Report Type:Expedited (15-DaCompany Report #B0096063A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 150 MG (TWICE Initial or Prolonged PER DAY) ORAL		Cerebrovascular Accident	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/22/01ISR Number: 3652748-4Report Type:Expedited (15-DaCompany Report #D0012242A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG (PER DAY) ORAL		Hemiparesis	Foreign Health  Professional	Bupropion Hydrochloride  Iodide Salt Zolpidem Tartrate Diclofenac Birth Control	PS   C C C C	Glaxo Wellcome Inc	ORAL

Date:01/22/01ISR Number: 3652749-6Report Type:Expedited (15-DaCompany Report #A0137061A  
Age:17 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other SEE TEXT, ORAL		Convulsion  Sinus Tachycardia  Suicide Attempt Tremor	Foreign	Wellbutrin Sr  Citalopram Hydrobromide Tablet (Citalopram Hydrobromide)	PS   SS	Glaxo Wellcome Inc	ORAL   ORAL

SEE TEXT,

ORAL

Date:01/22/01ISR Number: 3652889-1Report Type:Expedited (15-DaCompany Report #A0126045A  
Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 150 MG / AT NIGHT / ORAL		Epilepsy  Sudden Death	Health  Professional	Wellbutrin Sr  Carbamazepine Sertraline	PS  C	Glaxo Wellcome Inc	ORAL

Date:01/22/01ISR Number: 3652891-XReport Type:Expedited (15-DaCompany Report #A0134853A  
Age:27 YR Gender:Female I/FU:F

Outcome	PT
Death	Abdominal Adhesions
Hospitalization -	Abdominal Distension
Initial or Prolonged	Abdominal Pain
	Anaemia
	Anorexia
	Anxiety
	Back Pain
	Chest Discomfort
	Chest Pain
	Cough
	Crying
	Cystitis
	Diarrhoea
	Drug Toxicity
	Dyspnoea
	Fatigue
	Feeling Jittery
	Fistula
	Flatulence
	Headache

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Hypoaesthesia Hysterectomy Insomnia	Health				
THREE TIMES		Malaise	Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
PER DAY /		Metastases To Lymph Nodes					
ORAL		Nervousness					
		Neutropenia					
		Night Sweats		Triphasil	C		
		Orthostatic Hypotension		Hypericum	C		
		Overdose		Multivitamin	C		
		Palpitations					
		Paraesthesia					
		Post Coital Bleeding					
		Pyrexia					
		Rhinitis					
		Sinus Tachycardia					
		Squamous Cell Carcinoma Of The Cervix					
		Thyroxine Increased					
		Upper Respiratory Tract Infection					
		Vaginal Haemorrhage					
		Vomiting					
		Weight Decreased					

Date:01/22/01ISR Number: 3652981-1Report Type:Expedited (15-DaCompany Report #A0131639A

Age:73 YR Gender:Female I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/PER		Hyperkalaemia	Health	Zantac 150	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged DAY/ORAL		Intestinal Ischaemia	Professional				
		Megaloblasts Increased		Zyban Tablet - Zyban (Bupropion Hydrochloride)	SS		ORAL
150 MG/PER		Rectal Haemorrhage					
DAY/ORAL							



Theophylline C  
Multivitamin C

Date:01/23/01ISR Number: 3652286-9Report Type:Expedited (15-DaCompany Report #A0137018A  
Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Haemorrhagic Stroke	Health	Wellbutrin	PS	Glaxo Wellcome	
Initial or Prolonged		Professional	Ephedra	SS		

Date:01/24/01ISR Number: 3653401-3Report Type:Expedited (15-DaCompany Report #A0137270A  
Age:36 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Abortion Spontaneous		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:01/24/01ISR Number: 3653418-9Report Type:Expedited (15-DaCompany Report #B0095925A  
Age:38 YR Gender:Male I/FU:I

Outcome  
Life-Threatening  
Hospitalization -

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Hospitalization - Epistaxis  
Initial or Prolonged International Normalised  
Ratio Increased

Zyban	PS	Glaxo Wellcome	ORAL
Bezafibrate	C		
Indapamide	C		
Aspirin	C		
Bisoprolol	C		
Lisinopril	C		
Digoxin	C	Glaxo Wellcome	
Allopurinol	C	Glaxo Wellcome	
Warfarin	C	Glaxo Wellcome	

Date:01/25/01ISR Number: 3654009-6Report Type:Expedited (15-DaCompany Report #B0096574A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	5 DAY	Dizziness		Evening Primrose Oil	C		
		Sedation					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/25/01ISR Number: 3655712-4Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged 150MG DAILY  PO 2 WEEKS  PRIOR TO  ADMISSION	Atrial Fibrillation Blood Magnesium Decreased Hypocalcaemia  Hypovolaemia  Inappropriate  Antidiuretic Hormone  Secretion		Bupropion (Wellbutrin) 150mg Po Q Day	PS		ORAL

Date:01/25/01ISR Number: 3655720-3Report Type:Expedited (15-DaCompany Report #A036897  
 Age:58 YR Gender:Female I/FU:F

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL  300.00 MG  TOTAL: BID: ORA  L	Amnesia Blood Pressure Decreased  Pneumonia  Sedation  Speech Disorder  Toxicologic Test Abnormal	Consumer Health  Professional	Procardia Xl  Bupropion  Gabapentin Atenolol	PS  SS  SS	Pfizer Laboratories Div Pfizer Inc	ORAL  ORAL

Date:01/25/01ISR Number: 3655726-4Report Type:Direct  
 Age:41 YR Gender:Female I/FU:I

Company Report #

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability  40MG DAY ORAL	Agitation Disturbance In Attention  Feeling Jittery		Prozac 20mg Eli Lilly  Wellbutrin Sr 100mg	PS	Eli Lilly	ORAL

100MG DAY	Hypoaesthesia	Glaxo-Wellcome	SS	Glaxo-Wellcome	ORAL
ORAL	Intervertebral Disc				
	Disorder	Xanax	C		
	Muscle Rigidity	Minocycline	C		
	Muscle Spasms				
	Sensory Disturbance				

Date:01/25/01ISR Number: 3655735-5Report Type:Direct Company Report #  
Age:21 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening Hospitalization - EVERYDAY ORAL	Depression Intentional Misuse		Wellbutrin / Glaxowelcome	PS	Glaxowelcome	ORAL
Initial or Prolonged						

Date:01/25/01ISR Number: 3655782-3Report Type:Direct Company Report #  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Required 75 MG DAILY	Pruritus	Consumer	Wellbutrin	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage DAILY ORAL	Urticaria		Yellow #5/#6	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/25/01ISR Number: 3656344-4Report Type:Expedited (15-DaCompany Report #B0095557A

Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG / Prevent Permanent TWICE PER DAY Impairment/Damage / ORAL		Dermographism Urticaria Papular	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:01/25/01ISR Number: 3656356-0Report Type:Expedited (15-DaCompany Report #A0133472A

Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/ TWICE PER DAY / ORAL		Amnesia Asthma Convulsion Decreased Appetite	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Ecchymosis Fall Fatigue Fear Feeling Abnormal Flashback Injury Insomnia Nausea Nervousness Oedema Peripheral Speech Disorder Tremor		Salmeterol + Fluticasone Salbutamol Sulphate	C C		

Date:01/25/01ISR Number: 3656357-2Report Type:Expedited (15-DaCompany Report #B0096064A

Age:46 YR Gender:Male I/FU:I



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Disability

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Foreign	Bupropion			
150 MG/ORAL		Dizziness		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Fall		Paroxetine	C		
		Feeling Abnormal		Thioridazine	C		
		Gamma-Glutamyltransferase		Oestrone	C		
		Increased					
		Loss Of Consciousness					
		Macrocytosis					
		Spinal Compression					
		Fracture					

Date:01/25/01ISR Number: 3656359-6Report Type:Expedited (15-DaCompany Report #B0096199A  
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Angina Pectoris	Foreign	Bupropion			
Required		Bronchospasm		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Intervention to				Simvastatin	C		
150 MG/ ORAL				Isosorbide	C		
Prevent Permanent		Dyspnoea		Atenolol	C		
Impairment/Damage				Aspirin	C		

Date:01/25/01ISR Number: 3656360-2Report Type:Expedited (15-DaCompany Report #B0096202A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State	Foreign	Bupropion			
Required		Convulsion		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Intervention to		Difficulty In Walking					
150 MG/ PER							
Prevent Permanent		Dizziness					
DAY / ORAL		Speech Disorder					
Impairment/Damage							



Date:01/25/01ISR Number: 3656362-6Report Type:Expedited (15-DaCompany Report #B0096210A  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Glossodynia	Foreign	Bupropion			
150 MG / PER		Oral Intake Reduced		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
DAY / ORAL		Pharyngitis					
				Allopurinol	C		

Date:01/25/01ISR Number: 3656401-2Report Type:Expedited (15-DaCompany Report #D0013322A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dry Mouth	Foreign	Bupropion			
150 MG /		Obstructive Airways		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Disorder					
/ ORAL		Panic Reaction					
		Tachycardia					
		Tongue Oedema					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/25/01ISR Number: 3656622-9Report Type:Expedited (15-DaCompany Report #B0094941A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1 MON Initial or Prolonged		Haematemesis	Foreign	Zyban	PS	Glaxo Wellcome Inc	
		Rectal Haemorrhage Vomiting	Health Professional				

Date:01/26/01ISR Number: 3655075-4Report Type:Expedited (15-DaCompany Report #B0095016A

Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 10.5G per day 18 DAY Hospitalization - Initial or Prolonged Other		Brain Oedema Cardiac Arrest Convulsion Encephalopathy Hallucination Hepatic Failure Overdose Suicide Attempt	Health Professional	Zyban	PS	Glaxo Wellcome	ORAL

Date:01/26/01ISR Number: 3655081-XReport Type:Expedited (15-DaCompany Report #B0096570A

Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Variable dose 16 DAY		Blood Creatine Phosphokinase Increased		Zyban	PS	Glaxo Wellcome	ORAL
		Dizziness Dyspnoea Myalgia		Rabeprazole Salbutamol	C C	Glaxo Wellcome	ORAL

Date:01/26/01ISR Number: 3655082-1Report Type:Expedited (15-DaCompany Report #B0096577A

Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	57 DAY	Constipation Depression Hypothyroidism Lethargy Mood Swings Skin Nodule Weight Increased		Zyban  Dydrogesterone	PS  C	Glaxo Wellcome	ORAL  ORAL

Date:01/26/01ISR Number: 3655084-5Report Type:Expedited (15-DaCompany Report #B0096694A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG As Initial or Prolonged directed		Anxiety Blood Pressure Decreased Drug Interaction Face Oedema Paralysis	Consumer	Zyban  Deptran	PS  I	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/26/01ISR Number: 3655122-XReport Type:Expedited (15-DaCompany Report #A0136520A

Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
		Intentional Misuse	Company Representative	Quinidine	SS		
		Ovarian Cyst		Diphenhydramine Hcl	SS		
				Ethanol	SS		

Date:01/26/01ISR Number: 3655785-9Report Type:Direct

Company Report #

Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 150MG QD PO Intervention to Prevent Permanent Impairment/Damage		Skin Discolouration		Wellbutrin (150mg)	PS		ORAL

Date:01/29/01ISR Number: 3655880-4Report Type:Expedited (15-DaCompany Report #B0096583A

Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Eye Pain		Zyban	PS	Glaxo Wellcome	ORAL
		Vision Blurred		Timolol	C		
OPHTHALMIC				Diltiazem	C		ORAL
				Dorzolamide Hydrochloride	C		
OPHTHALMIC				Warfarin	C	Glaxo Wellcome	ORAL
				Gtn	C	Glaxo Wellcome	

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Blister					
per day	12 DAY	Ecchymosis					
		Pruritus					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alcohol Poisoning		Wellbutrin 300 Mg	PS		
QD		Grand Mal Convulsion					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin 75 Mg	PS		
1-AM		Loss Of Consciousness		Prozac	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/29/01ISR Number: 3656533-9Report Type:Expedited (15-DaCompany Report #A0137018A

Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Haemorrhagic Stroke	Health	Wellbutrin	PS	Glaxo Wellcome Inc	
Initial or Prolonged		Professional	Ma-Huang	SS		

Date:01/29/01ISR Number: 3656726-0Report Type:Expedited (15-DaCompany Report #HQ6358124JAN2001

Age:32 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Agitation	Health	Advil	PS	Whitehall	
Initial or Prolonged	Confusional State	Professional			Laboratories Inc Div	
	Convulsion	Other			American Home	
	Coordination Abnormal				Products Corp	ORAL
ORAL						
	Hallucination		Zyban (Amfebutamone			
	Intentional Misuse		Hydrochloride)	SS		ORAL
9 GRAMS, ORAL 1	DAY					
	Tremor					

Date:01/29/01ISR Number: 3656764-8Report Type:Expedited (15-DaCompany Report #B0095016A

Age:25 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Arrhythmia	Foreign	Bupropion			
Hospitalization -	Brain Oedema	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / UNK						
Initial or Prolonged	Cardiac Arrest	Professional				
/ ORAL						
Required	Coma					
Intervention to	Completed Suicide					
Prevent Permanent	Convulsion					
Impairment/Damage	Encephalopathy					
	Hallucination					
	Hepatic Failure					
	Intracranial Pressure					
	Increased					
	Overdose					

Date:01/29/01ISR Number: 3656771-5Report Type:Expedited (15-DaCompany Report #B0096694A  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Anxiety	Foreign	Bupropion			
Initial or Prolonged	Blood Pressure Decreased	Consumer	Hydrochloride	PS	Glaxo Wellcome Inc	
150 MG / AS						
	Drug Interaction					
DIRECTED /						
UNKNOWN	Face Oedema					
	Paralysis		Deptran Tablet (Deptran)	SS		
UNK / AS						
REQUIRED /						
UNKNOWN						

Date:01/29/01ISR Number: 3656772-7Report Type:Expedited (15-DaCompany Report #B0096577A  
Age:39 YR Gender:Female I/FU:I

Outcome	PT
Required	Constipation
Intervention to	Depression
Prevent Permanent	Hypothyroidism
Impairment/Damage	Lethargy
	Mean Cell Volume

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Increased Mood Swings Skin Nodule Tearfulness	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Weight Increased					
/ ORAL				Dydrogesterone	C		

Date:01/29/01ISR Number: 3656774-0Report Type:Expedited (15-DaCompany Report #B0096570A  
Age:42 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG /		Blood Creatine Phosphokinase Increased	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
VARIABLE DOSE			Dizziness					
/ ORAL			Dyspnoea					
			Myalgia		Sodium Rabeprazole Salbutamol Sulphate	C C		

Date:01/30/01ISR Number: 3656821-6Report Type:Expedited (15-DaCompany Report #B0094589A  
Age:57 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2TAB Per day	5	DAY	Abdominal Pain Upper		Zyban	PS	Glaxo Wellcome	ORAL
10MG Per day			Arthralgia		Losec	C		ORAL
12.5MG Twice			Chest Pain		Vioxx	C		ORAL
per day			Cold Sweat					
10MG Per day			Dizziness		Clarityn	C		ORAL



Dry Mouth  
Dry Throat  
Gastrointestinal Ulcer  
Heart Rate Increased  
Hyperhidrosis  
Myalgia  
Nausea  
Vision Blurred

Paracetamol

C

ORAL

Date:01/30/01ISR Number: 3656832-0Report Type:Expedited (15-DaCompany Report #B0096539A

Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Eyelid Oedema Nausea		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG Per day	2 DAY	Throat Irritation					

Date:01/30/01ISR Number: 3656842-3Report Type:Expedited (15-DaCompany Report #B0096982A

Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Nail Disorder		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Rash Generalised					
Other per day							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/30/01ISR Number: 3656843-5Report Type:Expedited (15-DaCompany Report #B0096984A  
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	14 DAY	Psoriasis		Zyban	PS	Glaxo Wellcome	ORAL
				Beclomethasone	C	Glaxo Wellcome	

RESPIRATORY  
 (INHALATION) 100MCG Per  
 day

Date:01/30/01ISR Number: 3657360-9Report Type:Direct Company Report #  
 Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG PO BID Initial or Prolonged		Grand Mal Convulsion		Wellbutrin	PS		ORAL
				Haloperidol	C		
				Ddavp	C		

Date:01/30/01ISR Number: 3657931-XReport Type:Expedited (15-DaCompany Report #2001041398GB  
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Agitation Confusional State	Foreign Health	Motrin	PS	Mcneil Consumer Healthcare	ORAL
		Convulsion Coordination Abnormal	Professional Other	Bupropion(Amfebutamo ne)	SS		

9 G.  
 Hallucination  
 Intentional Misuse  
 Tremor

Date:01/30/01ISR Number: 3658639-7Report Type:Expedited (15-DaCompany Report #B0096116A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required 150 MG/PER		Dizziness Ear Pain	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Intervention to DAY/ORAL		Headache	Professional				
Prevent Permanent Impairment/Damage		Panic Attack Photophobia		Ovysmen	C		

Date:01/30/01ISR Number: 3658641-5Report Type:Expedited (15-DaCompany Report #B0096573A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/ORAL		Epistaxis International Normalised Ratio Increased	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Bezafibrate	C		
				Indapamide	C		
				Aspirin	C		
				Bisoprolol	C		
				Lisinopril	C		
				Digoxin	C		
				Allopurinol	C		
				Warfarin Sodium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/30/01ISR Number: 3658649-XReport Type:Expedited (15-DaCompany Report #B0096574A

Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia Dizziness	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER DAY,ORAL		Sedation		Evening Primrose Oil	C		

Date:01/30/01ISR Number: 3659181-XReport Type:Expedited (15-DaCompany Report #A0137270A

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Study	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
ORAL			Health Professional				

Date:01/30/01ISR Number: 3659824-0Report Type:Expedited (15-DaCompany Report #D0013329A

Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization - UNK / UNK / Initial or Prolonged ORAL		Depression Suicide Attempt	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:01/30/01ISR Number: 3659825-2Report Type:Expedited (15-DaCompany Report #B0095925A

Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization - 150 MG /		Blood Pressure Abnormal Chest Pain	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Initial or Prolonged      Dyspnoea

TWICE PER DAY

Oedema

/ ORAL

Rash Maculo-Papular

Date:01/31/01ISR Number: 3657619-5Report Type:Expedited (15-DaCompany Report #A0137597A

Age:            Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Gingivitis Jaw Disorder		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:01/31/01ISR Number: 3657623-7Report Type:Expedited (15-DaCompany Report #A0138130A

Age:56 YR    Gender:Male      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG Twice		Agitation		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	4 DAY	Anger Angina Pectoris Blood Pressure Increased Cyanosis Skin Discolouration Speech Disorder		Nefazodone	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/31/01ISR Number: 3657624-9Report Type:Expedited (15-DaCompany Report #A0138288A  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	12 WK	Disturbance In Attention Headache Irritability Malaise Mood Swings Sleep Disorder Thyroid Function Test Abnormal		Zyban	PS	Glaxo Wellcome	ORAL

Date:01/31/01ISR Number: 3657650-XReport Type:Expedited (15-DaCompany Report #B0097007A  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		C-Reactive Protein Increased Erysipelas Hypersensitivity Streptococcal Infection		Zyban	PS	Glaxo Wellcome	

Date:01/31/01ISR Number: 3657651-1Report Type:Expedited (15-DaCompany Report #B0097054A  
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 14 DAY Initial or Prolonged 10MG Per day		Aggression Insomnia		Zyban Singulair	PS C	Glaxo Wellcome	 ORAL
600MG per day		Sensory Loss		Theophylline	C		ORAL
INTRA-ARTICULAR 50NG per day		Subarachnoid Haemorrhage		Salmeterol	C	Glaxo Wellcome	
RESPIRATORY				Flutide	C	Glaxo Wellcome	

(INHALATION) 1MG per day

Atrovent C Glaxo Wellcome  
Ventolin C Glaxo Wellcome

Date:01/31/01ISR Number: 3657652-3Report Type:Expedited (15-DaCompany Report #B0097265A  
Age:50 YR Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Atrial Fibrillation		Zyban	PS	Glaxo Wellcome	
UNKNOWN							
Hospitalization - Initial or Prolonged		Chest Pain					

Date:01/31/01ISR Number: 3658029-7Report Type:Periodic Company Report #C2000-0781.01  
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Burning Sensation Haematuria	Consumer	Metoprolol Tartrate	PS	Mylan Pharmaceuticals Inc	ORAL
12.5 MG BID		Oliguria					
ORAL		Pollakiuria Prostatitis		Wellbutrin Glaxo Wellcome	SS	Glaxo Wellcome	ORAL
100 MG QD				Triameterene/Hctz	C		
ORAL				Atorvastatin	C		
				Enalapril	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/31/01ISR Number: 3658131-XReport Type:Periodic  
Age:20 YR Gender:Female I/FU:I

Company Report #A0122448A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL Initial or Prolonged	Pulmonary Embolism	Health  Professional Other	Wellbutrin  Clozapine (Formulation Unknown) (Clozapine) Oral Contraceptive Lithium Salt Clonazepam Thyroxine Sodium	PS  SS C C C C	Glaxo Wellcome Inc	ORAL

Date:01/31/01ISR Number: 3658132-1Report Type:Periodic  
Age:26 YR Gender:Male I/FU:I

Company Report #A0117336A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL Initial or Prolonged	Cognitive Disorder  Depersonalisation Depressed Level Of Consciousness	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL

Date:01/31/01ISR Number: 3658133-3Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0113556A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL Initial or Prolonged	Unevaluable Event	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL

Date:01/31/01ISR Number: 3658134-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0111402A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						



Other ORAL	Convulsion	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
		Professional Company Representative				
Date:01/31/01ISR Number: 3658135-7Report Type:Periodic			Company Report #A0111156A			
Age:36 YR	Gender:Female	I/FU:I				
Outcome Dose Other 150 MG /  TWICE PER DAY  / ORAL	PT  Convulsion  Loss Of Consciousness	Report Source  Health  Professional  Company  Representative	Product  Wellbutrin     Oral Contraceptive Vitamin	Role  PS     C C	Manufacturer  Glaxo Wellcome Inc	Route  ORAL

Date:01/31/01ISR Number: 3658136-9Report Type:Periodic	Company Report #A0110920A					
Age:78 YR	Gender:Female	I/FU:I				
Outcome Dose Hospitalization - ORAL Initial or Prolonged  150 MG /	PT  Grand Mal Convulsion	Report Source  Health  Professional Other	Product  Wellbutrin    Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	Role  PS    SS	Manufacturer  Glaxo Wellcome Inc	Route  ORAL    ORAL

Freedom Of Information (FOI) Report

TWICE PER DAY

/ ORAL

Date:01/31/01ISR Number: 3658137-0Report Type:Periodic Company Report #A0129999A  
 Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Anxiety	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged ORAL		Bipolar Disorder Crying	Professional Other	Risperidone (Risperidone)	SS		ORAL
		Delusion		Human Insulin	C		
		Depression		Alendronate Sodium	C		
		Disturbance In Attention		Aspirin	C		
		Elevated Mood		Multivitamin	C		
		Fatigue		Paracetamol	C		
		Ideas Of Reference		Ibuprofen	C		
		Insomnia		Carmellose Sodium	C		
		Logorrhoea					
		Mania					
		Pressure Of Speech					
		Psychomotor Hyperactivity					
		Psychotic Disorder					
		Suicidal Ideation					
		Thinking Abnormal					

Date:01/31/01ISR Number: 3658138-2Report Type:Periodic Company Report #A0128383A  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1 TABLET / THREE TIMES PER ORAL		Amnesia	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
		Convulsion					
		Drug Interaction					
		Electroencephalogram Abnormal		Amitriptyline Hcl Tablet			

ORAL Hypoglycaemia (Amitriptyline Hcl) SS ORAL  
 Loss Of Consciousness

Date:01/31/01ISR Number: 3658139-4Report Type:Periodic Company Report #A0127311A  
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL Initial or Prolonged	Convulsion	Health  Professional Company Representative	Wellbutrin  Tramadol Hydrochloride	PS  C	Glaxo Wellcome Inc	ORAL

Date:01/31/01ISR Number: 3658140-0Report Type:Periodic Company Report #A0127100A  
 Age:30 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL Initial or Prolonged	Dermatitis  Face Oedema Pruritus Urticaria	Health  Professional	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL

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Freedom Of Information (FOI) Report

Date:01/31/01ISR Number: 3658141-2Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #A0109620A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion Overdose Suicide Attempt	Health Professional	Wellbutrin	PS	Glaxo Wellcome Inc	

Date:01/31/01ISR Number: 3658142-4Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0108729A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100 MG/ TWICE  PER DAY/ ORAL		Abdominal Distension  Abdominal Pain	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
3.75 MG		Convulsion Crying Decreased Appetite		Leuprorelin Acetate Injection (Leuprolide Acetate)	SS		
		Depression Dermatitis Dyspnoea Fatigue Headache Insomnia Mental Disorder Nausea Palpitations Photosensitivity Reaction Pruritus Tremor Urinary Retention Urticaria Weight Decreased		Loratadine	C		

Date:01/31/01ISR Number: 3658143-6Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #A0106624A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other                      Amnesia                      Consumer                      Wellbutrin                      PS                      Glaxo Wellcome Inc  
                                 Convulsion  
                                 Loss Of Consciousness

Date:01/31/01    ISR Number: 3658519-7    Report Type:Periodic                      Company Report #A0126047A  
Age:37 YR    Gender:Female                      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chills	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
75 MG/ TWICE		Confusional State	Professional				
PER DAY/ ORAL		Convulsion					
		Dizziness					
		Fall					
		Muscle Rigidity					
		Tongue Disorder					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/31/01ISR Number: 3658521-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0126044A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 75 MG/ SEE	Back Injury	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged TEXT/ ORAL	Convulsion					
	Deja Vu Difficulty In Walking Hallucination, Olfactory Injury		Clonazepam (Formulation Unknown) (Clonazepam)	SS		
.5 MG/ AS REQUIRED	Insomnia					
	Loss Of Consciousness Pain Road Traffic Accident		Lithium Salt	C		

Date:01/31/01ISR Number: 3658523-9Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #A0125633A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100 MG/ THREE	Convulsion	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged TIMES PER		Professional				
DAY/ ORAL		Company				
		Representative	Ethanol (Formulation Unknown) (Alcohol) Combivent	SS C		

Date:01/31/01ISR Number: 3658525-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0124191A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other ORAL	Convulsion	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
			Sertraline			

Hydrochloride C  
Carbamazepine C

Date:01/31/01ISR Number: 3658527-6Report Type:Periodic Company Report #A0123986A  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
250 MG/ PER							
DAY/ ORAL	2	MON					

Date:01/31/01ISR Number: 3658529-XReport Type:Periodic Company Report #A0108252A  
Age:16 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL		Grand Mal Convulsion	Professional				
		Loss Of Consciousness	Company				
		Mouth Haemorrhage	Representative				
		Muscle Twitching					
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/31/01ISR Number: 3658530-6Report Type:Periodic  
Age:49 YR Gender:Male I/FU:F

Company Report #A0102483A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	450 MG/ PER	Convulsion	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
Other	DAY/ ORAL	Epileptic Aura	Professional				
		Loss Of Consciousness	Company	Aurothiogluucose	C		
		Road Traffic Accident	Representative	Oxaprozin	C		
				Prednisone	C		
				Famotidine	C		

Date:02/01/01ISR Number: 3660626-XReport Type:Expedited (15-DaCompany Report #B0096688A  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required	150 MG/TWICE	Asthenia	Foreign	Bupropion			
Intervention to	PER DAY/ORAL	Blister		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Prevent Permanent	Impairment/Damage	Ecchymosis					
		Pruritus					

Date:02/01/01ISR Number: 3660741-0Report Type:Expedited (15-DaCompany Report #B0096583A  
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required	150 MG /	Eye Pain	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Intervention to	TWICE PER DAY	Vision Blurred					
Prevent Permanent	/ ORAL						
Impairment/Damage				Timolol	C		
				Diltiazem	C		
				Dorzolamide Hcl	C		
				Warfarin Sodium	C		
				Nitroglycerin	C		



Date:02/02/01ISR Number: 3658981-XReport Type:Expedited (15-DaCompany Report #A0130601A  
Age:36 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice Initial or Prolonged per day 2 WK Disability	Arthralgia Blindness Unilateral Brain Hypoxia Cardiac Arrest Cerebrovascular Accident Coma Convulsion Dermatitis Dyspnoea Haemorrhage Lymphadenopathy Medication Error Sarcoidosis		Zyban  No Concurrent Medications	PS  C	Glaxo Wellcome	ORAL

Date:02/02/01ISR Number: 3658983-3Report Type:Expedited (15-DaCompany Report #A0132722A  
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death 150MG Twice per day	Myocarditis		Zyban  Depoprovera	PS  SS	Glaxo Wellcome	ORAL
INTRAMUSCULAR	150MG See					

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Freedom Of Information (FOI) Report

text

Alfalfa	C	ORAL
Calcium	C	ORAL
Activated Charcoal	C	
Kelp	C	ORAL
Potassium	C	ORAL
Vitamin C With Rose		
Hips	C	
Antacid	C	ORAL
Herbal Multivitamin	C	ORAL

MON

Date:02/02/01ISR Number: 3658984-5Report Type:Expedited (15-DaCompany Report #A0135962A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice			Haemorrhage	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	2 WK		Nervous System Disorder				
			Sarcoidosis				

Date:02/02/01ISR Number: 3658986-9Report Type:Expedited (15-DaCompany Report #A0136852A  
Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Keratitis	Zyban	PS	Glaxo Wellcome	ORAL
1 MON			Keratoconjunctivitis				
			Sicca				
			Peripheral Nerve Injury				

Date:02/02/01ISR Number: 3658989-4Report Type:Expedited (15-DaCompany Report #A0138368A  
Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG Twice			Abnormal Behaviour	Wellbutrin	PS	Glaxo Wellcome	ORAL

Initial or Prolonged Condition Aggravated  
per day 6 MON  
Drug Abuser  
Drug Interaction  
Glycosuria  
Syncope

Marijuana SS  
No Concurrent Medications C

Date:02/02/01ISR Number: 3659028-1Report Type:Expedited (15-DaCompany Report #A0129782A

Age:22 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2250MG Single Initial or Prolonged dose 1 DAY		Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day 2 WK		Intentional Misuse		Zyban	SS	Glaxo Wellcome	ORAL
1 DAY				Prenatal Vitamins Marijuana	C C		

Date:02/02/01ISR Number: 3659035-9Report Type:Expedited (15-DaCompany Report #A0138561A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dementia Memory Impairment		Wellbutrin Sr Celexa	PS C	Glaxo Wellcome	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/02/01ISR Number: 3659037-2Report Type:Expedited (15-DaCompany Report #B0084724A  
Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Alopecia		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day 3 WK		Blood Pressure Fluctuation		Allopurinol	C	Glaxo Wellcome	
Disability 300MG Per day 8 YR		Dermatitis Exfoliative Drug Hypersensitivity Eosinophilia Erythema Pigmentation Disorder Psoriasis Rash Erythematous Skin Inflammation Viral Infection					
Other							

Date:02/02/01ISR Number: 3659042-6Report Type:Expedited (15-DaCompany Report #B0096692A  
Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day		Pyrexia		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Vasculitis		Magnyl	C		

Date:02/02/01ISR Number: 3659046-3Report Type:Expedited (15-DaCompany Report #B0097051A  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice		International Normalised Ratio Increased		Wellbutrin	PS	Glaxo Wellcome	ORAL
per day		Memory Impairment		Frusemide	C		ORAL
40MG per day				Warfarin	I	Glaxo Wellcome	ORAL
5MG per day							

Date:02/02/01ISR Number: 3659047-5Report Type:Expedited (15-DaCompany Report #B0097436A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day	12 DAY	Asthma	Zyban	PS	Glaxo Wellcome	ORAL
			Dysphagia				

Date:02/02/01ISR Number: 3659048-7Report Type:Expedited (15-DaCompany Report #B0097575A  
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	300MG Per day	17 DAY	Abdominal Pain Upper	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	40MG per day		Constipation	Isosorbide	C		ORAL
	50MG per day			Atenolol	C		ORAL
	30MG per day			Nifedipine	C		ORAL
	10MG per day			Simvastatin	C		ORAL
	15MG per day			Lansoprazole	C		ORAL
	75MG per day			Aspirin	C		ORAL
				Gaviscon	C		ORAL
				Glyceryl Trinitrate	C	Glaxo Wellcome	ORAL

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Freedom Of Information (FOI) Report

Date:02/02/01ISR Number: 3659049-9Report Type:Expedited (15-DaCompany Report #B0097832A

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Duration Lower Respiratory Tract Infection		Zyban	PS	Glaxo Wellcome	

Date:02/02/01ISR Number: 3660218-2Report Type:Expedited (15-DaCompany Report #B0096984A

Age:53 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150 MG / ORAL	Duration Psoriasis	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
			Beclomethasone Dipropion.	C		

Date:02/02/01ISR Number: 3660219-4Report Type:Expedited (15-DaCompany Report #B0096539A

Age:32 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150 MG / PER DAY / ORAL	Duration Abdominal Pain Eyelid Oedema Nausea	Foreign Study Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:02/02/01ISR Number: 3660249-2Report Type:Expedited (15-DaCompany Report #B0096982A

Age:38 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150 MG / Required TWICE PER DAY	Duration Nail Disorder Rash Generalised	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Intervention to  
/ ORAL  
Prevent Permanent  
Impairment/Damage

Date:02/02/01ISR Number: 3660258-3Report Type:Direct  
Age:36 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema Arthralgia Dermatitis Bullous Malaise Pruritus Urticaria		Wellbutrin	PS		

Date:02/02/01ISR Number: 3660261-3Report Type:Direct  
Age:18 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia		Zyban	PS		ORAL
PO		Paraesthesia Periorbital Oedema Pharyngeal Oedema Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/02/01ISR Number: 3660399-0Report Type:Expedited (15-DaCompany Report #B0094589A  
Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
2 TABLET /		Arthralgia	Health				
PER DAY /		Blood Pressure Increased	Professional				
ORAL		Chest Pain		Omeprazole	C		
		Cold Sweat		Rofecoxib	C		
		Dizziness		Loratadine	C		
		Dry Mouth		Paracetamol	C		
		Dry Throat					
		Feeling Hot And Cold					
		Heart Rate Increased					
		Myalgia					
		Nausea					
		Ulcer					
		Vision Blurred					

Date:02/05/01ISR Number: 3659239-5Report Type:Expedited (15-DaCompany Report #B0097367A  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Zyban	PS	Glaxo Wellcome	ORAL
Disability		Panic Attack					
150MG Per day	18 DAY	Tremor					

Date:02/05/01ISR Number: 3659240-1Report Type:Expedited (15-DaCompany Report #B0097417A  
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination		Zyban	PS	Glaxo Wellcome	ORAL
Other		Pruritus					
150MG Twice							
per day	42 DAY						



50MG Per day	Skin Disorder	Fluconazole	C	ORAL
	Sleep Disorder	Nystatin	C	
	Sleep Talking	Prednisolone	C	
1TAB As				
required		Nitrazepam	C	
10MG At night		Beclomethasone	C	Glaxo Wellcome
800MCG Twice				
per day		Co-Trimoxazole	C	Glaxo Wellcome
480MG Twice				
per day		Combivent	C	
RESPIRATORY				
(INHALATION)	200MCG Four			
times per day		Salbutamol	C	Glaxo Wellcome
1NEB As				
required		Amitriptyline Hydrochloride	C	Glaxo Wellcome
25MG Three				
times per day		Ranitidine	C	Glaxo Wellcome
150MG Twice				
per day		Co-Proxamol Ointment	C C	

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Freedom Of Information (FOI) Report

Date:02/05/01ISR Number: 3659241-3Report Type:Expedited (15-DaCompany Report #B0097573A  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	19 DAY	Influenza Like Illness		Zyban	PS	Glaxo Wellcome	ORAL
100MG Per day		Mouth Ulceration		Atenolol	C		ORAL
2.5MG Per day		Oedema Peripheral		Bendrofluazide	C	Glaxo Wellcome	ORAL
75MG Per day		Rash Generalised		Aspirin	C		ORAL

Date:02/05/01ISR Number: 3661253-0Report Type:Expedited (15-DaCompany Report #A0135962A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG / TWICE PER DAY	Nervous System Disorder Post Procedural	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Haemorrhage					
		Respiratory Disorder					
		Sarcoidosis					

Date:02/05/01ISR Number: 3661257-8Report Type:Expedited (15-DaCompany Report #A0130601A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG / TWICE PER DAY	Arthralgia Blindness Unilateral	Literature Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Disability	TWICE PER DAY	Brain Damage	Professional				
		Cardiac Arrest	Company				
		Cerebrovascular Accident	Representative				
		Chest Pain					
		Chest X-Ray Abnormal					

Coma  
 Convulsion  
 Dermatitis  
 Dyspnoea  
 Haemorrhage  
 Headache  
 Hemiplegia  
 Lung Infiltration  
 Lymphadenopathy  
 Nervous System Disorder  
 Respiratory Arrest

Date:02/05/01ISR Number: 3661276-1Report Type:Expedited (15-DaCompany Report #A0136852A

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eye Disorder	Consumer	Bupropion			
Other		Eye Pain		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL		Keratitis					
		Keratoconjunctivitis					
		Sicca					
		Sensory Loss					
		Viral Infection					

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Freedom Of Information (FOI) Report

Date:02/05/01ISR Number: 3661279-7Report Type:Expedited (15-DaCompany Report #A0138368A  
Age:15 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100 MG / Initial or Prolonged TWICER PER DAY / ORAL	Condition Aggravated Disturbance In Social Behaviour Drug Interaction Movement Disorder Syncope	Health Professional	Wellbutrin Sr Cannabis	PS SS	Glaxo Wellcome Inc	ORAL

Date:02/05/01ISR Number: 3661524-8Report Type:Expedited (15-DaCompany Report #A0138130A  
Age:56 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL 4 DAY	Agitation Anger Angina Pectoris Blood Pressure Increased Cyanosis Speech Disorder	Health Professional	Zyban Nefazodone Hydrochloride	PS C	Glaxo Wellcome Inc	ORAL

Date:02/05/01ISR Number: 3661525-XReport Type:Expedited (15-DaCompany Report #A0137597A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other ORAL	Bone Disorder Gingivitis	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL

Date:02/05/01ISR Number: 3662664-XReport Type:Expedited (15-DaCompany Report #B0097265A  
Age:50 YR Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Atrial Fibrillation	Foreign Health Professional Company Representative	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:02/05/01ISR Number: 3662688-2Report Type:Expedited (15-DaCompany Report #B0097054A  
Age:55 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG Initial or Prolonged		14 DAY	Aggression	Foreign	Zyban	PS	Glaxo Wellcome Inc	
			Headache		Montelukast Sodium	C		
			Insomnia		Theophylline	C		
			Motor Dysfunction		Salmeterol Xinafoate	C		
			Nausea		Fluticasone			
			Subarachnoid Haemorrhage		Propionate	C		
					Ipratropium Bromide	C		
					Salbutamol Sulphate	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/05/01ISR Number: 3662690-0Report Type:Expedited (15-DaCompany Report #B0097007A  
Age:37 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG Initial or Prolonged	C-Reactive Protein Increased Erysipelas Hypersensitivity Joint Swelling Pharyngitis Streptococcal Pyrexia Rash Pruritic	Foreign	Zyban	PS	Glaxo Wellcome Inc	

Date:02/05/01ISR Number: 3662693-6Report Type:Expedited (15-DaCompany Report #A0138288A  
Age:47 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG / TWICE PER DAY / ORAL	Hypothyroidism	Foreign Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Duration						12 WK

Date:02/05/01ISR Number: 3663077-7Report Type:Expedited (15-DaCompany Report #A0132722A  
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 150 MG/ TWICE PER DAY/ ORAL INTRAMUSCULAR	Myocarditis	Foreign Health Professional	Bupropion Hydrochloride Medroxyprogesterone (Medroxyprogesteron e Ace.)	PS SS	Glaxo Wellcome Inc	ORAL
Duration						150 MG/ SEE

TEXT/

INTRAMUSCULAR

Alfalfa C  
Calcium Salt C  
Activated Charcoal C  
Kelp C  
Potassium Salt C  
Ascorbic Acid + Rose C  
Hips C  
Antacid C  
Herbal Vitamins C

Date:02/06/01ISR Number: 3660025-0Report Type:Expedited (15-DaCompany Report #A0128480A

Age:83 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100MG Twice	Hip Fracture		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day 490 DAY	Liver Function Test Abnormal		Xanax	C		ORAL
.125MG Three times per day	Respiratory Failure					
162 DAY	White Blood Cell Count Increased		Pulmocare	C		OTHER
136 DAY			Vicodin	C		ORAL
181 DAY			Digoxin	C	Glaxo Wellcome	ORAL
40MG Four times per day 161 DAY			Megace	C		ORAL
SUBCUTANEOUS 5000IU Twice per day 185 DAY			Heparin	C		
192 DAY			Promod Supplement	C		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

150MG At night	190 DAY	Zantac	C	Glaxo Wellcome	ORAL
20MG Twice per day	102 DAY	Lasix	C		ORAL
20MEQ Twice per day	102 DAY	Kcl	C		ORAL
107 DAY		Feso4	C		ORAL

Date:02/06/01ISR Number: 3660035-3Report Type:Expedited (15-DaCompany Report #A0138183A  
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous		Bupropion	PS	Glaxo Wellcome	ORAL
200MG per day	98 DAY			Prozac	SS		ORAL
				Klonopin	SS		ORAL

Date:02/06/01ISR Number: 3660037-7Report Type:Expedited (15-DaCompany Report #A0138183B  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abortion Spontaneous		Bupropion	PS	Glaxo Wellcome	ORAL
200MG per day	98 DAY			Prozac	SS		ORAL
Congenital Anomaly		Complications Of Maternal Exposure To Therapeutic Drugs Skull Malformation		Klonopin	SS		ORAL

Date:02/06/01ISR Number: 3660051-1Report Type:Expedited (15-DaCompany Report #B0092315A  
Age:37 YR Gender:Female I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aneurysm Haemorrhagic Stroke		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG Twice per day	48 DAY	Headache Nausea					

Date:02/06/01ISR Number: 3660063-8Report Type:Expedited (15-DaCompany Report #B0097298A  
 Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Polydipsia		Zyban	PS	Glaxo Wellcome	ORAL

Date:02/06/01ISR Number: 3660069-9Report Type:Expedited (15-DaCompany Report #B0097835A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN Initial or Prolonged		Crohn'S Disease 7 DAY		Zyban	PS	Glaxo Wellcome	

Date:02/06/01ISR Number: 3660070-5Report Type:Expedited (15-DaCompany Report #B0097960A  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Angina Pectoris		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/06/01ISR Number: 3660394-1Report Type:Direct  
Age:16 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	100MH/BID/ORA	Libido Increased		Bupropion	PS	Teva	ORAL

L

				Paxil	C		
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Date:02/07/01ISR Number: 3660812-9Report Type:Expedited (15-DaCompany Report #A0138544A  
Age:15 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	450MG Per day 1 YR	Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Drug Interaction		Doxycycline	C		
				Acne Medication	C		
TOPICAL				Methylphenidate	I		
18MG Per day							

Date:02/07/01ISR Number: 3661187-1Report Type:Expedited (15-DaCompany Report #B0098045A  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	25MG per day	Blood Pressure Decreased		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Cerebral Ischaemia		Atenolol	C		
		Cerebrovascular Accident		Valsartan	C		
160MG per day		Condition Aggravated		Pravastatin	C		
		Coordination Abnormal					
		Dysarthria					
		Heart Rate Decreased					
		Hypertension					
		Palpitations					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypertensive Crisis	Health Professional	Effexor Xr	PS	Wyeth Ayerst Laboratories	ORAL
300 MG DAILY,			Company				
ORAL			Representative	Wellbutrin - Slow Release (Amfebutamone Hydrochloride)	SS		ORAL
200 MG DAILY,							
ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Asthenia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day 8 DAY		Clonic Convulsion Hyperreflexia Hypoaesthesia Muscle Spasms Pain In Extremity					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/08/01ISR Number: 3661882-4Report Type:Expedited (15-DaCompany Report #A0138874A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Retinal Disorder		Wellbutrin Sr Concurrent Medications	PS  C	Glaxo Wellcome	

Date:02/08/01ISR Number: 3661892-7Report Type:Expedited (15-DaCompany Report #B0091312A

Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Asthma Hypersensitivity		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG Per day	4 DAY	Lower Respiratory Tract Infection Respiratory Distress Urticaria					

Date:02/08/01ISR Number: 3662246-XReport Type:Direct

Age:48 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema		Wellbutrin Sr 150	PS		ORAL
150MG BID PO		Dermatitis					

Date:02/08/01ISR Number: 3662312-9Report Type:Direct

Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Purpura		Wellbutrin 100 Mg	PS		
100MG -> 150		Rash Papular					
Initial or Prolonged		Skin Lesion		Colace	C		
MG BID							

Vasculitis

Remeron	C
Synthroid	C
Hctz	C
Senekot	C
Levaquin	C

Date:02/08/01ISR Number: 3662603-1Report Type:Expedited (15-DaCompany Report #B0092315A  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Haemorrhagic Stroke Headache	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE PER DAY/ORAL		Intracranial Aneurysm Nausea	Professional				

Date:02/08/01ISR Number: 3662610-9Report Type:Expedited (15-DaCompany Report #B0097960A  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Angina Pectoris	Foreign Literature	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/08/01ISR Number: 3662613-4Report Type:Expedited (15-DaCompany Report #B0097835A

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 7 DAY	Crohn'S Disease	Foreign Health  Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:02/08/01ISR Number: 3662636-5Report Type:Expedited (15-DaCompany Report #B0097573A

Age:40 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required Intervention to 150 MG ORAL Prevent Permanent Impairment/Damage	Influenza Like Illness Mouth Ulceration  Oedema Peripheral Rash Generalised	Foreign	Bupropion Hydrochloride  Atenolol Bendrofluazide Aspirin	PS  C C C	Glaxo Wellcome Inc	ORAL

Date:02/08/01ISR Number: 3662640-7Report Type:Expedited (15-DaCompany Report #B0097417A

Age:66 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required 150 MG/ TWICE Intervention to PER DAY/ ORAL Prevent Permanent Impairment/Damage	Hallucination  Pruritus  Skin Disorder Sleep Disorder Sleep Talking	Foreign  Health  Professional Other	Zyban   Fluconazole Nystatin Prednisolone Nitrazepam Beclomethasone Dipropion Co-Trimoxazole Combivent Salbutamol Sulphate Amitriptyline Hcl Ranitidine Hydrochloride	PS   C C C C C C C C C C	Glaxo Wellcome Inc	ORAL

Co-Proxamol C  
Ointment C

Date:02/08/01ISR Number: 3662641-9Report Type:Expedited (15-DaCompany Report #B0097367A  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Headache	Foreign	Bupropion			
150 MG/ PER		Panic Attack	Study	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
DAY/ ORAL		Tremor	Health				
			Professional				

Date:02/08/01ISR Number: 3662859-5Report Type:Expedited (15-DaCompany Report #A0128480A  
Age:83 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Hip Fracture	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
100 MG /		Respiratory Failure	Professional				
Initial or Prolonged		White Blood Cell Count					
TWICE PER DAY		Increased		Alprazolam	C		
/ ORAL				Pulmocare	C		
				Vicodin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Digoxin	C
Megestrol Acetate	C
Heparin	C
Promod	C
Ranitidine	C
Frusemide	C
Postassium Chloride	C
Ferrous Sulfate	C

Date:02/08/01ISR Number: 3662872-8Report Type:Expedited (15-DaCompany Report #A0138544A  
 Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 450 MG/PER		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged DAY/ORAL	1 YR	Drug Interaction	Professional				
				Methylphenidate (Formulation Unknown)	SS		
18 MG/PER DAY				Doxycycline	C		
				Acne Medication	C		

Date:02/08/01ISR Number: 3662874-1Report Type:Expedited (15-DaCompany Report #A0138183A  
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL SEE TEXT		Abortion Spontaneous	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
			Professional	Fluoxetine Hydrochloride Capsule	SS		ORAL
ORAL				Clonazepam	SS		ORAL
ORAL							

Date:02/08/01ISR Number: 3662878-9Report Type:Expedited (15-DaCompany Report #A0138183B  
 Age: Gender: I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abortion Spontaneous	Study	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
ORAL SEE TEXT							
Congenital Anomaly		Complications Of Maternal Exposure To Therapeutic Drugs	Health Professional	Fluoxetine Hydrochloride Capsule	SS		ORAL
ORAL							
ORAL		Skull Malformation		Clonazepam Tablet	SS		ORAL

Date:02/09/01ISR Number: 3662612-2Report Type:Expedited (15-DaCompany Report #B0097298A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Polydipsia	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY)							ORAL

Date:02/09/01ISR Number: 3663043-1Report Type:Expedited (15-DaCompany Report #A0130601A  
Age:36 YR Gender:Female I/FU:F

Outcome  
Hospitalization -  
Initial or Prolonged  
Disability

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 WK	Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
		Blindness Unilateral					
		Brain Damage		No Concurrent Medications	C		
		Cardiac Arrest					
		Cerebrovascular Accident					
		Chest Pain					
		Coma					
		Convulsion					
		Dermatitis					
		Dyspnoea					
		Haemorrhage					
		Headache					
		Hypoxia					
		Inflammation					
		Lung Infiltration					
		Lymphadenopathy					
		Nervous System Disorder					
		Paralysis					
		Respiratory Arrest					
		Sarcoidosis					

Date:02/09/01ISR Number: 3663049-2Report Type:Expedited (15-DaCompany Report #A0139224A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Per day	26 DAY	Abortion Spontaneous		Bupropion	PS	Glaxo Wellcome	ORAL

Date:02/09/01ISR Number: 3663072-8Report Type:Expedited (15-DaCompany Report #B0098183A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Ventricular Fibrillation		Quomem	PS	Glaxo Wellcome	ORAL

Date:02/09/01ISR Number: 3663073-XReport Type:Expedited (15-DaCompany Report #B0098194A  
Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	Sudden Death		Zyban	PS	Glaxo Wellcome	ORAL
	16 DAY						

Date:02/09/01ISR Number: 3663074-1Report Type:Expedited (15-DaCompany Report #B0098198A  
Age:55 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	Sudden Death		Zyban	PS	Glaxo Wellcome	ORAL
	55 DAY						
	2MG Per day			Perindopril	C		ORAL
	286 DAY						
	500MG Twice			Metformin	C		
	per day						
	258 DAY						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/09/01ISR Number: 3663687-7Report Type:Expedited (15-DaCompany Report #B0084724A

Age:54 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY Disability / ORAL Required Intervention to Prevent Permanent Impairment/Damage	Alopecia Blood Pressure Fluctuation Body Temperature Decreased Dermatitis Exfoliative Drug Hypersensitivity Eosinophilia Rash Erythematous Skin Inflammation Viral Infection	Foreign Health Professional	Zyban  Allopurinol	PS  C	Glaxo Wellcome Inc	ORAL

Date:02/09/01ISR Number: 3663700-7Report Type:Expedited (15-DaCompany Report #B0097051A

Age:64 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required 150 MG/TWICE Intervention to PER DAY/ORAL Prevent Permanent Impairment/Damage	International Normalised Ratio Increased Memory Impairment	Foreign	Wellbutrin Sr  Warfarin Sodium (Formulation Unknown) (Warfarin Sodium)  Frusemide	PS  SS  C	Glaxo Wellcome Inc	ORAL  ORAL

Date:02/09/01ISR Number: 3663702-0Report Type:Expedited (15-DaCompany Report #B0096692A

Age:73 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization -	Pyrexia	Foreign	Bupropion			

Initial or Prolonged Vasculitis Hydrochloride PS Glaxo Wellcome Inc ORAL  
150 MG/PER

DAY/ORAL Magnyl C

Date:02/09/01ISR Number: 3663703-2Report Type:Expedited (15-DaCompany Report #B0097832A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG		Bronchospasm Dyspnoea	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
		Lower Respiratory Tract Infection Respiratory Depression					

Date:02/09/01ISR Number: 3663719-6Report Type:Expedited (15-DaCompany Report #B0097436A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150 MG/PER		Asthma Dysphagia	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

DAY/ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/09/01ISR Number: 3663720-2Report Type:Expedited (15-DaCompany Report #B0097575A  
Age:67 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG /PER DAY/ORAL	Abdominal Pain Upper Constipation	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
			Isosorbide	C		
			Atenolol	C		
			Nifedipine	C		
			Simvastatin	C		
			Lansoprazole	C		
			Aspirin	C		
			Gaviscon	C		
			Nitroglycerin	C		

Date:02/09/01ISR Number: 3663841-4Report Type:Expedited (15-DaCompany Report #B0098045A  
Age:56 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Brain Stem Ischaemia Condition Aggravated	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
	Heart Rate Decreased Hypertension Palpitations	Professional	Atenolol Valsartan Pravastatin	C C C		

Date:02/09/01ISR Number: 3664410-2Report Type:Expedited (15-DaCompany Report #A0138561A  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other	Cognitive Disorder Dementia Disturbance In Attention Memory Impairment	Health Professional Company Representative	Wellbutrin Sr Citalopram Hydrobromide	PS C	Glaxo Wellcome Inc	

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2250 MG / SINGLE DOSE / ORAL, 150 MG / PER DAY/ ORAL		Amphetamines Positive Convulsion Electrocardiogram Abnormal Intentional Misuse Suicide Attempt	Health Professional	Bupropion Hydrochloride          Prenatal Cannabis	PS           C C	Glaxo Wellcome Inc	ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day 5 DAY 2MG per day		Angina Pectoris Condition Aggravated Coronary Artery Occlusion		Bupropion      Atenolol + Chlorthalidone Doxazosin	PS      C C	Glaxo Wellcome	ORAL      ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/12/01ISR Number: 3663331-9Report Type:Expedited (15-DaCompany Report #B0097894A  
 Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day	15 DAY	Asthenopia Condition Aggravated	Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
			Gastrointestinal Candidiasis Insomnia Oral Candidiasis Pharyngolaryngeal Pain				

Date:02/12/01ISR Number: 3664055-4Report Type:Periodic Company Report #HQ9426222DEC1999  
 Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	600 MG	OVERDOSE	Health Professional	Effexor	PS	Wyeth Ayerst Laboratories	ORAL
				Prozac (Fluoxetine Hydrochloride)	SS		ORAL
				Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL

AMOUNT, ORAL

AMOUNT, ORAL

AMOUNT, ORAL



Date:02/12/01ISR Number: 3664151-1Report Type:Expedited (15-DaCompany Report #A0138874A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Retinal Disorder	Health Professional Company Representative	Wellbutrin Sr	PS	Glaxo Wellcome Inc	

Date:02/12/01ISR Number: 3664514-4Report Type:Expedited (15-DaCompany Report #B0091312A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Asthma Hypersensitivity	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG PER DAY ORAL		Lower Respiratory Tract Infection Respiratory Distress Urticaria					

Date:02/12/01ISR Number: 3664516-8Report Type:Expedited (15-DaCompany Report #A0138421A  
Age:38 YR Gender:Female I/FU:I

Outcome	PT
Disability	Asthenia Clonic Convulsion Hyperreflexia Hypoaesthesia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Muscle Spasms

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:02/13/01ISR Number: 3664546-6Report Type:Expedited (15-DaCompany Report #B0093437A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 3 WK Initial or Prolonged		Asthenia Confusional State Depression Dizziness Eating Disorder Hyperhidrosis Tremor Visual Disturbance Vomiting		Zyban	PS	Glaxo Wellcome	

Date:02/13/01ISR Number: 3664564-8Report Type:Expedited (15-DaCompany Report #B0098045A  
 Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 25MG per day		Cerebrovascular Accident Condition Aggravated		Zyban Atenolol	PS C	Glaxo Wellcome	ORAL
160MG per day		Coordination Abnormal Dysarthria Heart Rate Decreased Palpitations		Valsartan Pravastatin	C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Twice per day		Tinnitus		Zyban	PS	Glaxo Wellcome	ORAL
		Tremor					
		Visual Disturbance		Eformoterol Salbutamol	C C	Glaxo Wellcome	
RESPIRATORY (INHALATION)							
RESPIRATORY (INHALATION)				Beclomethasone	C	Glaxo Wellcome	
.625MCG Per day				Prempak-C	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice per day		Condition Aggravated		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	17 DAY	Left Ventricular Failure					
20MG per day		Tachycardia		Simvastatin	C		
40MG per day				Pantoprazole	C		
125MCG per day				Digoxin	C	Glaxo Wellcome	
6MG per day				Warfarin	C	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

50MG per day	Losartan	C	
3MG per day	Bumetanide	C	
25MG per day	Gtn	C	Glaxo Wellcome
	Spiroinolactone	C	

Date:02/13/01ISR Number: 3664568-5Report Type:Expedited (15-DaCompany Report #D0013784A  
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Torsade De Pointes		Zyban	PS	Glaxo Wellcome	ORAL
Life-Threatening		Ventricular Fibrillation		Dynacil	C		ORAL
.1MG per day				Digimerck	C		ORAL

Date:02/13/01ISR Number: 3664569-7Report Type:Expedited (15-DaCompany Report #D0013944A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	5 DAY	Initial or Prolonged	Anxiety	Zyban	PS	Glaxo Wellcome	ORAL
			Convulsion				
			Dizziness				
			Paralysis				

Date:02/13/01ISR Number: 3664641-1Report Type:Expedited (15-DaCompany Report #A0139224A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG/PER DAY/ORAL	Abortion Spontaneous	Study	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
			Health				
			Professional				

Date:02/13/01ISR Number: 3664730-1Report Type:Direct  
Age:36 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - Initial or Prolonged	Convulsion		Bupropion	PS		

Date:02/13/01ISR Number: 3665171-3Report Type:Expedited (15-DaCompany Report #A0130601A  
Age:36 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Arthralgia
Disability	Blindness Unilateral
Required	Cardiac Arrest
Intervention to Prevent Permanent Impairment/Damage	Cerebrovascular Accident
	Chest Pain
	Coma
	Convulsion
	Dermatitis
	Dyspnoea
	Haemorrhage
	Headache
	Hemiplegia
	Hypoxia
	Inflammation
	Lung Infiltration

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Lymphadenopathy Nervous System Disorder Respiratory Arrest	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Literature Health	Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG TWICE			Professional				
PER DAY ORAL			Company Representative				

Date:02/13/01ISR Number: 3665225-1Report Type:Expedited (15-DaCompany Report #B0098194A  
Age:21 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Sudden Death	Foreign	Bupropion Hydrochloride	PS		ORAL
Dose							
Death							
150 MG PER							
DAY ORAL							

Date:02/13/01ISR Number: 3665226-3Report Type:Expedited (15-DaCompany Report #B0098198A  
Age:55 YR Gender: I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Sudden Death	Foreign	Bupropion Hydrochloride	PS		ORAL
Dose							
Death							
150 MG PER							
DAY ORAL				Perindopril Metformin	C C		

Date:02/13/01ISR Number: 3665233-0Report Type:Expedited (15-DaCompany Report #B0098183A  
Age: Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose							

Hospitalization - Initial or Prolonged 150 MG ORAL	Ventricular Fibrillation	Foreign Health  Professional	Bupropion Hydrochloride	PS	ORAL
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Date:02/14/01ISR Number: 3664783-0Report Type:Expedited (15-DaCompany Report #B0098813A  
Age:57 YR Gender:Female I/FU:F

Outcome Dose Disability 150MG Twice per day	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Cutaneous Vasculitis Pruritus Rash Erythematous Urticaria		Zyban	PS	Glaxo Wellcome	ORAL

Date:02/14/01ISR Number: 3664784-2Report Type:Expedited (15-DaCompany Report #D0013945A  
Age:31 YR Gender:Female I/FU:I

Outcome Dose Hospitalization - 150MG Twice Initial or Prolonged per day UNKNOWN	Duration 16 DAY	PT	Report Source	Product	Role	Manufacturer	Route
		Arthralgia Erythema Multiforme Haemoglobin Decreased Pruritus		Zyban  Orelox Contraceptives	PS  C C	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/01ISR Number: 3666053-3Report Type:Expedited (15-DaCompany Report #HQ7096312FEB2001

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 75 MG Initial or Prolonged (FREQUENCY UNKNOWN), ORAL	Blood Pressure Increased	Health	Effexor	PS		ORAL
Duration 100 MG 1X PER 1 DAY, ORAL	Drug Interaction	Professional	Wellbutrin - Slow Release (Amfebutamone)	SS		ORAL
			Thyroxine Sodium (Levothyroxine Sodium)	C		
			Amitriptyline (Amitriptyline)	C		
			Atorvastatin Calcium (Atorvastatin Calcium)	C		
			Lansoprazole (Lansoprazole)	C		
			Aspirin (Acetylsalicylic Acid)	C		
			Oxygen (Oxygen)	C		

Date:02/15/01ISR Number: 3665588-7Report Type:Expedited (15-DaCompany Report #A0138400A

Age:31 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 1TAB Twice	Abdominal Pain Upper	Health	Combivir	PS	Glaxo Wellcome	ORAL



Initial or Prolonged per day	Chest Pain	Professional				
100MG Twice	Clostridium Colitis		Wellbutrin	SS	Glaxo Wellcome	ORAL
per day	Condition Aggravated					
200MG Twice	Gallbladder Disorder		Viramune	SS		ORAL
per day	Leukocytoclastic					
	Vasculitis		Azithromycin	C		

Date:02/15/01ISR Number: 3665594-2Report Type:Expedited (15-DaCompany Report #A0139732A  
Age:42 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice	Cyanosis		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	Dermatitis					
7 DAY	Pharyngeal Oedema					
	Urticaria					

Date:02/15/01ISR Number: 3665595-4Report Type:Expedited (15-DaCompany Report #A0139738A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2 YR	Optic Neuritis		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
	Visual Acuity Reduced		Serzone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/15/01ISR Number: 3665601-7Report Type:Expedited (15-DaCompany Report #B0088230A  
Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Acidosis	Foreign	Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - dose 50TAB Single	1 DAY	Coma	Health				
Initial or Prolonged 24TAB Single		Hypotension	Professional	Anadin Extra	SS		ORAL
dose	1 DAY	Intentional Misuse					
		Vomiting					

Date:02/15/01ISR Number: 3665603-0Report Type:Expedited (15-DaCompany Report #B0091526A  
Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Confusional State		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL

Date:02/15/01ISR Number: 3665604-2Report Type:Expedited (15-DaCompany Report #B0091930A  
Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Unevaluable Event		Zyban	PS	Glaxo Wellcome	

Date:02/15/01ISR Number: 3665610-8Report Type:Expedited (15-DaCompany Report #B0098808A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Schizophreniform Disorder		Zyban Quetiapine	PS C	Glaxo Wellcome	

Date:02/15/01ISR Number: 3665611-XReport Type:Expedited (15-DaCompany Report #B0098815A  
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	Myocardial Infarction		Zyban	PS	Glaxo Wellcome	ORAL
		Tongue Oedema		Aspirin	C		ORAL
				Metformin	C		ORAL
	1G Twice per day						
	160MG Per day			Gliclazide	C		ORAL

Date:02/15/01ISR Number: 3665612-1Report Type:Expedited (15-DaCompany Report #B0098877A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day 12 DAY	Gastritis		Zyban	PS	Glaxo Wellcome	ORAL
		Haematemesis					

Date:02/15/01ISR Number: 3665980-0Report Type:Direct Company Report #  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hyperhidrosis Hypoglycaemia Mental Impairment		Bupropion	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/15/01ISR Number: 3665996-4Report Type:Direct  
Age:19 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 3750 MG ONCE Other PO	Grand Mal Convulsion Laboratory Test Abnormal Overdose Road Traffic Accident		Wellbutrin Xr (150mg)	PS		ORAL

Date:02/15/01ISR Number: 3666446-4Report Type:Expedited (15-DaCompany Report #B0097894A  
Age:58 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability 150 MG (PER DAY), ORAL	Asthenopia Gastrointestinal Candidiasis Insomnia Oral Candidiasis Pharyngolaryngeal Pain	Foreign Study Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:02/15/01ISR Number: 3666451-8Report Type:Expedited (15-DaCompany Report #B0089984A  
Age:66 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required Intervention to Prevent Permanent Impairment/Damage 150 MG (TWICE PER DAY) ORAL	Angina Pectoris	Foreign Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)  Atenolol + Chlorthalidone Doxazosin	PS  C C		ORAL

Date:02/16/01ISR Number: 3667536-2Report Type:Expedited (15-DaCompany Report #B0098443A  
Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Tinnitus Tremor Visual Disturbance	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL				Eformoterolc Salbutamol Sulphate Baclomethasone Dipropion Prempak-C	C C C C		

Date:02/16/01ISR Number: 3667988-8Report Type:Expedited (15-DaCompany Report #B0098045A  
Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Pressure Increased Cerebral Ischaemia Cerebrovascular Accident	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS	Glaxo Wellcome Inc	ORAL
ORAL		Coordination Abnormal Dysarthria		Atenolol Valsartan Pravastatin	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/16/01ISR Number: 3668002-0Report Type:Expedited (15-DaCompany Report #B0098449A  
Age:66 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY  / ORAL	Condition Aggravated  Left Ventricular Failure  Tachycardia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Simvastatin	C		
			Pantoprazole	C		
			Digoxin	C		
			Warfarin Sodium	C		
			Losartan Potassium	C		
			Bumetanide	C		
			Nitroglycerin	C		
			Spironolactone	C		

Date:02/16/01ISR Number: 3668008-1Report Type:Expedited (15-DaCompany Report #D0013784A  
Age:67 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death ORAL Life-Threatening	Torsade De Pointes  Ventricular Fibrillation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Fosinopril Sodium	C		
			Digitoxin	C		

Date:02/16/01ISR Number: 3668010-XReport Type:Expedited (15-DaCompany Report #B0093437A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG 3 WK Initial or Prolonged	Asthenia  Confusional State Depression Dizziness Headache Hyperhidrosis Oral Intake Reduced	Foreign  Consumer	Zyban	PS	Glaxo Wellcome Inc	

Tremor  
Visual Disturbance  
Vomiting

Date:02/16/01ISR Number: 3668013-5Report Type:Expedited (15-DaCompany Report #D0013944A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG/ ORAL 5 DAY	Anxiety	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged	Convulsion Dizziness Paralysis	Health Professional				

Date:02/16/01ISR Number: 3668073-1Report Type:Expedited (15-DaCompany Report #B0098813A  
Age:57 YR Gender:Female I/FU:I

Outcome	PT
Disability	Arthralgia Cutaneous Vasculitis Discomfort Pruritus Rash Erythematous Rash Macular

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Freedom Of Information (FOI) Report

Urticaria

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY / ORAL		Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:02/16/01ISR Number: 3668323-1Report Type:Expedited (15-DaCompany Report #D0013945A  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY / ORAL		Arthralgia C-Reactive Protein Increased Difficulty In Walking Erythema Multiforme Haemoglobin Decreased Joint Swelling Pruritus	Foreign Health Professional	Zyban  Cefpodoxime Proxetil Birth Control	PS  C C	Glaxo Wellcome Inc	ORAL

Date:02/19/01ISR Number: 3666538-XReport Type:Expedited (15-DaCompany Report #A0139794A  
Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation Completed Suicide		Zyban	PS	Glaxo Wellcome	ORAL

Date:02/19/01ISR Number: 3666547-0Report Type:Expedited (15-DaCompany Report #B0098962A  
Age: Gender:Male I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Cholelithiasis		Zyban	PS	Glaxo Wellcome	ORAL

Date:02/19/01ISR Number: 3666548-2Report Type:Expedited (15-DaCompany Report #B0098974A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Per day 11 DAY 10MG Twice per day 60MG per day 120MG Three times per day 75MG per day	11 DAY	Agitation Angina Pectoris Palpitations		Zyban Nicorandil Isosorbide Moduretic Diltiazem Aspirin	PS C C C C C	Glaxo Wellcome	ORAL ORAL ORAL ORAL ORAL ORAL

Date:02/19/01ISR Number: 3666549-4Report Type:Expedited (15-DaCompany Report #B0098976A  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Variable dose 75MG Per day		Angina Pectoris Nausea Sleep Disorder		Zyban Aspirin	PS C	Glaxo Wellcome	ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

25MG Per day	Atenolol	C		ORAL
400MCG per day	Gtn	C	Glaxo Wellcome	ORAL
40MG per day	Pravastatin	C		ORAL

Date:02/20/01ISR Number: 3666802-4Report Type:Expedited (15-DaCompany Report #A0123999A  
 Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	13 DAY	Positive	Antinuclear Antibody	Zyban	PS	Glaxo Wellcome	ORAL
		Anxiety Constipation Dizziness Hypoaesthesia Insomnia Paraesthesia Pollakiuria		Vitamin	C		

Date:02/20/01ISR Number: 3666805-XReport Type:Expedited (15-DaCompany Report #A0139668A  
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day	25 DAY	Abnormal Behaviour		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
		Dyskinesia Feeling Jittery Headache Memory Impairment Neck Pain Thinking Abnormal					

Date:02/20/01ISR Number: 3666806-1Report Type:Expedited (15-DaCompany Report #A0139749A  
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG Twice		Abnormal Behaviour		Zyban	PS	Glaxo Wellcome	ORAL
		Astrocytoma		Wellbutrin Sr	SS	Glaxo Wellcome	ORAL
		Body Temperature					
per day		Increased		Klonopin	SS		
6 MON							
		Brain Neoplasm Malignant		Luvox	SS		
6 MON							
		Coma		Triphasil	C		
		Convulsion		Aspirin	C		
		Eye Movement Disorder		Biotin	C		
		Grand Mal Convulsion		Vitamin C	C		
		Loss Of Consciousness		Zinc	C		
		Medication Error		Mvi	C		
		Ocular Hyperaemia					

Date:02/20/01ISR Number: 3666813-9Report Type:Expedited (15-DaCompany Report #B0093945A  
 Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Circulatory Collapse		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	9 DAY	Dizziness					

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Freedom Of Information (FOI) Report

Date:02/20/01ISR Number: 3666818-8Report Type:Expedited (15-DaCompany Report #D0010841A  
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG per day	9 DAY	Amnesia	Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged			Cardiac Arrest				
Other			Epilepsy				

Date:02/20/01ISR Number: 3667605-7Report Type:Periodic Company Report #S00-USA-02047-01  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	40 MG QD PO		Convulsion	Celexa (Citalopram)	PS	Forest Laboratories Inc	ORAL
			Paraesthesia	Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL
	150 MG BID PO			Hydroxyzine	C		
				Alprazolam	C		

Date:02/20/01ISR Number: 3667669-0Report Type:Periodic Company Report #S00-USA-01104-01  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Agitation	Celexa	PS	Forest Laboratories Inc	
			Anxiety	Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL
	150 MG QD PO		Asthenia				
			Depersonalisation				
			Dizziness				
			Drug Withdrawal Syndrome				
			Hypertension				
			Palpitations				
			Sexual Dysfunction				
			Tachycardia				
			Thinking Abnormal				

Date:02/21/01ISR Number: 3667492-7Report Type:Expedited (15-DaCompany Report #A0139670A  
Age:74 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Cardiac Failure		Wellbutrin	PS	Glaxo Wellcome	ORAL
2 MON						
Initial or Prolonged	Congestive		Rocephin	C		
INTRAVENOUS	1G Per day					
	Convulsion		Pepcid	C		
	Loss Of Consciousness		Kcl	C		
			Captopril	C	Glaxo Wellcome	
			Depakote	C		
			Buspar	C		
			Nicoderm Patch	C		
			Seroquel	C		
75MG Per day			Lasix	C		
			Lanoxin	C	Glaxo Wellcome	
			Albuterol	C	Glaxo Wellcome	

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Freedom Of Information (FOI) Report

Date:02/21/01ISR Number: 3667497-6Report Type:Expedited (15-DaCompany Report #B0091711A

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	PT	Report Source	Product	Role	Manufacturer	Route
Initial or Prolonged	Dermatitis Exfoliative		Zyban	PS	Glaxo Wellcome	
	Eye Allergy		Prozac	C		
	Pharyngitis		Corsodyl (Mouthwash)	C		
	Rash Erythematous					
	Rash Pruritic					

Date:02/21/01ISR Number: 3667498-8Report Type:Expedited (15-DaCompany Report #B0093052A

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	PT	Report Source	Product	Role	Manufacturer	Route
Initial or Prolonged	Eye Allergy		Zyban	PS	Glaxo Wellcome	ORAL
	Rash Pruritic		Fluoxetine	C		
			Chlorhexidine	C		

Date:02/21/01ISR Number: 3667499-XReport Type:Expedited (15-DaCompany Report #B0097367A

Age:48 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Disability	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	Headache		Bupropion			
18 DAY	Panic Attack		Hydrochloride	PS	Glaxo Wellcome	ORAL
	Sedation					
	Tremor					

Date:02/21/01ISR Number: 3667500-3Report Type:Expedited (15-DaCompany Report #B0097894A

Age:58 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Disability	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	Asthenopia		Bupropion			
15 DAY	Candidiasis		Hydrochloride	PS	Glaxo Wellcome	ORAL
	Dizziness					
	Insomnia					

Throat Tightness

Date:02/21/01ISR Number: 3668101-3Report Type:Expedited (15-DaCompany Report #2000030406-1

Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Aggression Dizziness		Paxil	PS	Smithkline Beecham Pharmaceuticals	ORAL
10 MILLIGRAMS		Drug Ineffective					
1.0 DAILY		Drug Withdrawal Syndrome					
ORAL	2	WK					
		Emotional Disorder Feeling Abnormal		Wellbutrin (Bupropion Hcl)	SS		ORAL
100		Hyperacusis					
MILLIGRAMS		Irritability					
2.0 DAILY		Malaise					
ORAL		Mood Swings Muscle Spasms		Paxil Smithkline Beecham	SS		
2		Nausea Photophobia		Paxil Smithkline Beecham	SS		ORAL
40 MILLIGRAMS		Sexual Dysfunction					
1.0 DAILY		Temperature Intolerance					
ORAL	2	YR		Paxil Smithkline Beecham	SS		

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20 MILLIGRAMS				Paxil Smithkline Beecham	SS	ORAL
1.0 DAILY						
ORAL	1	MON		Paxil Smithkline Beecham	SS	
				Paxil Smithkline Beecham	SS	ORAL
30 MILLIGRAMS						
1.0 DAILY						
ORAL				Paxil Smithkline Beecham	SS	

Date:02/21/01ISR Number: 3668441-8Report Type:Direct  
Age:28 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Feeling Abnormal		Biaxin 500 Mg	PS		ORAL
1 TAB BID		Feeling Drunk					
ORAL							
		Pain		Wellbutrin Sr	SS		ORAL
1 TAB BID		Paraesthesia					
ORAL							

Date:02/21/01ISR Number: 3668516-3Report Type:Expedited (15-DaCompany Report #A0139738A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Optic Neuritis	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL	2 YR	Visual Acuity Reduced	Consumer	Nefazodone Hydrochloride	C		



Date:02/21/01ISR Number: 3668518-7Report Type:Expedited (15-DaCompany Report #A0139732A  
Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG / TWICE PER DAY / ORAL	Pharyngeal Oedema Rash Generalised Skin Discolouration Urticaria	Foreign Consumer	Zyban Tablet- Zyban (Bupropion Hydrochloride)	PS	Glaxo Wellcome Inc	ORAL

Date:02/21/01ISR Number: 3668520-5Report Type:Expedited (15-DaCompany Report #B0098808A  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required Intervention to Prevent Permanent 150 MG Impairment/Damage	Schizophrenia	Foreign Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride) Quetiapine	PS C	Glaxo Wellcome Inc	

Date:02/21/01ISR Number: 3668522-9Report Type:Expedited (15-DaCompany Report #B0098815A  
Age:62 YR Gender:Male I/FU:I

Outcome	PT
Death	Lethargy Malaise Myocardial Infarction



Hospitalization - Initial or Prolonged	Unevaluable Event	Foreign Health Professional	Zyban Tablet- Zyban (Bupropion Hydrochloride)	PS	Glaxo Wellcome Inc	
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Date:02/21/01ISR Number: 3668536-9Report Type:Expedited (15-DaCompany Report #B0088230A  
Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged 50 TABLET  /SINGLE  DOSE/ORAL		Acidosis Coma Hypotension  Intentional Misuse  Vomiting	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS	Glaxo Wellcome Inc	ORAL
24 TABLET  /SINGLE  DOSE/ORAL				Anadin Extra Tablet (Anadin Extra)	SS		ORAL

Date:02/21/01ISR Number: 3668862-3Report Type:Expedited (15-DaCompany Report #A0138400A  
Age:31 YR Gender:Female I/FU:F

Outcome Hospitalization - Initial or Prolonged	PT Abdominal Pain Upper Aspartate
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Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1 TABLET / TWICE PER DAY / ORAL	Aminotransferase Increased Blood Amylase Decreased Chest Pain Gallbladder Disorder Leukocytoclastic Vasculitis	Health Professional Other	Combivir	PS	Glaxo Wellcome Inc	ORAL
100 MG / TWICE PER DAY / ORAL	Lipase Increased Rash Pruritic		Wellbutrin Tablet (Bupropion Hydrochloride)	SS		ORAL
200 MG / TWICE PER DAY / ORAL			Nevirapine Tablet (Nevirapine)	SS		ORAL
			Azithromycin	C		
Date:02/21/01ISR Number: 3669009-XReport Type:Periodic Company Report #JRFUSA2000001990						
Age:67 YR Gender:Female I/FU:I						

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - SEE IMAGE Initial or Prolonged		Depression Mania Sedation	Consumer Health Professional	Risperdal Risperdal Wellbutrin (Amfebutamone Hydrochloride)	PS SS SS SS	Janssen Research Fdn	ORAL    ORAL
ORAL				Humulin Insulin 70/30 (Humulin 70/30) Fosamax (Alendronate Sodium)	C C		

Aspirin (Acetylsalicylic Acid)	C
Multivitamin (Multiple Vitamins)	C
Acetaminophen (Paracetamol)	C
Motrin (Ibuprofen)	C
Celluvisc (Celluvisc)	C

Date:02/21/01ISR Number: 3701984-7Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #GBR001652

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
15 MG OD		Decreased Appetite Nonspecific Reaction	Consumer Other	Meridia	PS	Knoll Pharmaceutical Co Sub Basf Corp	
15 MG OD				Meridia	SS		
				Wellbutrin	SS		
				Loestrin	C		
				Cipro	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/22/01ISR Number: 3668146-3Report Type:Expedited (15-DaCompany Report #A0123857A  
 Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
MON		Drug Toxicity Emphysema Hepatic Steatosis Overdose Vomiting					

Date:02/22/01ISR Number: 3668149-9Report Type:Expedited (15-DaCompany Report #A0139732A  
 Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Pharyngeal Oedema		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day 8 DAY		Rash Generalised					
Disability 150MG Twice		Skin Discolouration		Zantac	C	Glaxo Wellcome	ORAL
Other per day		Urticaria					

Date:02/22/01ISR Number: 3668152-9Report Type:Expedited (15-DaCompany Report #A0140093A  
 Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 400MG per day 348 DAY		Abortion Spontaneous		Wellbutrin	PS	Glaxo Wellcome	ORAL
UNKNOWN				Ambien	C		

Date:02/22/01ISR Number: 3668153-0Report Type:Expedited (15-DaCompany Report #A0140271A  
 Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Emphysema		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day		Fatigue					
10MG Per day				Singulair	C		
RESPIRATORY				Advair	C	Glaxo Wellcome	
( INHALATION)							
RESPIRATORY				Combivent	C		
( INHALATION)							

Date:02/22/01ISR Number: 3668155-4Report Type:Expedited (15-DaCompany Report #A0140316A  
Age:26 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction		Zyban	PS	Glaxo Wellcome	ORAL

Date:02/22/01ISR Number: 3668170-0Report Type:Expedited (15-DaCompany Report #B0097835A  
Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Crohn'S Disease		Zyban	PS	Glaxo Wellcome	ORAL
1TAB Twice							
Initial or Prolonged							
per day	18 DAY			Omeprazole	C		ORAL
Disability							
20MG per day				Diltiazem	C		ORAL
Other							
120MG Per day				Simvastatin	C		ORAL
10MG per day							
				Kliofem	C		ORAL
				Motival	C		ORAL
				Folic Acid	C		
5MG Per day							

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5MG per day

				Dipyridamole	C		
				Prednisolone	C		

Date:02/22/01ISR Number: 3668171-2Report Type:Expedited (15-DaCompany Report #B0098194A  
 Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	16 DAY	Sudden Death	Zyban	PS	Glaxo Wellcome	ORAL

Date:02/22/01ISR Number: 3668180-3Report Type:Expedited (15-DaCompany Report #B0099236A  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Suicide Attempt	Zyban	PS	Glaxo Wellcome	

Date:02/22/01ISR Number: 3668181-5Report Type:Expedited (15-DaCompany Report #B0099239A  
 Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Twice		Cerebrovascular Accident	Zyban	PS	Glaxo Wellcome	
	per day		Clumsiness				
			Dysgraphia	Cipramil	C		
			Paralysis	Eformoterol	C		
				Hypertension Med	C		

Date:02/22/01ISR Number: 3669958-2Report Type:Expedited (15-DaCompany Report #A0139794A  
 Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Death	Agitation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL	Completed Suicide	Consumer				

Date:02/22/01ISR Number: 3669960-0Report Type:Expedited (15-DaCompany Report #B0098974A  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Agitation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Angina Pectoris					
Intervention to		Palpitations		Nicorandil	C		
Prevent Permanent				Isosorbide	C		
Impairment/Damage				Moduretic	C		
				Diltiazem	C		
				Aspirin	C		

Date:02/22/01ISR Number: 3669975-2Report Type:Expedited (15-DaCompany Report #B0098976A  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability		Angina Pectoris	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG		Nausea					
VARIABLE DOSE		Sleep Disorder					
ORAL				Aspirin	C		
				Atenolol	C		
				Nitroglycerin	C		

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Freedom Of Information (FOI) Report

Pravastatin C

Date:02/22/01ISR Number: 3669984-3Report Type:Expedited (15-DaCompany Report #B0098962A  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL	Cholelithiasis	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:02/23/01ISR Number: 3668794-0Report Type:Expedited (15-DaCompany Report #B0091583A  
Age:46 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death 150MG Per day	Circulatory Collapse		Bupropion Hydrochloride Alcohol	PS C	Glaxo Wellcome	ORAL

Date:02/23/01ISR Number: 3668800-3Report Type:Expedited (15-DaCompany Report #B0098567A  
Age:52 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 15 DAY Initial or Prolonged	Depression Dermatitis Eyelid Oedema Face Oedema Pruritus Psoriasis Tongue Oedema		Zyban	PS	Glaxo Wellcome	ORAL

Date:02/23/01ISR Number: 3668805-2Report Type:Expedited (15-DaCompany Report #B0099220A  
Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Agitation		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	50 DAY	Confusional State					
		Convulsion		Aspirin	C		
		Electroencephalogram Abnormal		Alcohol	C		
		Grand Mal Convulsion					
		Hyperglycaemia					
		Lactic Acidosis					
		Nervousness					
		Salivary Hypersecretion					
		Thrombocythaemia					
		Urinary Incontinence					

Date:02/23/01ISR Number: 3668806-4Report Type:Expedited (15-DaCompany Report #B0099224A  
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day	6 DAY	Depression		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Hallucination					
		Headache					
		Subarachnoid Haemorrhage					

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Freedom Of Information (FOI) Report

Date:02/23/01ISR Number: 3668808-8Report Type:Expedited (15-DaCompany Report #B0099526A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Astrocytoma		Zyban	PS	Glaxo Wellcome	
UNKNOWN							
Hospitalization - Initial or Prolonged		Body Temperature Increased Coma Dysphagia Infection Memory Impairment Urinary Incontinence Vomiting		Prozac	C		

Date:02/23/01ISR Number: 3669200-2Report Type:Direct Company Report #  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Angioneurotic Oedema		Zyban	PS	Glaxo	ORAL
ORAL	2 WK						
Intervention to Prevent Permanent Impairment/Damage		Erythema Multiforme Urticaria		Plendil Centrum Nsaid	C C C		

Date:02/23/01ISR Number: 3669870-9Report Type:Expedited (15-DaCompany Report #A0139749A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abnormal Behaviour Anaplastic Astrocytoma	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL							
		Convulsion Eye Movement Disorder Hyperpyrexia Loss Of Consciousness		Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	SS		ORAL
150 MG/ TWICE PER DAY/ ORAL		Medication Error					
		Ocular Hyperaemia		Clonazepam			

(Formulation	
Unknown)	
(Clonazepam)	SS
Fluvoxamine Maleate	
(Formulation	
Unknown)	
(Fluvoxamine	
Maleate)	SS
Triphasil	C
Aspirin	C
Biotin	C
Ascorbic Acid	C
Zinc Salt	C
M.V.I.	C

Date:02/23/01ISR Number: 3669911-9Report Type:Expedited (15-DaCompany Report #D0010841A  
Age:24 YR Gender:Male I/FU:F

Outcome  
Life-Threatening  
Hospitalization -  
Initial or Prolonged  
Required  
Intervention to

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Prevent Permanent  
Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Cardiac Arrest	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Convulsion Electroencephalogram Abnormal	Health Professional				

Date:02/23/01ISR Number: 3669948-XReport Type:Expedited (15-DaCompany Report #A0123999A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL		Antinuclear Antibody Positive	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Anxiety Constipation Dizziness Hypoaesthesia Insomnia Paraesthesia Pollakiuria		Vitamin	C		

Date:02/23/01ISR Number: 3669950-8Report Type:Expedited (15-DaCompany Report #A0139668A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG/ TWICE PER DAY/ ORAL		Aggression	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
		Feeling Jittery Headache Movement Disorder Neck Pain Thinking Abnormal					

Date:02/23/01ISR Number: 3670092-6Report Type:Expedited (15-DaCompany Report #B0093945A  
Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY		Circulatory Collapse Dizziness	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
/ ORAL							

Date:02/26/01ISR Number: 3669369-XReport Type:Expedited (15-DaCompany Report #A0133472A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Anxiety Asthma Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY (INHALATION) per day	500MCG	Decreased Appetite Twice Ecchymosis		Advair	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)		Fall Insomnia Oedema Peripheral Panic Disorder Tremor		Ventolin	C	Glaxo Wellcome	

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Freedom Of Information (FOI) Report

Date:02/26/01ISR Number: 3669373-1Report Type:Expedited (15-DaCompany Report #A0140573A  
Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Blood Potassium Decreased		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice		Caesarean Section					
Hospitalization -	21 DAY	Cardiovascular Disorder		Prenatal Vitamins	C		
per day		Hepatic Failure		Proventil	C	Glaxo Wellcome	
Initial or Prolonged		Hepatic Steatosis					
		Premature Baby					

Date:02/26/01ISR Number: 3669376-7Report Type:Expedited (15-DaCompany Report #B0098045A  
Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Cerebrovascular Accident		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Coordination Abnormal		Atenolol	C		
25MG per day		Dysarthria		Valsartan	C		
160MG per day		Hypercholesterolaemia		Pravastatin	C		
		Hypertension					
		Palpitations					

Date:02/26/01ISR Number: 3669381-0Report Type:Expedited (15-DaCompany Report #B0099487A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Alopecia Areata		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Costochondritis					
per day	15 DAY	Dyspnoea					
		Hyperventilation					
		Palpitations					



Date:02/26/01ISR Number: 3669382-2Report Type:Expedited (15-DaCompany Report #B0099502A  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Haemorrhagic Stroke		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Ruptured Cerebral					
per day		Aneurysm		Estracombi	C		
TRANSDERMAL							

Date:02/26/01ISR Number: 3669383-4Report Type:Expedited (15-DaCompany Report #B0099512A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Blood Pressure Increased		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day 5 DAY		Cerebral Cyst		Omeprazole	C		
Initial or Prolonged		Congenital Central		Bisoprolol	C		ORAL
20MG per day		Nervous System Anomaly		Enalapril	C		ORAL
Other		Diplopia					
2.5MG per day		Gait Disturbance		Clopidogrel	C		ORAL
10MG Three		Headache		Citalopram	C		ORAL
times per day		Pain					
75MG per day		Vomiting					
20MG per day							

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Freedom Of Information (FOI) Report

Date:02/26/01ISR Number: 3669384-6Report Type:Expedited (15-DaCompany Report #B0099514A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	300MG Three	Feeling Abnormal		Zyban	PS	Glaxo Wellcome	ORAL
	times per day 5	Palpitations					
	DAY	Tearfulness		Prempak C	C		ORAL
		Visual Disturbance					

Date:02/26/01ISR Number: 3669385-8Report Type:Expedited (15-DaCompany Report #B0099522A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	300MG Per day 13	Aggression		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	DAY	Attention					
		Deficit/Hyperactivity					
		Disorder					
		Meningitis					

Date:02/26/01ISR Number: 3669386-XReport Type:Expedited (15-DaCompany Report #B0099529A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	26 DAY	Candidiasis		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged		Dermatitis					
		Malaise					
		Mouth Ulceration					
		Paraesthesia					

Date:02/26/01ISR Number: 3670467-5Report Type:Expedited (15-DaCompany Report #A0139670A  
Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization -	Cardiac Failure	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG / UNK						
Initial or Prolonged	Congestive					
/ ORAL	2 MON					
	Convulsion		Ceftriaxone Sodium	C		
	Loss Of Consciousness		Famotidine	C		
			Potassium Chloride	C		
			Captopril	C		
			Semisodium Valproate	C		
			Buspirone			
			Hydrochloride	C		
			Nicotine	C		
			Quetiapine Fumarate	C		
			Frusemide	C		
			Digoxin	C		
			Salbutamol Sulphate	C		

Date:02/26/01ISR Number: 3670475-4Report Type:Expedited (15-DaCompany Report #A0123857A  
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL		Drug Toxicity					
(DURATION:		Emphysema					
MONTHS)		Hepatic Steatosis					
		Overdose					
		Vomiting					

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Freedom Of Information (FOI) Report

Date:02/26/01ISR Number: 3670480-8Report Type:Expedited (15-DaCompany Report #A0140093A

Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Study	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
SEE TEXT/ ORAL		Complications Of Maternal	Health				
		Exposure To Therapeutic Drugs	Professional	Zolipdem Tartrate	C		

Date:02/26/01ISR Number: 3670499-7Report Type:Expedited (15-DaCompany Report #A103444

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Drug Ineffective	Consumer	Zoloft	PS	Pfizer	
Intervention to		Ecchymosis				Pharmaceuticals Inc	
100.00 MG		Epistaxis					
Prevent Permanent							
TOTAL: DAILY							
Impairment/Damage		Haemorrhagic Stroke		Wellbutrin	SS		
900.00 TOTAL		Syncope		Coumadin	SS		
				Toprol	C		
				Zebeta	C		

Date:02/26/01ISR Number: 3670639-XReport Type:Expedited (15-DaCompany Report #B0091583A

Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Circulatory Collapse	Foreign	Bupropion			
150 MG (PER		Sudden Death	Literature	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
DAY) ORAL				Ethanol	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 25 MG / SEE	Agitation	Health	Lamictal	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged TEXT / ORAL 10 WK Required	Arthralgia	Professional				
Intervention to Prevent Permanent Impairment/Damage ORAL	Dermatitis Erythema Multiforme Headache Hypersensitivity  Influenza Like Illness Nausea Pruritus Stevens-Johnson Syndrome Weight Increased	Company Representative	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)  S-Adenosyl-L-Methion ine	SS   C		ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Dermatitis	Foreign	Zyban	PS	Glaxo Wellcome Inc	
	Dermatitis Exfoliative	Health	Fluoxetine			
	Hypersensitivity	Professional	Hydrochloride	C		
	Pharyngitis		Chlorhexidine			
	Rash Pruritic		Gluconate	C		

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Date:02/26/01ISR Number: 3671309-4Report Type:Expedited (15-DaCompany Report #B0093052A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Eye Allergy	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL							
Initial or Prolonged		Rash Pruritic		Fluoxetine	C		
				Chlorhexidine	C		

Date:02/26/01ISR Number: 3671394-XReport Type:Expedited (15-DaCompany Report #A0140316A

Age:26 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL							

Date:02/26/01ISR Number: 3671396-3Report Type:Expedited (15-DaCompany Report #A0140271A

Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Emphysema	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Fatigue					
PER DAY /							
ORAL				Montelukast Sodium	C		
				Salmeterol			
				+Fluaticasone	C		
				Combivent	C		

Date:02/26/01ISR Number: 3671397-5Report Type:Expedited (15-DaCompany Report #B0099239A

Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Required Intervention to 150 MG /TWICE Prevent Permanent PER DAY Impairment/Damage	Cerebrovascular Accident Clumsiness  Dysgraphia  Facial Palsy	Foreign Health  Professional  Company Representative	Bupropion Hydrochloride   Citalopram Hydrobromide Eformoterol Antihypertensive	PS    C C C	Glaxo Wellcome Inc
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Date:02/26/01ISR Number: 3671399-9Report Type:Expedited (15-DaCompany Report #B0099236A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG		Suicide Attempt	Foreign Health  Professional Company Representative	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:02/26/01ISR Number: 3671428-2Report Type:Expedited (15-DaCompany Report #B0099220A  
Age:41 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Agitation Confusional State Grand Mal Convulsion

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Hyperglycaemia Lactic Acidosis Leukocytosis	Foreign Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Nervousness	Professional	Aspirin	C		
		Salivary Hypersecretion		Ethanol	C		
		Thrombocythaemia					
		Urinary Incontinence					

Date:02/26/01ISR Number: 3671430-0Report Type:Expedited (15-DaCompany Report #B0099224A  
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ PER Initial or Prolonged DAY/ ORAL		Depression	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Hallucination					
		Headache					
		Subarachnoid Haemorrhage					

Date:02/26/01ISR Number: 3671433-6Report Type:Expedited (15-DaCompany Report #B0099526A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - Initial or Prolonged		Astrocytoma	Foreign	Zyban	PS	Glaxo Wellcome Inc	
		Coma	Consumer	Fluoxetine			
		Dysphagia		Hydrochloride	C		
		Eating Disorder					
		Infection					
		Memory Impairment					
		Urinary Incontinence					
		Vomiting					

Date:02/26/01ISR Number: 3671443-9Report Type:Expedited (15-DaCompany Report #A0139732A  
Age:42 YR Gender:Female I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dermatitis Pharyngeal Oedema	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Disability PER DAY/ORAL Required Intervention to Prevent Permanent Impairment/Damage		Skin Discolouration  Urticaria	Professional	  Ranitidine Hydrochloride	  C		

Date:02/26/01ISR Number: 3671453-1Report Type:Expedited (15-DaCompany Report #B0097835A  
Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Crohn'S Disease Intestinal Dilatation	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Disability (TWICE PER Required DAY) ORAL Intervention to Prevent Permanent Impairment/Damage			Professional	  Omeprazole Diltiazem Simvastatin Kliofem Motival Folic Acid Dipyridamole	  C C C C C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Prednisolone C

Date:02/26/01ISR Number: 3671454-3Report Type:Expedited (15-DaCompany Report #B0098194A  
Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (PER DAY)	ORAL		Professional				

Date:02/26/01ISR Number: 3671494-4Report Type:Expedited (15-DaCompany Report #B0097367A  
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Headache Panic Attack	Foreign Study	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (PER DAY)	ORAL	Sedation Tremor	Health Professional				

Date:02/26/01ISR Number: 3671495-6Report Type:Expedited (15-DaCompany Report #B0097894A  
Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Asthenopia Candidiasis	Foreign Study	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (PER DAY)	ORAL	Condition Aggravated Discomfort Dizziness Gastrointestinal Disorder Insomnia Oral Candidiasis Throat Tightness	Health Professional				

Date:02/26/01ISR Number: 3671600-1Report Type:Expedited (15-DaCompany Report #A0140463A  
Age:26 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Euphoric Mood Feeling Abnormal	Foreign Literature	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:02/26/01ISR Number: 3671610-4Report Type:Expedited (15-DaCompany Report #B0098567A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Depression Dermatitis  Difficulty In Walking Eyelid Oedema Face Oedema Pruritus Psoriasis Tongue Oedema	Foreign Consumer	Bupropion Hydrochloride	PS		ORAL



Life-Threatening PO	Blood Pressure Decreased	Health	Zyban	PS	ORAL
Hospitalization - Initial or Prolonged Required	Chest Discomfort Urticaria	Professional			
Intervention to Prevent Permanent Impairment/Damage					

Date:02/28/01ISR Number: 3671094-6Report Type:Expedited (15-DaCompany Report #B0099500A  
 Age:78 YR Gender:Male I/FU:F

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 3 DAY Initial or Prolonged	Atrial Fibrillation	Foreign	Zyban	PS	Glaxo Wellcome	ORAL
5MG Per day Other	Chest Pain	Consumer	Finasteride	C		ORAL
20MG Per day 8 DAY	Dyspnoea		Omeprazole	C		ORAL
	Left Ventricular Failure					

Date:02/28/01ISR Number: 3671096-XReport Type:Expedited (15-DaCompany Report #B0099525A  
 Age:60 YR Gender:Male I/FU:I

Outcome Hospitalization - Initial or Prolonged	PT Benign Respiratory Tract Neoplasm Convulsion Grand Mal Convulsion
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Freedom Of Information (FOI) Report

Muscle Spasms

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300MG Per day	1 WK	Foreign Consumer	Zyban	PS	Glaxo Wellcome	ORAL

Date:02/28/01ISR Number: 3671097-1Report Type:Expedited (15-DaCompany Report #B0099531A  
Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	26 DAY	Arthritis Erythema Marginatum	Foreign Consumer	Zyban	PS	Glaxo Wellcome	ORAL

Date:02/28/01ISR Number: 3671099-5Report Type:Expedited (15-DaCompany Report #B0099756A  
Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death		Alcohol Interaction Drowning	Foreign Consumer	Zyban Alcohol	PS SS	Glaxo Wellcome	ORAL

Date:02/28/01ISR Number: 3671100-9Report Type:Expedited (15-DaCompany Report #B0099804A  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day Initial or Prolonged 7.5MG As required	18 DAY	Angina Pectoris Chest Pain Dyspnoea	Foreign Consumer	Zyban Arthrotec Zopiclone Oestrone Antibiotics	PS C C C C	Glaxo Wellcome	ORAL ORAL ORAL ORAL
8 DAY							

Date:02/28/01ISR Number: 3671101-0Report Type:Expedited (15-DaCompany Report #B0099815A  
Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	Sudden Death	Foreign	Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY			Consumer	Salbutamol	C	Glaxo Wellcome	
(INHALATION)	2PUFF As						
required				Thyroxine	C	Glaxo Wellcome	ORAL
100MCG per							
day				Glyceryl Trinitrate	C	Glaxo Wellcome	ORAL
				Candesartan	C		
4G Per day				Fluoxetine	C		
20MG Per day				Omeprazole	C		
20MG Per day				Frusamide	C		
40MG Per day				Didronel	C		ORAL
1TAB Per day							

Date:02/28/01ISR Number: 3671102-2Report Type:Expedited (15-DaCompany Report #B0099834A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day 2 DAY	Rectal Haemorrhage		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged				Dihydrocodeine	C		ORAL
				Enalapril	C		
2.5MG Per day				Bendrofluazide	C	Glaxo Wellcome	
2.5MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

200MG Twice  
per day

Cimetidine C

Date:02/28/01ISR Number: 3671103-4Report Type:Expedited (15-DaCompany Report #B0099836A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day	14 DAY	Initial or Prolonged	Radiculitis Brachial	Zyban	PS	Glaxo Wellcome	ORAL
			Turner'S Syndrome				

Date:02/28/01ISR Number: 3671104-6Report Type:Expedited (15-DaCompany Report #B0099837A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Per day	6 DAY		Angina Pectoris	Zyban	PS	Glaxo Wellcome	ORAL
			Anxiety	Simvastatin	C		ORAL
			Tachycardia	Indomethacin	C		ORAL
				Gtn	C	Glaxo Wellcome	
				Gaviscon	C		ORAL
				Enalapril	C		ORAL
				Aspirin	C		ORAL

Date:02/28/01ISR Number: 3671849-8Report Type:Periodic Company Report #001-0945-M0001223  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Amnesia Blood Pressure Decreased Laboratory Test Abnormal Pneumonia	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
300 MG (150 MG, BID), PER			Sedation	Zyban (Amfebutamone Hydrochloride)	SS		ORAL



Speech Disorder

ORAL

(Nifedipine) SS  
(Atenolol) SS

Date:02/28/01ISR Number: 3675771-2Report Type:Periodic Company Report #A005685  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression Dermatitis	Health Professional	Zoloft	PS	Pfizer Pharmaceuticals Inc	
25MG TOTAL							
DAILY		Drug Ineffective					
				Wellbutrin Sr	SS		
100 MG TOTAL							
DAILY							

Date:02/28/01ISR Number: 3676137-1Report Type:Periodic Company Report #A012674  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety Cough	Consumer	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
ORAL							
150.00 MG		Drug Ineffective		Bupropion	SS		ORAL
TOTAL:BID:ORA		Dyspnoea					
L		Malaise		Atenolol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/01ISR Number: 3676281-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A028832

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Liver Function Test Abnormal	Health Professional	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
ORAL				Neurontin Wellbutrin	SS SS		

Date:02/28/01ISR Number: 3677286-4Report Type:Periodic  
 Age:47 YR Gender:Female I/FU:I

Company Report #A025527

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coordination Abnormal Dizziness	Health Professional	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
50.00 MG		Drug Ineffective					
TOTAL:DAILY:0		Increased Appetite					
RAL		Libido Decreased Weight Increased		Wellbutrin Sustained Release	SS		ORAL
100.00 MG							
TOTAL:DAILY:0							
RAL							

Date:02/28/01ISR Number: 3677394-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A004071

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction Hypertension	Health Professional	Zoloft	PS	Pfizer Pharmaceuticals Inc	
				Wellbutrin	SS		

Date:02/28/01ISR Number: 3677626-6Report Type:Periodic Company Report #A025366  
Age:38 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 50.00 MG	Grand Mal Convulsion	Consumer	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
TOTAL DAILY						
ORAL 200.00 MG	3 DAY		Wellbutrin Sr	SS		ORAL
TOTAL BID						
ORAL	6 MON					

Date:02/28/01ISR Number: 3677830-7Report Type:Periodic Company Report #9800371  
Age:63 YR Gender:Male I/FU:F

Outcome	PT
Other	Abnormal Dreams Alopecia Constipation Depersonalisation Dizziness Drug Ineffective Drug Withdrawal Syndrome Haematuria Insomnia Neurosis Nocturia Sexual Dysfunction

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Weight Decreased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
100.00 MG		Consumer Health	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
TOTAL:DAILY:0		Professional				
RAL			Wellbutrin	SS		
87.50 MG						
TOTAL:0						
			Aspirin	C		
			Buspar	C		
			Valium	C		

Date:02/28/01ISR Number: 3678092-7Report Type:Periodic Company Report #A040753  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
Other		Apathy Asthenia					
250.00 MG		Drug Ineffective					
TOTAL:DAILY:0		Intentional Misuse					
RAL		Medication Error		Wellbutrin	SS		ORAL
600.00 MG		Nervousness					
TOTAL:0							

Date:02/28/01ISR Number: 3683295-1Report Type:Periodic Company Report #A016975  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Anxiety	Consumer	Zoloft	PS	Pfizer	ORAL
	Asthenia				Pharmaceuticals Inc	
50.00 MG						
	Attention					
TOTAL:DAILY:O						
	Deficit/Hyperactivity					
RAL						
	Disorder		Zidovudine	SS		
	Depression		Sustiva	SS		
	Drug Ineffective		Wellbutrin	SS		
	Drug Level Below		Paxil	C		
	Therapeutic		Serzone	C		
	Headache		Effexor	C		
	Hostility		Norpramin	C		
	Hypercholesterolaemia		Prevacid	C		
	Insomnia					
	Laboratory Test Abnormal					
	Leukopenia					
	Libido Decreased					
	Myalgia					
	Nausea					
	Sedation					
	Testicular Disorder					
	Thinking Abnormal					

Date:03/01/01ISR Number: 3671689-XReport Type:Expedited (15-DaCompany Report #A0138368A  
Age:15 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Condition Aggravated
Initial or Prolonged	Disturbance In Social
Other	Behaviour

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Interaction Glycosuria Movement Disorder	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	6 MON	Syncope	Health Professional	Wellbutrin	PS	Glaxo Wellcome	ORAL
25MG As required				Marijuana Benadryl	SS C		ORAL

Date:03/01/01ISR Number: 3671691-8Report Type:Expedited (15-DaCompany Report #A0140271A  
Age:54 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT Other	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Emphysema	Foreign Health Professional	Zyban	PS	Glaxo Wellcome	ORAL
10MG Per day				Singulair	C		
RESPIRATORY (INHALATION)				Advair	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)				Combivent	C		

Date:03/01/01ISR Number: 3671692-XReport Type:Expedited (15-DaCompany Report #A0141153A  
Age:61 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT Other	Report Source	Product	Role	Manufacturer	Route
150MG Per day	7 DAY	Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL

Date:03/01/01ISR Number: 3671700-6Report Type:Expedited (15-DaCompany Report #B0098813A  
Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Arthralgia	Foreign	Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Cutaneous Vasculitis	Health				
per day	16 DAY						
20MG per day		Rash Macular	Professional	Lisinopril	C		
1TAB per day		Urticaria		Dytenzide	C		

Date:03/01/01ISR Number: 3671706-7Report Type:Expedited (15-DaCompany Report #B0099637A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Intestinal Obstruction		Zyban	PS	Glaxo Wellcome	ORAL

Date:03/01/01ISR Number: 3672077-2Report Type:Expedited (15-DaCompany Report #A0140573A  
Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Blood Potassium Decreased	Study	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Caesarean Section	Health				
Hospitalization - TWICE PER DAY		Complications Of Maternal	Professional				
Initial or Prolonged / ORAL		Exposure To Therapeutic Drugs		Prenatal	C		
		Hepatic Failure		Salbutamol Sulphate	C		
		Hepatic Steatosis					
		Premature Labour					
		Shock					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/01ISR Number: 3672411-3Report Type:Expedited (15-DaCompany Report #B0098045A  
Age:56 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Cerebral Ischaemia Cerebrovascular Accident	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
	Condition Aggravated Heart Rate Decreased Hypertension Palpitations	Professional	Atenolol Valsartan Pravastatin	C C C		

Date:03/01/01ISR Number: 3672412-5Report Type:Expedited (15-DaCompany Report #A0133472A  
Age:37 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG (TWICE PER DAY) ORAL	Anxiety Asthma Convulsion	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
	Decreased Appetite Disturbance In Attention Dizziness Ecchymosis Fatigue Feeling Abnormal Flashback Insomnia Nausea Nervousness Oedema Peripheral Panic Reaction Tremor		Salmeterol + Fluticasone Salbutamol Sulphate	C C		

Date:03/01/01ISR Number: 3672414-9Report Type:Expedited (15-DaCompany Report #B0099487A  
Age:35 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening	Alopecia Areata	Foreign	Bupropion			



150 MG (TWICE PER DAY) ORAL	Costochondritis Dyspnoea Hyperventilation Palpitations	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
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Date:03/01/01ISR Number: 3672415-0Report Type:Expedited (15-DaCompany Report #B0099502A  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Haemorrhagic Stroke Ruptured Cerebral Aneurysm	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL				Estracombi	C		

Date:03/01/01ISR Number: 3672416-2Report Type:Expedited (15-DaCompany Report #B0099514A  
Age:51 YR Gender:Female I/FU:I

Outcome Disability	PT Feeling Abnormal Palpitations
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Sedation Tearfulness Visual Disturbance	Report Source	Product	Role	Manufacturer	Route
300 MG/THREE TIMES PER DAY ORAL			Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Prempak-C	C		

Date:03/01/01ISR Number: 3672417-4Report Type:Expedited (15-DaCompany Report #B0099512A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/PER Required DAY/ORAL		Blood Pressure Increased Cough	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Intervention to Prevent Permanent Impairment/Damage		Diplopia Gait Disturbance Headache Vomiting		Omeprazole Bisoprolol Enalapril Clopidogrel Citalopram	C C C C C		

Date:03/01/01ISR Number: 3672418-6Report Type:Expedited (15-DaCompany Report #B0099522A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG/PER Initial or Prolonged DAY/ORAL		Aggression Attention	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Deficit/Hyperactivity Disorder Meningitis					

Date:03/01/01ISR Number: 3672419-8Report Type:Expedited (15-DaCompany Report #B0099529A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG	Ageusia Dehydration  Dermatitis Dry Mouth Mouth Ulceration Nausea Oral Candidiasis Oral Pain	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:03/02/01ISR Number: 3672194-7Report Type:Expedited (15-DaCompany Report #B0099829A  
Age:38 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG As Initial or Prolonged directed 13 DAY	Arthralgia  Blood Pressure Increased  Heart Rate Increased Hypersensitivity Inflammation Syncope		Zyban	PS	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/01ISR Number: 3672200-XReport Type:Expedited (15-DaCompany Report #D0014384A

Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Zyban	PS	Glaxo Wellcome	ORAL
4 WK		Sedation					

Date:03/02/01ISR Number: 3673056-1Report Type:Expedited (15-DaCompany Report #B0099518A

Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / UNK		Agitation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged / ORAL		Dizziness					
		Dyskinesia					
		Hypertension					
		Nausea					
		Tremor					

Date:03/02/01ISR Number: 3673061-5Report Type:Expedited (15-DaCompany Report #B0099524A

Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ TWICE		Cardiac Disorder	Foreign	Zyban	PS	Glaxo Wellcome Inc	
Initial or Prolonged PER DAY /		Chest Pain	Consumer				

UNKNOWN

Date:03/02/01ISR Number: 3673064-0Report Type:Expedited (15-DaCompany Report #B0099508A

Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Disability	Dizziness	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER						
	Ear Discomfort					
DAY / ORAL			Influenza Vaccine	C		
	Headache					
	Malaise					
	Palpitations					

Date:03/05/01ISR Number: 3673112-8Report Type:Expedited (15-DaCompany Report #A0141089A  
 Age:46 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Suicide Attempt		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day						
Initial or Prolonged			Alcohol	SS		
			Flovent	C	Glaxo Wellcome	
			Serevent	C	Glaxo Wellcome	
			Atrovent	C	Glaxo Wellcome	
			Proventil	C	Glaxo Wellcome	

Date:03/05/01ISR Number: 3673118-9Report Type:Expedited (15-DaCompany Report #B0095116A  
 Age:59 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Cerebrovascular Accident		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Lipids Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/05/01ISR Number: 3674046-5Report Type:Expedited (15-DaCompany Report #A0138368A  
Age:15 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100 MG /TWICE	Abnormal Behaviour	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged PER DAY /ORAL	Drug Dependence	Professional				
Required Intervention to Prevent Permanent Impairment/Damage	Drug Interaction Glycosuria Movement Disorder Syncope		Cannabis (Cannabis) Diphenhydramine Hcl	SS C		

Date:03/05/01ISR Number: 3674449-9Report Type:Expedited (15-DaCompany Report #B0099531A  
Age:28 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required 150 MG /	Arthritis	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Intervention to TWICE PER DAY Prevent Permanent / ORAL Impairment/Damage	Erythema Marginatum					

Date:03/05/01ISR Number: 3674450-5Report Type:Expedited (15-DaCompany Report #B0099837A  
Age:53 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150 MG / PER	Angina Pectoris	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY / ORAL	Anxiety					
	Tachycardia		Simvastatin Indomethacin Nitroglycerin Gaviscon Enalapril Aspirin	C C C C C C		

Date:03/05/01ISR Number: 3674451-7Report Type:Expedited (15-DaCompany Report #B0099836A  
Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300 MG / PER Initial or Prolonged DAY / ORAL	Radiculitis Brachial	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/05/01ISR Number: 3674508-0Report Type:Expedited (15-DaCompany Report #B0099500A  
Age:78 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / PER Initial or Prolonged DAY / ORAL	Atrial Fibrillation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Required Intervention to Prevent Permanent Impairment/Damage	Chest Pain Dyspnoea Left Ventricular Failure		Finasteride Omeprazole	C C		

Date:03/05/01ISR Number: 3674510-9Report Type:Expedited (15-DaCompany Report #B0099834A  
Age:51 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Rectal Haemorrhage	Foreign	Zyban Tablet - Zyban (Bupropion			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

150 MG /  
 PER DAY /  
 ORAL

Hydrochloride) PS ORAL

Dihydrocodeine C  
 Enalapril C  
 Bendrofluazide C  
 Cimetidine C

Date:03/05/01ISR Number: 3674512-2Report Type:Expedited (15-DaCompany Report #B0099815A  
 Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

150 MG /  
 PER DAY /  
 ORAL

Salbutamol Sulphate C  
 Thyroxine Sodium C  
 Nitroglycerin C  
 Candesartan  
 Cilexetil C  
 Fluoxetine C  
 Omeprazole C  
 Frusemide C  
 Disodium Etidronate C

Date:03/05/01ISR Number: 3674514-6Report Type:Expedited (15-DaCompany Report #B0099804A  
 Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Angina Pectoris Dyspnoea	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

300 MG /  
 PER DAY /



ORAL

Arthrotec	C
Zopiclone	C
Oestrone	C
Antibiotics	C

Date:03/05/01ISR Number: 3674520-1Report Type:Expedited (15-DaCompany Report #B0099525A

Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG / PER DAY /		Grand Mal Convulsion Lung Disorder Neoplasm	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL	1 WK						

Date:03/05/01ISR Number: 3674524-9Report Type:Expedited (15-DaCompany Report #B0099756A

Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death UNK / UNK / ORAL		Alcohol Interaction Drowning	Foreign Literature	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Ethanol (Formulation Unknown) (Alcohol)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/05/01ISR Number: 3674586-9Report Type:Expedited (15-DaCompany Report #A0140271A

Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chronic Obstructive	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Airways Disease	Health				
TWICE PER DAY		Exacerbated	Professional				
/ ORAL		Cough		Montelukast Sodium	C		
		Dyspnoea Exacerbated		Salmeterol +			
		Emphysema		Fluticasone	C		
		Fatigue		Combivent	C		

Date:03/05/01ISR Number: 3674588-2Report Type:Expedited (15-DaCompany Report #B0098813A

Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Arthralgia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Cutaneous Vasculitis	Health				
TWICE PER DAY		Pruritus	Professional				
/ ORAL		Rash Erythematous		Lisinopril	C		
		Rash Macular		Dytenzide	C		
		Urticaria					

Date:03/05/01ISR Number: 3674590-0Report Type:Expedited (15-DaCompany Report #A0141153A

Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER			Consumer				
DAY / ORAL							

Date:03/05/01ISR Number: 3674592-4Report Type:Expedited (15-DaCompany Report #B0099637A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL Initial or Prolonged		Intestinal Obstruction	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/06/01ISR Number: 3673996-3Report Type:Expedited (15-DaCompany Report #A0141087A  
Age:88 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100MG Twice per day		Coordination Abnormal Fall Gait Disturbance		Wellbutrin Eskalith Coumadin Procardia Zestril	PS C C C C	Glaxo Wellcome Glaxo Wellcome	ORAL

Date:03/06/01ISR Number: 3674002-7Report Type:Expedited (15-DaCompany Report #B0098194A  
Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Per day 16 DAY		Headache Loss Of Consciousness Sudden Death Vision Blurred		Zyban Chloroquine Diphenhydramine	PS C C	Glaxo Wellcome	ORAL



Date:03/06/01ISR Number: 3674008-8Report Type:Expedited (15-DaCompany Report #B0100066A  
Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	7 DAY	Influenza Like Illness	Zyban	PS	Glaxo Wellcome	
			Pulmonary Embolism				

Date:03/06/01ISR Number: 3674009-XReport Type:Expedited (15-DaCompany Report #B0100069A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	5 DAY	Initial or Prolonged	Chest Pain	Zyban	PS	Glaxo Wellcome	
			Gastrooesophageal Reflux Disease	Sertraline	C		
			Hyperventilation				
			Malaise				
			Nausea				
			Tremor				
			Vomiting				
			Weight Decreased				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/06/01ISR Number: 3674010-6Report Type:Expedited (15-DaCompany Report #B0100194A  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day	9 DAY	Insomnia	Zyban	PS	Glaxo Wellcome	ORAL
	2.5MG per day		Irritability	Bendrofluazide	C	Glaxo Wellcome	ORAL
			Panic Reaction				

Date:03/06/01ISR Number: 3674011-8Report Type:Expedited (15-DaCompany Report #B0100196A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Dehydration Hypersensitivity Pharyngolaryngeal Pain	Zyban	PS	Glaxo Wellcome	

Date:03/06/01ISR Number: 3674012-XReport Type:Expedited (15-DaCompany Report #B0100202A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Hypertensive Crisis	Zyban	PS	Glaxo Wellcome	

Date:03/06/01ISR Number: 3674013-1Report Type:Expedited (15-DaCompany Report #B0100203A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Abnormal Behaviour Chest Discomfort Circulatory Collapse Dyspnoea Panic Attack	Zyban	PS	Glaxo Wellcome	

Date:03/06/01ISR Number: 3674633-4Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
2 QD		Alanine Aminotransferase		Zyban	PS		
		Increased					
		Aspartate					
		Aminotransferase					
		Increased					
		Blood Creatine					
		Phosphokinase Increased					
		Gamma-Glutamyltransferase					
		Increased					
		Monocytosis					
		Oedema Peripheral					
		Serum Ferritin Increased					

Date:03/06/01ISR Number: 3674675-9Report Type:Direct  
Age:32 YR Gender:Female I/FU:I

Company Report #

Outcome	PT
Hospitalization -	Anaphylactoid Reaction
Initial or Prolonged	Chest Discomfort
	Dyspnoea

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Pruritus Urticaria				
Dose	Duration		Report Source	Product	Role	Manufacturer
PO				Bupropion (Wellbutrin)	PS	Wellbutrin
				Clonazepam	C	
				Paroxetine	C	

Date:03/06/01ISR Number: 3675238-1Report Type:Direct  
Age:31 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose						
Death		Convulsion		Wellbutrin Sr Glaxco-Welcome	PS	Glaxco-Welcome

Date:03/06/01ISR Number: 3675388-XReport Type:Expedited (15-DaCompany Report #D0014384A  
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose						
Death		Completed Suicide Sedation	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc
150 MG / UNK			Professional			ORAL
/ ORAL			Other			

Date:03/06/01ISR Number: 3675394-5Report Type:Expedited (15-DaCompany Report #B0099829A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose						
Hospitalization - Initial or Prolonged		Arthralgia Blood Pressure Increased	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc
150 MG / AS			Other			
DIRECTED /		Dizziness				
		Ear Infection				



Heart Rate Increased  
Hypersensitivity

Date:03/07/01ISR Number: 3674830-8Report Type:Expedited (15-DaCompany Report #A0136085A  
Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day		Gastrointestinal Disorder					
60MG Per day		Joint Stiffness		Adalat Xl	C		ORAL
300MG Per day		Pain In Extremity		Synthroid	C	Glaxo Wellcome	ORAL
40MG Per day				Lipitor	C		ORAL
				Hytrin	C		

Date:03/07/01ISR Number: 3674832-1Report Type:Expedited (15-DaCompany Report #A0136852A  
Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eye Pain		Zyban	PS	Glaxo Wellcome	ORAL
1 MON							
		Keratitis					
		Keratoconjunctivitis					
		Sicca					
		Nervous System Disorder					
		Sensory Loss					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/07/01ISR Number: 3674839-4Report Type:Expedited (15-DaCompany Report #A0141476A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	4 WK	Anaemia		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
		Fall Haemoglobin Decreased Hypokalaemia Syncope Tremor		Birth Control Pills	C		

Date:03/07/01ISR Number: 3674841-2Report Type:Expedited (15-DaCompany Report #A0141486A  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Atrial Fibrillation Cardiac Failure Congestive		Zyban	PS	Glaxo Wellcome	ORAL

Date:03/07/01ISR Number: 3674842-4Report Type:Expedited (15-DaCompany Report #B0082488A  
Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG Twice Other per day	31 DAY	Retinal Tear Retinitis Vasculitis		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL

Date:03/07/01ISR Number: 3674845-XReport Type:Expedited (15-DaCompany Report #B0091526A  
Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Confusional State		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL

Date:03/07/01ISR Number: 3674870-9Report Type:Expedited (15-DaCompany Report #B0100045A

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Myocardial Infarction		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged		Myocarditis					

Date:03/07/01ISR Number: 3674871-0Report Type:Expedited (15-DaCompany Report #B0100365A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Renal Failure Acute		Zyban	PS	Glaxo Wellcome	
UNKNOWN							
Initial or Prolonged							

Date:03/07/01ISR Number: 3675547-6Report Type:Expedited (15-DaCompany Report #A0141089A

Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Drug Toxicity	Health	Bupropion			
Initial or Prolonged		Suicide Attempt	Professional	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG PER							

DAY ORAL

Ethanol (Formulation

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Unknown) (Alcohol) C  
 Fluticasone  
 Propionate C  
 Salmeterol Xinafoate C  
 Ipratropium Bromide C  
 Salbutamol Sulphate C

Date:03/07/01ISR Number: 3675724-4Report Type:Expedited (15-DaCompany Report #B0095116A  
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Cerebrovascular Accident Lipids Increased	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/08/01ISR Number: 3675836-5Report Type:Expedited (15-DaCompany Report #B0098815A  
 Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Per day 1 WK		Chest Discomfort		Zyban	PS	Glaxo Wellcome	ORAL
1G Twice per day		Chest Pain Decreased Appetite		Aspirin Metformin	C C		ORAL ORAL
160MG Per day		Dehydration Dermatitis		Gliclazide	C		ORAL
		Dizziness Dry Mouth Dry Skin Dyspnoea Fatigue Headache Insomnia Lethargy Malaise Myocardial Infarction Swelling					

Date:03/08/01ISR Number: 3675838-9Report Type:Expedited (15-DaCompany Report #B0099525A  
Age:60 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Condition Aggravated		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day 8 DAY						
Initial or Prolonged	Convulsion					
	Grand Mal Convulsion					
	Neoplasm					

Date:03/08/01ISR Number: 3675845-6Report Type:Expedited (15-DaCompany Report #B0099781A  
Age:51 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Deep Vein Thrombosis		Zyban	PS	Glaxo Wellcome	
UNKNOWN	4 DAY					
Initial or Prolonged	Malaise		No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/08/01ISR Number: 3675847-XReport Type:Expedited (15-DaCompany Report #B0100349A

Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain		Zyban	PS	Glaxo Wellcome	ORAL
5 DAY		Haemoptysis		Beclomethasone	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)	250MCG per						
day				Gaviscon	C		ORAL
				Loratadine	C		
10MG Per day				Salmeterol	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)	2PUFF Twice						
per day							

Date:03/08/01ISR Number: 3675849-3Report Type:Expedited (15-DaCompany Report #B0100357A

Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Urticaria		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Vasculitis					
per day	22 DAY			Mefenamic Acid	C		ORAL
500MG Three							
times per day							

Date:03/08/01ISR Number: 3675850-XReport Type:Expedited (15-DaCompany Report #B0100358A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Depression		Zyban	PS	Glaxo Wellcome	

Haemorrhagic Stroke

Date:03/08/01ISR Number: 3675852-3Report Type:Expedited (15-DaCompany Report #B0100501A  
 Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Angina Pectoris		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged per day	8 DAY	Dizziness Dry Mouth Nausea					

Date:03/08/01ISR Number: 3675854-7Report Type:Expedited (15-DaCompany Report #B0100520A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Circulatory Collapse Dizziness Hypoaesthesia Musculoskeletal Stiffness Nausea Photophobia Retching		Zyban	PS	Glaxo Wellcome	

Date:03/08/01ISR Number: 3675855-9Report Type:Expedited (15-DaCompany Report #B0100605A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT
Hospitalization - Initial or Prolonged		Convulsion Overdose

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sedation

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
22TAB Per day			Zyban	PS	Glaxo Wellcome	ORAL

Date:03/08/01ISR Number: 3675856-0Report Type:Expedited (15-DaCompany Report #D0014564A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 7TAB		Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
cumulative dose		Gamma-Glutamyltransferase Increased					
		Liver Function Test Abnormal Respiratory Arrest		Amoxicillin	C		

Date:03/08/01ISR Number: 3676575-7Report Type:Expedited (15-DaCompany Report #B0100069A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG		Chest Pain	Foreign	Zyban	PS	Glaxo Wellcome Inc	
Initial or Prolonged		Gastrooesophageal Reflux Disease Hyperventilation Pallor Tremor Vomiting Weight Decreased	Literature Health Professional	Sertraline	C		

Date:03/08/01ISR Number: 3676579-4Report Type:Expedited (15-DaCompany Report #B0100196A  
Age: Gender:Male I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG		Blood Pressure Decreased	Foreign	Zyban	PS	Glaxo Wellcome Inc	
Initial or Prolonged		Dehydration Dermatitis Hypersensitivity Oedema Peripheral	Consumer				

Date:03/08/01ISR Number: 3676581-2Report Type:Expedited (15-DaCompany Report #B0100194A  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 150 MG / PER		Insomnia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Intervention to DAY / ORAL		Irritability	Health				
Prevent Permanent Impairment/Damage		Panic Reaction	Professional	Bendrofluazide	C		

Date:03/08/01ISR Number: 3676583-6Report Type:Expedited (15-DaCompany Report #B0100203A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG		Chest Discomfort	Foreign	Zyban	PS	Glaxo Wellcome Inc	
Initial or Prolonged		Circulatory Collapse Panic Attack Respiratory Disorder	Literature Consumer				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/08/01ISR Number: 3676587-3Report Type:Expedited (15-DaCompany Report #B0100202A  
Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG Initial or Prolonged		Hypertensive Crisis	Foreign	Zyban	PS	Glaxo Wellcome Inc	
			Health Professional Other				

Date:03/08/01ISR Number: 3676599-XReport Type:Expedited (15-DaCompany Report #B0098194A  
Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG/ PER DAY/ ORAL		Headache Loss Of Consciousness Sudden Death Vision Blurred	Foreign Literature Health Professional	Bupropion Hydrochloride Chloroquine Diphenhydramine	PS  C C	Glaxo Wellcome Inc	ORAL

Date:03/08/01ISR Number: 3676623-4Report Type:Expedited (15-DaCompany Report #B0100066A  
Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG (PER DAY)		Influenza Like Illness Pulmonary Embolism	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:03/08/01ISR Number: 3676628-3Report Type:Expedited (15-DaCompany Report #B0100055A  
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Arthralgia	Foreign	Bupropion			

Initial or Prolonged      Blood Pressure Decreased      Consumer      Hydrochloride      PS      Glaxo Wellcome Inc      ORAL  
150 MG ORAL  
  
Dermatitis  
Heart Rate Decreased  
Palpitations  
Vomiting

Date:03/08/01ISR Number: 3676629-5Report Type:Expedited (15-DaCompany Report #B0100063A  
Age:              Gender:Male              I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG		Suicide Attempt	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:03/08/01ISR Number: 3676630-1Report Type:Expedited (15-DaCompany Report #B0099532A  
Age:              Gender:Female              I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Intermittent Claudication Retinal Detachment	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/08/01ISR Number: 3676632-5Report Type:Expedited (15-DaCompany Report #B0099946A  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hearing Impaired	Foreign	Bupropion			
Other		Influenza Like Illness	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL			Professional				

Date:03/08/01ISR Number: 3676686-6Report Type:Expedited (15-DaCompany Report #A0141087A  
Age:88 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Balance Disorder	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Coordination Abnormal	Professional				
100 MG /		Difficulty In Walking					
TWICE PER DAY		Fall		Lithium Carbonate	C		
/ ORAL		Gait Disturbance		Warfarin Sodium	C		
				Nifedipine	C		
				Lisinopril	C		

Date:03/09/01ISR Number: 3676495-8Report Type:Direct Company Report #  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arrhythmia		Bupropion	PS		ORAL
75 MG BID							
ORAL				Amitriptyline Hcl	C		
				Insulin Nph Human	C		
				Insulin Reg Human	C		
				Metformin Hcl	C		
				Zolpidem	C		

Date:03/09/01ISR Number: 3676855-5Report Type:Expedited (15-DaCompany Report #A0141757A  
Age:77 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 200MG Twice			Blood Creatine Increased	Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	18 MON		Blood Urea Increased				
			Dehydration				
			Electrocardiogram Qt Prolonged				
			Nausea				
			Vomiting				

Date:03/09/01ISR Number: 3676856-7Report Type:Expedited (15-DaCompany Report #B0096694A  
Age:66 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice			Anxiety	Zyban	PS	Glaxo Wellcome	
Initial or Prolonged per day			Blood Pressure Decreased				
2UNIT per day			Drug Interaction	Seretide	C	Glaxo Wellcome	
			Face Oedema	Aspirin	C		
			Paraesthesia	Deptran	I		
10MG Three times per day			Paralysis				
			Transient Ischaemic Attack				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/09/01ISR Number: 3676872-5Report Type:Expedited (15-DaCompany Report #B0100339A

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Conversion Disorder		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged	Fall		Atorvastatin	C		
10MG per day	Feeling Abnormal		Enalapril	C		
2.5MG per day	Headache					
	Lethargy					
	Loss Of Consciousness					
	Malaise					
	Movement Disorder					
	Syncope					
	Tremor					

Date:03/09/01ISR Number: 3676873-7Report Type:Expedited (15-DaCompany Report #B0100353A

Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Life-Threatening	Depression		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day 33 DAY	Overdose		Paracetamol	SS		
Hospitalization -						
Initial or Prolonged						

Date:03/09/01ISR Number: 3676878-6Report Type:Expedited (15-DaCompany Report #B0100670A

Age:43 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Condition Aggravated		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day 17 DAY	Mouth Ulceration		Diclofenac	C		ORAL
50MG per day 12 DAY	Vulval Ulceration		Co-Codamol	C		ORAL

Date:03/09/01ISR Number: 3676879-8Report Type:Expedited (15-DaCompany Report #B0100671A

Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day 65 DAY		Left Ventricular Failure	Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -	75MG per day			Amitriptyline	C		ORAL
Initial or Prolonged	20MG per day			Temazepam	C		ORAL
				Human Insulin	C		
SUBCUTANEOUS							

Date:03/09/01ISR Number: 3676880-4Report Type:Expedited (15-DaCompany Report #B0100677A  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice		Agitation	Zyban	PS	Glaxo Wellcome	ORAL
	per day		Back Pain				
			Insomnia				

Date:03/09/01ISR Number: 3676882-8Report Type:Expedited (15-DaCompany Report #B0100678A  
Age:45 YR Gender:Female I/FU:I

Outcome	PT
Life-Threatening	Acute Psychosis
Hospitalization -	Bite
Initial or Prolonged	Convulsion
	Muscle Contractions
	Involuntary

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oxygen Saturation Decreased					
		Respiratory Depression					
		Tongue Disorder		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice per day	18 DAY	Urinary Incontinence					
800MG per day	130 DAY			Lithium	C		ORAL
150MG per day	170 DAY			Clomipramine	C		ORAL
1TAB per day				Folic Acid	C		ORAL

Date:03/09/01ISR Number: 3676883-XReport Type:Expedited (15-DaCompany Report #B0100682A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Glucose Decreased		Zyban	PS	Glaxo Wellcome	
Hospitalization - 150MG Twice Initial or Prolonged per day		Circulatory Collapse					
		Dizziness Hypoaesthesia Visual Disturbance		Coffee	C		ORAL

Date:03/09/01ISR Number: 3676884-1Report Type:Expedited (15-DaCompany Report #B0100684A  
 Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Zyban	PS	Glaxo Wellcome	
				Lovan	C		
				Alcohol	C		

Date:03/12/01ISR Number: 3678075-7Report Type:Expedited (15-DaCompany Report #A0141841A  
 Age:23 YR Gender:Male I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
2 WK							

Date:03/12/01ISR Number: 3678079-4Report Type:Expedited (15-DaCompany Report #B0100043A  
 Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coronary Artery Disease		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day		Myocardial Infarction		Carbimazole	C		ORAL
5MG Per day				Enalapril	C		ORAL
2.5MG Per day				Clarithromycin	C		ORAL
1000MG Per							
day	8 DAY			Doxycycline	C		ORAL
100MG Per day	8 DAY						

Date:03/12/01ISR Number: 3678083-6Report Type:Expedited (15-DaCompany Report #B0100749A  
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Fibrinogen		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day	14 DAY	Increased					
		Haematoma					
		Mean Cell Volume Abnormal					
		Red Blood Cell Count					
		Decreased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/12/01ISR Number: 3678085-XReport Type:Expedited (15-DaCompany Report #B0100755A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Zyban	PS	Glaxo Wellcome	

Date:03/12/01ISR Number: 3678086-1Report Type:Expedited (15-DaCompany Report #B0100763A  
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged		Anaphylactic Reaction Aphasia Circulatory Collapse Dermatitis Respiratory Arrest Tremor		Zyban	PS	Glaxo Wellcome	

Date:03/12/01ISR Number: 3680290-3Report Type:Expedited (15-DaCompany Report #A0141476A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL 4 WK		Anaemia Convulsion Fall Hypokalaemia Syncope	Health Professional	Wellbutrin Sr Oral Contraceptive	PS C	Glaxo Wellcome Inc	ORAL

Date:03/12/01ISR Number: 3680291-5Report Type:Expedited (15-DaCompany Report #A0136852A  
Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG/ ORAL		Eye Infection Viral Keratitis Keratoconjunctivitis	Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/12/01ISR Number: 3680666-4Report Type:Expedited (15-DaCompany Report #A0136085A  
 Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Arthritis	Health				
PER DAY ORAL		Gastrointestinal Disorder	Professional	Nifedipine	C		
		Joint Stiffness		Thyroxine Sodium	C		
				Atorvastatin Calcium	C		
				Terazosin			
				Hydrochloride	C		

Date:03/12/01ISR Number: 3680670-6Report Type:Expedited (15-DaCompany Report #B0091526A  
 Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Confusional State	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL			Health				
Initial or Prolonged			Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/12/01ISR Number: 3680689-5Report Type:Expedited (15-DaCompany Report #B0100365A

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Renal Failure Acute	Foreign Health Professional Other	Zyban	PS	Glaxo Wellcome Inc	

Date:03/12/01ISR Number: 3680693-7Report Type:Expedited (15-DaCompany Report #A0141486A

Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - ORAL Initial or Prolonged	Atrial Fibrillation  Cardiac Failure Congestive	Foreign  Health Professional Other	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/12/01ISR Number: 3680695-0Report Type:Expedited (15-DaCompany Report #B0100045A

Age: Gender:I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG Initial or Prolonged	Myocarditis	Foreign  Health Professional Other	Zyban	PS	Glaxo Wellcome Inc	

Date:03/12/01ISR Number: 3680834-1Report Type:Expedited (15-DaCompany Report #B0082488A

Age:32 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ ORAL	Maculopathy  Retinal Exudates	Foreign  Study	Zyban	PS	Glaxo Wellcome Inc	ORAL

Required	Retinal Oedema	Health
Intervention to	Retinal Tear	Professional
Prevent Permanent	Retinal Vasculitis	
Impairment/Damage	Retinitis	
	Visual Disturbance	

Date:03/12/01ISR Number: 3680843-2Report Type:Expedited (15-DaCompany Report #A0140463A  
Age:27 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia Cardiac Arrest	Foreign Literature	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL	10 DAY	Euphoric Mood Feeling Abnormal Heart Rate Increased Palpitations Psychomotor Hyperactivity					

Date:03/13/01ISR Number: 3679122-9Report Type:Expedited (15-DaCompany Report #A0134853A  
Age:27 YR Gender:Female I/FU:F

Outcome  
Death  
Hospitalization -

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Freedom Of Information (FOI) Report

Initial or Prolonged

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Three		Abdominal Distension		Wellbutrin	PS	Glaxo Wellcome	ORAL
times per day	6	Abdominal Pain					
1TAB Per day		Abscess		Triphasil	C		ORAL
	DAY	Anaemia		St. John'S Wort	C		
		Anorexia		Vitamins	C		
		Anxiety					
		Back Pain					
		Cellulitis					
		Cervix Carcinoma					
		Chest Pain					
		Constipation					
		Crepitations					
		Cystitis					
		Depression					
		Drug Toxicity					
		Dyspnoea					
		Embolism					
		Fistula					
		Hypoaesthesia					
		Hypokalaemia					
		Hyponatraemia					
		Hypovolaemia					
		Insomnia					
		Malaise					
		Menorrhagia					
		Muscle Spasms					
		Nervousness					
		Neutropenia					
		Night Sweats					
		Oedema					
		Oral Candidiasis					
		Orthostatic Hypotension					
		Palpitations					
		Paraesthesia					
		Pollakiuria					
		Post Coital Bleeding					
		Postoperative Adhesion					
		Proteinuria					
		Pyrexia					
		Rectal Haemorrhage					

Respiratory Disorder  
Rhinitis  
Sepsis  
Small Intestinal  
Obstruction  
Squamous Cell Carcinoma  
Tachycardia  
Urinary Tract Infection  
Urine Analysis Abnormal  
Weight Decreased

Date:03/13/01ISR Number: 3679125-4Report Type:Expedited (15-DaCompany Report #A0138336A  
Age:68 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Anxiety
Initial or Prolonged	Blood Creatine

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Phosphokinase Increased  
Dyspnoea  
Rhabdomyolysis

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day			Wellbutrin	PS	Glaxo Wellcome	ORAL
25MG Per day			Benazepril Vioxx	C C		ORAL ORAL

Date:03/13/01ISR Number: 3679126-6Report Type:Expedited (15-DaCompany Report #A0141472A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Eye Disorder		Zyban	PS	Glaxo Wellcome	ORAL

Date:03/13/01ISR Number: 3679127-8Report Type:Expedited (15-DaCompany Report #A0141548A  
Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Chest Pain		Zyban	PS	Glaxo Wellcome	ORAL
UNKNOWN per day	10MG Twice	Dyspnoea Fatigue Haemoptysis		Accupril	C		
		Lower Respiratory Tract Infection					

Date:03/13/01ISR Number: 3679133-3Report Type:Expedited (15-DaCompany Report #B0098808A  
Age:38 YR Gender:Male I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Zyban	PS	Glaxo Wellcome	
150MG Twice		Depression					
per day				Quetiapine	C		

Date:03/13/01ISR Number: 3679138-2Report Type:Expedited (15-DaCompany Report #B0100348A  
 Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		C-Reactive Protein		Zyban	PS	Glaxo Wellcome	ORAL
150MG Three		Increased					
times per day	9	DAY		Catapressan	C		
		Confusional State		Penicillin	C	Glaxo Wellcome	
		Drug Dependence		Netromycin			
		Drug Withdrawal Syndrome		(Netilmicin)	C		
		Dysphonia		Haloperidol	C		
		Hallucination, Visual		Morphine Chloride	C		
		Psychotic Disorder					
		Restlessness					
		Sepsis					

Date:03/13/01ISR Number: 3679148-5Report Type:Expedited (15-DaCompany Report #B0100733A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Akinesia		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day	12	DAY					
100MG per day		Dysarthria		Danazol	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/13/01ISR Number: 3679149-7Report Type:Expedited (15-DaCompany Report #B0100762A  
Age:19 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Zyban	PS	Glaxo Wellcome	
		Intentional Misuse		Ibuprofen	SS		
		Suicide Attempt		Alcohol	SS		

Date:03/13/01ISR Number: 3679151-5Report Type:Expedited (15-DaCompany Report #B0100897A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Haemorrhagic Stroke		Zyban	PS	Glaxo Wellcome	
150MG As		Headache					
directed							

Date:03/13/01ISR Number: 3679152-7Report Type:Expedited (15-DaCompany Report #B0101145A  
Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Circulatory Collapse		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day 9	DAY	Dizziness					
Initial or Prolonged							

Date:03/13/01ISR Number: 3679860-8Report Type:Direct Company Report #  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Movement Disorder		Wellbutrin Sr			
		Oedema Mouth		(150mg)	PS		
1T BID	17 DAY	Oedema Peripheral		Motrin	C		
		Rash Maculo-Papular		Adalat	C		
		Tenderness		Prinivil	C		
		Tremor					

Date:03/13/01ISR Number: 3680170-3Report Type:Expedited (15-DaCompany Report #B0100677A  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation Back Pain	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Sleep Disorder					
PER DAY/ORAL		Sudden Death					

Date:03/13/01ISR Number: 3680172-7Report Type:Expedited (15-DaCompany Report #B0100684A  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
150 MG				Fluoxetine Hydrochloride	C		
				Ethanol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/13/01ISR Number: 3680173-9Report Type:Expedited (15-DaCompany Report #B0100682A

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Blindness Transient	Foreign	Bupropion			
Initial or Prolonged	Blood Glucose Decreased	Consumer	Hydrochloride	PS	Glaxo Wellcome Inc	
	Circulatory Collapse		Coffee	C		
	Dizziness					
	Hypoaesthesia					

Date:03/13/01ISR Number: 3680175-2Report Type:Expedited (15-DaCompany Report #B0100678A

Age:45 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Dyskinesia	Foreign	Bupropion			
Hospitalization -	Oxygen Saturation		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE						
Initial or Prolonged	Abnormal					
PER DAY/ORAL						
	Respiratory Depression		Lithium Salt	C		
	Urinary Incontinence		Clomipramine Hcl	C		
			Folic Acid	C		

Date:03/13/01ISR Number: 3680176-4Report Type:Expedited (15-DaCompany Report #B0100339A

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Conversion Disorder	Foreign	Bupropion			
Initial or Prolonged	Fall	Consumer	Hydrochloride	PS	Glaxo Wellcome Inc	
	Headache		Atorvastatin Calcium	C		
	Lethargy		Enalapril	C		
	Loss Of Consciousness					
	Malaise					
	Movement Disorder					
	Syncope					
	Tremor					

Date:03/13/01ISR Number: 3680179-XReport Type:Expedited (15-DaCompany Report #B0100670A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Mouth Ulceration	Foreign	Bupropion			
Intervention to		Vulval Ulceration		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
300 MG/PER							
Prevent Permanent							
DAY/ORAL							
Impairment/Damage				Diclofenac	C		
				Co-Codamol	C		

Date:03/13/01ISR Number: 3680181-8Report Type:Expedited (15-DaCompany Report #B0100353A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Depression	Foreign	Bupropion			
Hospitalization -		Intentional Misuse		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
300 MG /PER							
Initial or Prolonged							
DAY/ ORAL				Paracetamol			
				(Acetaminophen)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/13/01ISR Number: 3680182-XReport Type:Expedited (15-DaCompany Report #B0100671A  
 Age:71 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Dyspnoea	Foreign	Bupropion			
Hospitalization - 150 MG PER	Electrocardiogram		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged DAY/ ORAL	Abnormal					
	Left Ventricular Failure		Amitriptyline	C		
	Myocardial Ischaemia		Temazepam	C		
	Pulmonary Oedema		Human Insulin	C		

Date:03/13/01ISR Number: 3680418-5Report Type:Expedited (15-DaCompany Report #B0099781A  
 Age:51 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 4 DAY	Deep Vein Thrombosis	Foreign	Zyban	PS	Glaxo Wellcome Inc	
Initial or Prolonged	Malaise	Health Professional				

Date:03/13/01ISR Number: 3680419-7Report Type:Expedited (15-DaCompany Report #B0100501A  
 Age:35 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150 MG /	Angina Pectoris	Foreign	Zyban	PS	Glaxo Wellcome Inc	
Initial or Prolonged TWICE PER DAY	Dizziness					
	Dry Mouth					
	Nausea					
	Tachycardia					

Date:03/13/01ISR Number: 3680421-5Report Type:Expedited (15-DaCompany Report #B0100349A  
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Burning Sensation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							
Intervention to Prevent Permanent Impairment/Damage		Chest Discomfort Chest Pain Haemoptysis		Beclomethasone Dipropion Gaviscon Loratadine Salmeterol Xinafoate	C C C C		

Date:03/13/01ISR Number: 3680423-9Report Type:Expedited (15-DaCompany Report #D0014564A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Convulsion	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							
		Gamma-Glutamyltransferase Increased Liver Function Test Abnormal Respiratory Arrest	Health Professional Company Representative Other	Amoxicillin	C		

Date:03/13/01ISR Number: 3680425-2Report Type:Expedited (15-DaCompany Report #B0100520A  
Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Circulatory Collapse Dizziness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
150 MG		Hypoaesthesia Musculoskeletal Stiffness Nausea Photophobia Retching	Foreign	Zyban	PS	Glaxo Wellcome Inc	
			Health Professional Other				

Date:03/13/01ISR Number: 3680427-6Report Type:Expedited (15-DaCompany Report #B0100605A  
Age: Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 22 TABLET Initial or Prolonged /PER DAY/ORAL		Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Health Professional				

Date:03/13/01ISR Number: 3680428-8Report Type:Expedited (15-DaCompany Report #B0100358A  
Age: Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Death 150 MG		Foreign	Zyban	PS	Glaxo Wellcome Inc	
		Literature Consumer Other				

Date:03/13/01ISR Number: 3680430-6Report Type:Expedited (15-DaCompany Report #B0100357A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Required 150 MG / Intervention to TWICE PER DAY		Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Other				



Prevent Permanent  
/ ORAL  
Impairment/Damage

Mefenamic Acid C

Date:03/13/01ISR Number: 3680936-XReport Type:Expedited (15-DaCompany Report #B0096694A  
Age:66 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY	Anxiety	Foreign	Zyban	PS	Glaxo Wellcome Inc	
/	Blood Pressure Decreased	Health				
	Drug Interaction	Professional				
10 MG / THREE TIMES PER DAY	Face Oedema		Deptran Tablet	SS		
	Transient Ischaemic Attack		Salmeterol + Fluticasone Aspirin	C C		

Date:03/13/01ISR Number: 3680988-7Report Type:Expedited (15-DaCompany Report #B0099525A  
Age:60 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG PER DAY ORAL	Grand Mal Convulsion Respiratory Tract Neoplasm	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/13/01ISR Number: 3680995-4Report Type:Expedited (15-DaCompany Report #B0098815A

Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Chest Discomfort Chest Pain	Foreign Literature	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG PER DAY ORAL		Decreased Appetite	Consumer				
		Dehydration		Aspirin	C		
		Dermatitis		Metformin	C		
		Dizziness		Gliclazide	C		
		Dry Mouth					
		Dry Skin					
		Dyspnoea					
		Fatigue					
		Headache					
		Insomnia					
		Lethargy					
		Myocardial Infarction					
		Oedema					
		Pain					
		Tongue Oedema					

Date:03/13/01ISR Number: 3681042-0Report Type:Expedited (15-DaCompany Report #A0116641A

Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation Arrhythmia	Foreign Literature	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE PER DAY ORAL		Coronary Artery Disease	Health				
		Dizziness	Professional				
		Feeling Abnormal					
		Myocardial Infarction					
		Sudden Death					
		Ventricular Fibrillation					

Date:03/13/01ISR Number: 3682633-3Report Type:Expedited (15-DaCompany Report #A0141757A

Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 200 MG / Initial or Prolonged TWICE PER DAY		Blood Creatinine Increased	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
/ ORAL	18	MON Blood Urea Increased Dehydration Electrocardiogram Qt Prolonged Vomiting					

Date:03/14/01ISR Number: 3680050-3Report Type:Expedited (15-DaCompany Report #A0139116A  
Age:48 YR Gender:Male I/FU:F

Outcome	PT
Other	Asthenia Dyspnoea Erythema Multiforme Hypoaesthesia Leukocytoclastic Vasculitis Neck Pain Paraesthesia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Skin Exfoliation Urticaria				
Dose	Duration		Report Source	Product	Role	Manufacturer
150MG Twice per day	3 WK			Zyban	PS	Glaxo Wellcome
				Sorbitol	C	
				Centrum Vitamin	C	
				Plendil	C	

Date:03/14/01ISR Number: 3680052-7Report Type:Expedited (15-DaCompany Report #A0140680A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose						
Other		Constipation		Zyban	PS	Glaxo Wellcome
150MG Twice per day	19 DAY	Dizziness Dyspnoea Feeling Jittery Headache Hypertension Visual Disturbance				

Date:03/14/01ISR Number: 3680053-9Report Type:Expedited (15-DaCompany Report #A0141508A  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose						
Hospitalization - 22 DAY Initial or Prolonged		Amnesia Anorexia Burning Sensation Confusional State Crying Dry Mouth Dyspnoea Feeling Abnormal Feeling Jittery Heart Rate Increased		Wellbutrin Alcohol Paxil Remeron Depakote Klonopin Premarin	PS SS C C C C C	Glaxo Wellcome

Insomnia  
Loss Of Consciousness  
Road Traffic Accident  
Suicide Attempt  
Thirst

Date:03/14/01ISR Number: 3680057-6Report Type:Expedited (15-DaCompany Report #A0141931A  
Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1TAB Twice Initial or Prolonged per day	Cardiovascular Disorder  Electroencephalogram  Abnormal Grand Mal Convulsion Injury Memory Impairment Skin Injury		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/14/01ISR Number: 3681038-9Report Type:Expedited (15-DaCompany Report #A0141841A  
Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2 WK	Completed Suicide	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL			Professional Company Representative				

Date:03/14/01ISR Number: 3682730-2Report Type:Expedited (15-DaCompany Report #B0100043A  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coronary Artery Occlusion	Foreign	Bupropion			
300 MG / PER		Ear Infection	Literature	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
DAY / ORAL		Myocardial Infarction					
				Carbimazole	C		
				Enalapril	C		
				Clarithromycin	C		
				Doxycycline	C		

Date:03/14/01ISR Number: 3682733-8Report Type:Expedited (15-DaCompany Report #B0100763A  
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 150 MG		Anaphylactic Shock	Foreign	Bupropion			
Initial or Prolonged		Aphasia	Literature	Hydrochloride	PS	Glaxo Wellcome Inc	
		Circulatory Collapse					
		Dermatitis					
		Paralysis					
		Throat Tightness					
		Tremor					
		Vomiting					

Date:03/14/01ISR Number: 3682736-3Report Type:Expedited (15-DaCompany Report #B0100749A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Blood Fibrinogen	Foreign	Bupropion			
Intervention to		Increased		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
300 MG / PER							
Prevent Permanent		Haematoma					
DAY / ORAL							
Impairment/Damage		Mean Cell Volume Abnormal					
		Red Blood Cell Count					
		Decreased					

Date:03/14/01ISR Number: 3682740-5Report Type:Expedited (15-DaCompany Report #B0100755A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cardiac Arrest	Foreign	Bupropion			
Death			Health	Hydrochloride	PS	Glaxo Wellcome Inc	
150 MG							
			Professional				
			Company				
			Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/15/01ISR Number: 3680997-8Report Type:Expedited (15-DaCompany Report #A0141153A  
Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day 7 DAY	Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
		Condition Aggravated Constipation Pain In Extremity		Zyban	SS	Glaxo Wellcome	ORAL

Date:03/15/01ISR Number: 3680998-XReport Type:Expedited (15-DaCompany Report #A0141870A  
Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Grand Mal Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged		Ischaemic Stroke		Celexa	C		
				Multiple Medications	C		

Date:03/15/01ISR Number: 3681004-3Report Type:Expedited (15-DaCompany Report #B0092315A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice per day	Headache Nausea		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
	48 DAY	Ruptured Cerebral Aneurysm Subarachnoid Haemorrhage					

Date:03/15/01ISR Number: 3681005-5Report Type:Expedited (15-DaCompany Report #B0096809A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Chest Pain		Zyban	PS	Glaxo Wellcome	ORAL



Initial or Prolonged  
per day

Dyspnoea  
Nausea

Estracombi

C

TRANSDERMAL

Date:03/15/01ISR Number: 3681006-7Report Type:Expedited (15-DaCompany Report #B0097263A  
Age:39 YR Gender:Female I/FU:F

Outcome  
Life-Threatening

- PT
- Amnesia
- Constipation
- Depression
- Diarrhoea
- Dysgeusia
- Dysphonia
- Feeling Abnormal
- Headache
- Insomnia
- Lethargy
- Malaise
- Mouth Ulceration
- Pain
- Panic Attack
- Personality Change
- Psychiatric Symptom
- Sedation
- Social Avoidant Behaviour
- Tongue Oedema

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vision Blurred

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
			Zyban	PS	Glaxo Wellcome	ORAL

Date:03/15/01ISR Number: 3681011-0Report Type:Expedited (15-DaCompany Report #B0099529A  
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	29 DAY	Dehydration		Zyban	PS	Glaxo Wellcome	
		Dizziness Face Oedema Hypersensitivity Mouth Ulceration Oral Candidiasis Oral Pain Stevens-Johnson Syndrome					

Date:03/15/01ISR Number: 3681015-8Report Type:Expedited (15-DaCompany Report #B0100994A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 4 WK		Circulatory Collapse Diarrhoea		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
		Psychotic Disorder Rectal Haemorrhage					

Date:03/15/01ISR Number: 3681016-XReport Type:Expedited (15-DaCompany Report #B0101024A  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Per day	8 DAY	Dizziness		Zyban	PS	Glaxo Wellcome	ORAL

Dry Mouth  
Fatigue  
Hordeolum  
Insomnia  
Migraine  
Mood Swings  
Thirst

Date:03/15/01ISR Number: 3681017-1Report Type:Expedited (15-DaCompany Report #B0101026A  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Twice per day	18 DAY	Sudden Death		Zyban  Bisoprolol Diclofenac Ranitidine Aspirin	PS  C C C C	Glaxo Wellcome  Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/15/01ISR Number: 3681018-3Report Type:Expedited (15-DaCompany Report #B0101033A  
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	42 DAY	Death		Zyban	PS	Glaxo Wellcome	ORAL

Date:03/15/01ISR Number: 3681019-5Report Type:Expedited (15-DaCompany Report #B0101048A  
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day 3 DAY	Asthma		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -	1TAB per day			Maxsoten	C		
Initial or Prolonged	100MG per day			Ciprofibrate	C		
	150MG per day			Allopurinol	C	Glaxo Wellcome	
	50UG Twice			Salmeterol	C	Glaxo Wellcome	
per day							
	500UG Twice			Fluticasone	C	Glaxo Wellcome	
per day							
	2PUFF Three			Duovent	C		
times per day							
				Methylprednisolone	C		

Date:03/15/01ISR Number: 3681607-6Report Type:Direct Company Report #  
Age:17 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to	150MG BID	Grand Mal Convulsion		Bupropion Hcl Tablets 75mg Geneva	PS	Geneva	ORAL

Prevent Permanent  
ORAL  
Impairment/Damage

Dexedrine C  
Celexa C  
Risperdal C

Date:03/15/01ISR Number: 3682063-4Report Type:Expedited (15-DaCompany Report #B0100348A  
Age:30 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to 150 MG/THREE Prevent Permanent TIMES PER Impairment/Damage DAY/ORAL	Anxiety C-Reactive Protein Increased Confusional State Drug Withdrawal Syndrome Dysphonia Hallucination, Visual Psychotic Disorder Restlessness Sepsis	Foreign	Bupropion Hydrochloride  Clonidine Hydrochloride Benzylpenicillin Netilmicin Sulphate Haloperidol Morphine Hydrochloride	PS      C C C C C	Glaxo Wellcome Inc	ORAL

Date:03/15/01ISR Number: 3682064-6Report Type:Expedited (15-DaCompany Report #A0141548A  
Age:64 YR Gender:Female I/FU:I

Outcome	PT
Other	Chest Pain Condition Aggravated Dyspnoea Fatigue Lower Respiratory Tract

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Infection

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
			Quinapril Hydrochloride	C		

Date:03/15/01ISR Number: 3682066-XReport Type:Expedited (15-DaCompany Report #B0100897A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Haemorrhagic Stroke Headache	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
150 MG/AS DIRECTED			Professional				

Date:03/15/01ISR Number: 3682067-1Report Type:Expedited (15-DaCompany Report #B0101145A  
Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG / PER Initial or Prolonged DAY / ORAL		Circulatory Collapse Dizziness	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/15/01ISR Number: 3682068-3Report Type:Expedited (15-DaCompany Report #B0100762A  
Age:19 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG		Overdose Suicide Attempt	Foreign Literature	Zyban Ibuprofen	PS SS	Glaxo Wellcome Inc	

Date:03/15/01ISR Number: 3682072-5Report Type:Expedited (15-DaCompany Report #B0100733A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 300 MG / PER DAY / ORAL		Asthenia Dysarthria Haemorrhagic Stroke Transient Ischaemic Attack	Foreign	Zyban  Danazol	PS  C	Glaxo Wellcome Inc	ORAL

Date:03/15/01ISR Number: 3682074-9Report Type:Expedited (15-DaCompany Report #B0098808A  
 Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 150 MG / Intervention to TWICE PER DAY Prevent Permanent Impairment/Damage		Agitation Depression Schizophrenia	Foreign Health Professional	Zyban  Quetiapine	PS  C	Glaxo Wellcome Inc	

Freedom Of Information (FOI) Report

Date:03/15/01ISR Number: 3682412-7Report Type:Expedited (15-DaCompany Report #A0134853A

Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abdominal Distension	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG / Hospitalization - THREE TIMES		Aggression	Professional				
Initial or Prolonged PER DAY/ ORAL		Anaemia					
		Anxiety		Triphasil	C		
		Ascites		Hypericum	C		
		Back Pain		Multivitamin	C		
		Bacteriuria					
		Candidiasis					
		Cellulitis					
		Cervix Carcinoma Stage I					
		Chest Discomfort					
		Chest Pain					
		Constipation					
		Cough					
		Crepitations					
		Crying					
		Cystitis					
		Decreased Appetite					
		Depression					
		Drug Level Below Therapeutic					
		Dyspnoea					
		Feeling Jittery					
		Flatulence					
		Gastro-Intestinal Fistula					
		Headache					
		Hypokalaemia					
		Hyponatraemia					
		Hypoxia					
		Implant Site Infection					
		Insomnia					
		Micturition Urgency					
		Muscle Spasms					
		Nervousness					
		Night Sweats					
		Oedema					
		Oral Candidiasis					
		Orthostatic Hypotension					
		Overdose					



Palpitations  
Paraesthesia  
Pneumonia  
Pollakiuria  
Post Coital Bleeding  
Postoperative Adhesion  
Rectal Haemorrhage  
Rhinitis  
Sinus Tachycardia  
Small Intestinal  
Obstruction  
Thyroxine Increased  
Urinary Tract Infection  
Vaginal Haemorrhage  
Vomiting

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/15/01ISR Number: 3682428-0Report Type:Expedited (15-DaCompany Report #A0138336A  
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG (TWICE Initial or Prolonged PER DAY) ORAL		Anxiety	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
		Dyspnoea	Professional				
		Rhabdomyolysis	Company Representative	Benezepril Rofecoxib	C C		

Date:03/15/01ISR Number: 3682432-2Report Type:Expedited (15-DaCompany Report #A0141472A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG (TWICE PER DAY) ORAL		Eye Disorder Visual Disturbance	Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:03/16/01ISR Number: 3681942-1Report Type:Direct Company Report #  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1 TAB EVERY DAY ORAL		Fatigue Nausea		Wellbutrin Sr 150mg	PS		ORAL

Date:03/16/01ISR Number: 3682004-XReport Type:Expedited (15-DaCompany Report #A0142204A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 14 MON		Intracranial Aneurysm		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL

Date:03/16/01ISR Number: 3682014-2Report Type:Expedited (15-DaCompany Report #B0087305A  
Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Blood Alcohol Increased Completed Suicide Drowning Drug Abuser Drug Interaction Toxicologic Test Abnormal	Zyban Alcohol	PS I	Glaxo Wellcome	ORAL

Date:03/16/01ISR Number: 3682015-4Report Type:Expedited (15-DaCompany Report #B0089264A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Complications Of Maternal Exposure To Therapeutic	Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG Twice per day	43 DAY		Drugs Premature Baby Stillbirth Twin Pregnancy				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/16/01ISR Number: 3682030-0Report Type:Expedited (15-DaCompany Report #B0101035A  
 Age:29 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice	Dyspnoea		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	Movement Disorder					
10 DAY	Paraesthesia		Salbutamol	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)	Tongue Oedema					
RESPIRATORY (INHALATION)			Beclomethasone	C	Glaxo Wellcome	
2UNIT Twice per day						

Date:03/16/01ISR Number: 3683189-1Report Type:Expedited (15-DaCompany Report #A0140680A  
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG/ TWICE	Blood Pressure Increased	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL	Constipation	Consumer				
	Dyspnoea					
	Headache					
	Tremor					
	Visual Disturbance					

Date:03/16/01ISR Number: 3683664-XReport Type:Expedited (15-DaCompany Report #A0141508A  
 Age:29 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - ORAL	Anorexia	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged UNKNOWN	Burning Sensation		Ethanol (Alcohol)	SS		

Confusional State  
Crying  
Dry Mouth  
Dyspnoea  
Feeling Abnormal  
Feeling Jittery  
Heart Rate Increased  
Insomnia  
Loss Of Consciousness  
Mental Impairment  
Road Traffic Accident  
Suicide Attempt  
Thirst

Paroxetine C  
Hydrochloride C  
Mirtazapine C  
Semisodium Valproate C  
Clonazepam C  
Conjugated Estrogens C

Date:03/16/01ISR Number: 3683665-1Report Type:Expedited (15-DaCompany Report #A0139116A  
Age:48 YR Gender:Male I/FU:I

Outcome PT  
Required Angioneurotic Oedema  
Intervention to Asthenia  
Prevent Permanent Dyspnoea  
Impairment/Damage Erythema  
Erythema Multiforme  
Hypoaesthesia  
Leukocytoclastic  
Vasculitis  
Nausea  
Neck Pain  
Paraesthesia  
Periorbital Oedema

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Skin Exfoliation Throat Irritation Urticaria					
150 MG /TWICE PER DAY/ORAL		Xerosis	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Sorbitol	C		
				Centrum	C		
				Felodipine	C		

Date:03/16/01ISR Number: 3683667-5Report Type:Expedited (15-DaCompany Report #A0141931A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1 TABLET Initial or Prolonged /TWICE PER DAY/ORAL		Angiopathy  Cerebral Ischaemia  Electroencephalogram  Abnormal Grand Mal Convulsion Injury Memory Impairment	Health  Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:03/19/01ISR Number: 3683049-6Report Type:Expedited (15-DaCompany Report #B0100993A  
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged text 50MCG Twice per day		Asthma  Chronic Obstructive Airways Disease  Exacerbated		Zyban  Salmeterol	PS  C	Glaxo Wellcome  Glaxo Wellcome	ORAL

4U As  
 required  
 200MCG Twice  
 per day

Salbutamol C Glaxo Wellcome  
 Becotide C Glaxo Wellcome

Date:03/19/01ISR Number: 3683971-0Report Type:Expedited (15-DaCompany Report #PHBS2001BE02687  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Foreign Health	Neoral	PS	Novartis Pharmaceuticals Corp	ORAL
ORAL		Drug Level Below					
		Therapeutic	Professional	Zyban (Amfebutamone)	SS		ORAL
ORAL							

Date:03/20/01ISR Number: 3684148-5Report Type:Direct Company Report #  
 Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Wellbutrin (Generic)	PS		
100 MG BID		Headache		Bupropion	C		
		Insomnia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/20/01ISR Number: 3684572-0Report Type:Expedited (15-DaCompany Report #B0095941A  
Age:28 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 4 WK Initial or Prolonged	Malaise		Zyban	PS	Glaxo Wellcome	ORAL
	Neutrophil Count Decreased Tonsillitis Weight Decreased White Blood Cell Count Decreased					

Date:03/20/01ISR Number: 3684579-3Report Type:Expedited (15-DaCompany Report #B0101536A  
Age:37 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 1200MG per Initial or Prolonged day	Accidental Overdose		Zyban	PS	Glaxo Wellcome	
	Cerebrovascular Accident					
	Dizziness		No Concurrent Medication	C		

Date:03/20/01ISR Number: 3685076-1Report Type:Direct  
Age:55 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Fall		Bupropion	PS		
	Medication Error		Risperidone	SS		
			Clonazepam	SS		
			Venlafaxine	SS		

Date:03/20/01ISR Number: 3685079-7Report Type:Direct  
Age:55 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						



Hospitalization - Confusional State  
Initial or Prolonged Delirium  
Fall  
Overdose  
Urinary Incontinence

Bupropion 150mg Q Am PS  
Effexor 150mg Bid SS  
Risperidone C  
Clonazepam C

Date:03/20/01ISR Number: 3685228-0Report Type:Direct  
Age:44 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis		Zyban (150mg) (Catalytica Pharmaceuticals)	PS	Catalytica Pharmaceuticals	ORAL
150MG PO BID							

Date:03/20/01ISR Number: 3685235-8Report Type:Direct  
Age:52 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash Pruritic		Zyban (150mg) (Catalytica Pharmaceuticals)	PS	Catalytica Pharmaceuticals	ORAL
150MG PO BID							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nicoderm C

Date:03/20/01ISR Number: 3685613-7Report Type:Expedited (15-DaCompany Report #A0141870A  
Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coma	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - Initial or Prolonged		Grand Mal Convulsion Ischaemic Stroke	Professional Company Representative	Citalopram Hydrobromide Multiple Medication	C C		

Date:03/20/01ISR Number: 3685655-1Report Type:Expedited (15-DaCompany Report #10405  
Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 18 MG 1X/1 Initial or Prolonged DAY, ORAL		Convulsion	Health	Concerta	PS	Alza Corp	ORAL
450 MG 1X/1 DAY, PO		Drug Interaction	Professional Other	Wellbutrin	SS		ORAL
				Doxycycline Acne Cream	C C		

Date:03/20/01ISR Number: 3686287-1Report Type:Expedited (15-DaCompany Report #B0097263A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening ORAL		Amnesia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Constipation Depression Diarrhoea Disturbance In Attention Dysphonia	Health Professional				

Feeling Abnormal  
Headache  
Insomnia  
Lethargy  
Malaise  
Mouth Haemorrhage  
Mouth Ulceration  
Palpitations  
Panic Attack  
Personality Change  
Psychiatric Symptom  
Rectal Tenesmus  
Sedation  
Social Avoidant Behaviour  
Tongue Oedema  
Vision Blurred  
Weight Increased

Date:03/20/01ISR Number: 3686288-3Report Type:Expedited (15-DaCompany Report #B0101024A  
Age:29 YR Gender:Female I/FU:I

Outcome PT  
Disability Dizziness  
Dry Mouth  
Fatigue

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Hordeolum Insomnia Migraine	Report Source	Product	Role	Manufacturer	Route
150 MG/PER DAY/ORAL		Mood Swings Thirst	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/20/01ISR Number: 3686289-5Report Type:Expedited (15-DaCompany Report #B0101033A  
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG/ORAL		Sudden Death	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/20/01ISR Number: 3686290-1Report Type:Expedited (15-DaCompany Report #B0101026A  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG/TWICE PER DAY/ORAL		Sudden Death	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Bisoprolol	C
Diclofenac	C
Ranitidine	
Hydrochloride	C
Aspirin	C

Date:03/20/01ISR Number: 3686291-3Report Type:Expedited (15-DaCompany Report #B0096809A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL		Chest Pain Condition Aggravated	Foreign Health	Zyban	PS	Glaxo Wellcome Inc	ORAL

Dyspnoea  
Hiatus Hernia  
Malaise

Professional

Estracombi

C

Date:03/20/01ISR Number: 3686292-5Report Type:Expedited (15-DaCompany Report #B0101048A  
Age:66 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150 MG/PER Hospitalization - DAY/ORAL Initial or Prolonged	Asthma Malaise	Foreign Health Professional	Zyban  Maxsoten Ciprofibrate Allopurinol Salmeterol Xinafoate Fluticasone Propionate Duovent Methylprednisolone	PS  C C C C C C C	Glaxo Wellcome Inc	ORAL

Date:03/20/01ISR Number: 3686293-7Report Type:Expedited (15-DaCompany Report #B0100994A  
Age:36 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Circulatory Collapse Diarrhoea Psychotic Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Rectal Haemorrhage

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
ORAL	4 WK		Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:03/20/01ISR Number: 3686297-4Report Type:Expedited (15-DaCompany Report #B0092315A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG/ TWICE		Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL		Headache	Health				
		Intraventricular Haemorrhage	Professional				
		Nausea					
		Nervous System Disorder					
		Ruptured Cerebral Aneurysm					
		Subarachnoid Haemorrhage					

Date:03/20/01ISR Number: 3686298-6Report Type:Expedited (15-DaCompany Report #B0099529A  
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN	150 MG/ TWICE		Foreign	Zyban	PS	Glaxo Wellcome Inc	
Initial or Prolonged PER DAY/ UNKNOWN		Dry Mouth	Health				
		Face Oedema	Professional				
		Hypersensitivity					
		Mouth Ulceration					
		Oral Candidiasis					
		Sensory Loss					
		Stevens-Johnson Syndrome					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Chest Pain	Health				
DAY/ ORAL		Constipation	Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	SS		ORAL
150 MG/ UNK/							
ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complications Of Maternal Exposure To Therapeutic	Foreign Study	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Drugs	Health				
PER DAY/ ORAL		Premature Baby Stillbirth Twin Pregnancy	Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/20/01ISR Number: 3686542-5Report Type:Expedited (15-DaCompany Report #B0101035A  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/TWICE PER DAY/ ORAL		Dyspnoea Paraesthesia Tongue Oedema	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Salbutamol Sulphate Beclomethasone Dipropion	C C		

Date:03/20/01ISR Number: 3686543-7Report Type:Expedited (15-DaCompany Report #A0142204A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL	14 MON	Intracranial Aneurysm	Foreign Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:03/20/01ISR Number: 3686615-7Report Type:Expedited (15-DaCompany Report #B0087305A  
Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death ORAL		Blood Alcohol Increased	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Completed Suicide Drowning Drug Abuser Drug Interaction Drug Level Above Therapeutic	Health Professional	Ethanol (Formulation Unknown) (Alcohol)	SS		

Date:03/21/01ISR Number: 3685954-3Report Type:Expedited (15-DaCompany Report #A0131385A  
Age:24 YR Gender:Male I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Back Pain	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice							
Hospitalization -		Confusional State	Professional				
per day							
Initial or Prolonged		Eosinophilia		Xanax	C		
Other		Headache					
		Hepatitis Viral					
		Liver Function Test					
		Abnormal					
		Multi-Organ Failure					
		Neutrophilia					
		Petechiae					
		Pyrexia					
		Stevens-Johnson Syndrome					
		Thrombocytopenia					
		Toxic Epidermal					
		Necrolysis					
		Viral Infection					

Date:03/21/01ISR Number: 3685956-7Report Type:Expedited (15-DaCompany Report #A0136852A  
Age:53 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Corneal Erosion
Initial or Prolonged	Hypoaesthesia Eye
	Keratitis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Peripheral Nerve Injury

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1	MON	Health	Zyban	PS	Glaxo Wellcome	ORAL
		Professional	Ventolin	C	Glaxo Wellcome	
			Singulair	C		
			Ecotrin	C		
			Paxil	C		

Date:03/21/01ISR Number: 3685961-0Report Type:Expedited (15-DaCompany Report #A0141624A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Brain Damage Convulsion Mental Impairment		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:03/21/01ISR Number: 3685972-5Report Type:Expedited (15-DaCompany Report #B0088230A

Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Acidosis	Health	Zyban	PS	Glaxo Wellcome	ORAL
50TAB Single							
Hospitalization -		Coma	Professional				
dose	1 DAY						
Initial or Prolonged		Convulsion		Anadin Extra	SS		ORAL
24TAB Single							
dose	1 DAY	Hyperhidrosis					
		Hypotension		Alcohol	SS		
		Intentional Misuse					
		Irritability					
		Pyrexia					
		Tachycardia					
		Vomiting					

Date:03/21/01ISR Number: 3685973-7Report Type:Expedited (15-DaCompany Report #B0088388A  
Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Accident		Zyban	PS	Glaxo Wellcome	ORAL
3 WK		Circulatory Collapse Ecchymosis Petit Mal Epilepsy		Lithium	C		

Date:03/21/01ISR Number: 3685974-9Report Type:Expedited (15-DaCompany Report #B0091168A  
Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Acidosis	Consumer	Bupropion	PS	Glaxo Wellcome	ORAL
1 DAY							
Hospitalization -		Hyperhidrosis		Paracetamol	SS		
Initial or Prolonged		Hypotension		Alcohol	SS		
Other		Overdose Pyrexia Tachycardia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/21/01ISR Number: 3685975-0Report Type:Expedited (15-DaCompany Report #B0094744A  
Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	1 DAY	Acidosis	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -	1 DAY	Convulsion		Anadin	SS		ORAL
Initial or Prolonged		Hypotension					
Other		Overdose					
		Tachycardia					

Date:03/21/01ISR Number: 3685979-8Report Type:Expedited (15-DaCompany Report #B0099522A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	100MG Twice	Attention	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	per day 13 DAY	Deficit/Hyperactivity					
Disability		Disorder					
		Encephalitis					
		Meningitis Aseptic					
		Pleocytosis					

Date:03/21/01ISR Number: 3685980-4Report Type:Expedited (15-DaCompany Report #B0099526A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	UNKNOWN	Astrocytoma	Consumer	Zyban	PS	Glaxo Wellcome	
Hospitalization -		Coma		Prozac	C		
Initial or Prolonged		Dysphagia					
		Infection					
		Nervous System Disorder					
		Sedation					
		Urinary Incontinence					
		Vomiting					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Intermittent Claudication		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day							
60MG In the		Vitreous Haemorrhage		Isib	C		ORAL
morning							
				Co-Codamol	C		
400MCG At				Cerivastatin	C		
night							
75MG Per day				Clopidogrel	C		
40MG In the				Frusemide	C		
morning							
4MG In the				Lacidipine	C	Glaxo Wellcome	
morning							
20MG In the				Fluoxetine	C		
morning							
75MG Per day				Aspirin	C		
				Gtn	C	Glaxo Wellcome	
RESPIRATORY				Beclomethasone	C	Glaxo Wellcome	
(INHALATION)	2PUFF Twice						
per day							
150MG Twice				Ranitidine	C	Glaxo Wellcome	
per day							
20MG At night				Amitriptyline	C		

Freedom Of Information (FOI) Report

7.5G At night

Senna

C

Date:03/21/01ISR Number: 3685983-XReport Type:Expedited (15-DaCompany Report #B0100785A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 7 DAY		Aggression		Zyban	PS	Glaxo Wellcome	ORAL
		Crying					
		Depression					
		Diarrhoea					
		Diplopia					
		Disturbance In Attention					
		Dizziness					
		Eye Pain					
		Fatigue					
		Headache					
		Nausea					
		Suicidal Ideation					
		Vomiting					

Date:03/21/01ISR Number: 3685984-1Report Type:Expedited (15-DaCompany Report #B0100789A  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Twice per day		Anxiety Arthralgia		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
		Hyperhidrosis					
		Nausea					
		Peripheral Coldness					
		Tremor					

Date:03/21/01ISR Number: 3685987-7Report Type:Expedited (15-DaCompany Report #B0101027A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day 3 DAY	Drug Interaction		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -	120MG Twice	Grand Mal Convulsion		Gamolenic Acid	SS		ORAL
Initial or Prolonged per day				Diclofenac	C		ORAL
50MG Three times per day				Co-Codamol	C		ORAL

Date:03/21/01ISR Number: 3685993-2Report Type:Expedited (15-DaCompany Report #B0101331A  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice per day	Coronary Artery Disease Myocardial Infarction		Zyban	PS	Glaxo Wellcome	ORAL

Date:03/21/01ISR Number: 3685994-4Report Type:Expedited (15-DaCompany Report #B0101340A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG As	Atrial Fibrillation		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged directed		Drug Interaction					
200MCG per		Thyroxine Increased		Thyroxine	SS	Glaxo Wellcome	ORAL

Freedom Of Information (FOI) Report

day

Date:03/21/01ISR Number: 3685995-6Report Type:Expedited (15-DaCompany Report #B0101341A  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day 5 DAY	Atrial Fibrillation		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - RESPIRATORY	Initial or Prolonged (INHALATION)	Cardiac Failure		Salbutamol	C	Glaxo Wellcome	
RESPIRATORY	(INHALATION)	Congestive		Beclomethasone	C	Glaxo Wellcome	
		Dizziness					
		Headache					
	10 DAY						

Date:03/21/01ISR Number: 3685996-8Report Type:Expedited (15-DaCompany Report #B0101342A  
 Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice	Cardiac Arrest		Zyban	PS	Glaxo Wellcome	ORAL
	per day 7 DAY	Circulatory Collapse					
	125MG per day	Dyspnoea		Ramipril	C		ORAL
	10MG per day	Lower Respiratory Tract		Amlodipine	C		ORAL
	20MG Twice	Infection		Isosorbide	C		ORAL
	per day			Simvastatin	C		ORAL
	10MG per day			Combivent	C		
RESPIRATORY	(INHALATION)						



Date:03/21/01ISR Number: 3685997-XReport Type:Expedited (15-DaCompany Report #B0101343A  
Age:68 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	40	DAY					
50MG Twice				Diclofenac	C		ORAL
per day							

Date:03/21/01ISR Number: 3685998-1Report Type:Expedited (15-DaCompany Report #B0101345A  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Respiratory Disorder		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day							
RESPIRATORY				Combivent	C		
(INHALATION)							

Date:03/21/01ISR Number: 3686000-8Report Type:Expedited (15-DaCompany Report #B0101349A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Interaction		Zyban	PS	Glaxo Wellcome	ORAL
29 DAY							
173 DAY				Warfarin	I	Glaxo Wellcome	ORAL

Date:03/21/01ISR Number: 3686001-XReport Type:Expedited (15-DaCompany Report #B0101353A  
Age:67 YR Gender:Female I/FU:I

Outcome	PT
Death	Cerebrovascular Accident Dyspnoea

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Oedema

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day			Zyban	PS	Glaxo Wellcome	ORAL
UNKNOWN	75MG per day		Aspirin	C		
UNKNOWN	2PUFF Four times per day		Salbutamol	C	Glaxo Wellcome	
UNKNOWN	250MCG Twice per day		Fluticasone	C	Glaxo Wellcome	
UNKNOWN	75MG Twice per day		Diclofenac	C		
UNKNOWN	50MCG Twice per day		Salmeterol	C	Glaxo Wellcome	
UNKNOWN	200MCG Twice per day		Oxitropium	C		
UNKNOWN	10MG Per day		Imipramine	C		
UNKNOWN	40MG Per day		Frusemide	C		

Date:03/21/01ISR Number: 3686003-3Report Type:Expedited (15-DaCompany Report #B0101358A  
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Hypertension		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice per day		Myocardial Ischaemia					

80MG Per day				Valsartan	C		ORAL
5MG Per day				Bendrofluazide	C	Glaxo Wellcome	ORAL
Date:03/21/01ISR Number: 3686004-5Report Type:Expedited (15-DaCompany Report #B0101365A Age:48 YR Gender:Male I/FU:F							
Outcome	PT	Report Source	Product	Role	Manufacturer	Route	
Dose	Duration						
Hospitalization - 10 DAY	Blood Pressure Increased		Zyban	PS	Glaxo Wellcome		
Initial or Prolonged	Blood Viscosity Increased Embolism Facial Palsy						
Date:03/21/01ISR Number: 3686005-7Report Type:Expedited (15-DaCompany Report #B0101366A Age:38 YR Gender:Female I/FU:F							
Outcome	PT	Report Source	Product	Role	Manufacturer	Route	
Dose	Duration						
Hospitalization - 150MG As	Anaphylactic Reaction	Consumer	Zyban	PS	Glaxo Wellcome	ORAL	
Initial or Prolonged directed	Blood Pressure Increased						
500MG Three times per day	Chills		Amoxicillin	SS			
.05ML Four times per day	Heart Rate Increased						
	Urticaria		Budesonide	C		NASAL	
			Xylometazoline	C			
Date:03/21/01ISR Number: 3686006-9Report Type:Expedited (15-DaCompany Report #B0101413A Age:28 YR Gender:Female I/FU:I							
Outcome	PT	Report Source	Product	Role	Manufacturer	Route	
Dose	Duration						
Life-Threatening 150MG Twice per day	Alcoholism		Zyban	PS	Glaxo Wellcome	ORAL	
	Grand Mal Convulsion						
	Status Epilepticus		Tramadol	C		ORAL	
			Alcohol	C			



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/21/01ISR Number: 3686007-0Report Type:Expedited (15-DaCompany Report #B0101420A  
 Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice		Myocardial Infarction	Zyban	PS	Glaxo Wellcome	ORAL
	per day						

Date:03/21/01ISR Number: 3686008-2Report Type:Expedited (15-DaCompany Report #B0101422A  
 Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day	92 DAY	Therapeutic Response	Zyban	PS	Glaxo Wellcome	ORAL
	20MG per day		Unexpected	Amitriptyline	C		ORAL
	30MG per day		Vitreous Haemorrhage	Isosorbide	C		ORAL
	8U per day			Co-Codamol	C		ORAL
	400MCG per day			Cerivastatin	C		ORAL
	75MG per day			Clopidogrel	C		ORAL
	40MG per day			Frusemide	C		ORAL
	40MG per day			Lacidipine	C	Glaxo Wellcome	ORAL
	20MG per day			Fluoxetine	C		ORAL
	75MG per day			Aspirin	C		ORAL
	300MG per day			Ranitidine	C	Glaxo Wellcome	ORAL
	7.5MG per day			Senna	C		ORAL
	RESPIRATORY			Salbutamol	C	Glaxo Wellcome	
	( INHALATION)						

1000MG per				Naproxen	C		ORAL
day							
400MG per day				Trimethoprim	C	Glaxo Wellcome	ORAL
				Nicorandil	C		ORAL

Date:03/21/01ISR Number: 3686013-6Report Type:Expedited (15-DaCompany Report #D0014384A  
Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health	Zyban	PS	Glaxo Wellcome	ORAL
300MG per day	6 WK	Sedation	Professional				

Date:03/21/01ISR Number: 3687101-0Report Type:Expedited (15-DaCompany Report #A038609  
Age:18 YR Gender:Male I/FU:F

Outcome	PT
Required	Confusional State
Intervention to	Crying
Prevent Permanent	Depression
Impairment/Damage	Disorientation
	Dizziness
	Drug Interaction
	Emotional Disorder
	Feeling Jittery
	Hallucination, Visual
	Hyperhidrosis
	Inappropriate Affect
	Mood Swings
	Nasopharyngitis
	Psychomotor Hyperactivity
	Serotonin Syndrome
	Thinking Abnormal
	Tremor

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vision Blurred

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
75.00 MG		Health Professional	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
TOTAL:DAILY:ORAL						
150.00 MG			Wellbutrin	SS		
TOTAL: BID						
			Nyquil	SS		
UNKNOWN						

Date:03/21/01ISR Number: 3687618-9Report Type:Expedited (15-DaCompany Report #B0100993A  
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 ORAL		Asthma	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Chronic Obstructive Airways Disease Exacerbated	Health Professional	Salmeterol Xinafoate Salbutamol Sulphate Beclomethasone Dipropion	C C C		

Date:03/22/01ISR Number: 3687226-XReport Type:Expedited (15-DaCompany Report #A0142820A  
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Insomnia		Zyban	PS	Glaxo Wellcome	ORAL
		Lung Neoplasm Malignant					
		Pruritus					

Date:03/22/01ISR Number: 3687231-3Report Type:Expedited (15-DaCompany Report #B0100196A  
Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice Initial or Prolonged per day 20 DAY	Blood Pressure Decreased Cough Dehydration Dyspnoea Fatigue Hypersensitivity Oedema Peripheral Pharyngolaryngeal Pain Pyrexia Rash Pruritic Urticaria		Zyban	PS	Glaxo Wellcome	

Date:03/22/01ISR Number: 3687232-5Report Type:Expedited (15-DaCompany Report #B0100198A  
Age:36 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Dyspnoea Hyperventilation		Zyban Medroxyprogesterone Acetate Amitriptyline Hydrochloride Fluoxetine Hydrochloride Pizotifen Malate	PS C C C C	Glaxo Wellcome	ORAL ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Orlistat C

Date:03/22/01ISR Number: 3687236-2Report Type:Expedited (15-DaCompany Report #B0101350A  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day							
60MG Twice				Isosorbide	C		
per day							
20MG Twice				Nifedipine	C		
per day							
				Alverine	C		
				Aspirin	C		

Date:03/22/01ISR Number: 3687238-6Report Type:Expedited (15-DaCompany Report #B0101552A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction		Zyban	PS	Glaxo Wellcome	ORAL
		Sudden Death					

Date:03/22/01ISR Number: 3687239-8Report Type:Expedited (15-DaCompany Report #B0101558A  
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death		Zyban	PS	Glaxo Wellcome	ORAL
14 DAY							
1MG At night				Lormetazepam	C		ORAL
100MCG per				Thyroxine	C	Glaxo Wellcome	ORAL

day  
 80MG per day  
 Gliclazide C ORAL

Date:03/22/01ISR Number: 3687240-4Report Type:Expedited (15-DaCompany Report #B0101569A  
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG		Anxiety		Zyban	PS	Glaxo Wellcome	ORAL
Variable dose	9 DAY	Asthma					
RESPIRATORY (INHALATION)		Dyspnoea		Fluticasone	C	Glaxo Wellcome	
	500MCG	Hallucination Twice Insomnia					
per day RESPIRATORY (INHALATION)		Panic Attack		Salbutamol	C	Glaxo Wellcome	
1TAB per day				Trinordiol	C		ORAL

Date:03/22/01ISR Number: 3687242-8Report Type:Expedited (15-DaCompany Report #B0101671A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Distension Nephritis Renal Disorder Swelling		Zyban	PS	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/23/01ISR Number: 3688206-0Report Type:Expedited (15-DaCompany Report #A0142629A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	7 DAY		Zyban	PS	Glaxo Wellcome	ORAL
Other			Akathisia				
			Depersonalisation	Multivitamin	C		
			Mental Disorder	Calcium	C		
			Tremor				

Date:03/23/01ISR Number: 3688212-6Report Type:Expedited (15-DaCompany Report #B0099781A

Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	UNKNOWN	4 DAY		Zyban	PS	Glaxo Wellcome	
Hospitalization -			Haemorrhoids				
Initial or Prolonged			Malaise	No Concurrent Medication	C		

Date:03/23/01ISR Number: 3688982-7Report Type:Expedited (15-DaCompany Report #B0099532A

Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150 MG /PER			Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Other			Intermittent Claudication				
			Periorbital Haematoma				
			Retinal Detachment				
			Vitreous Haemorrhage	Isosorbide Monitrate	C		
				Co-Codamol	C		
				Cerivastatin	C		
				Clopidogrel	C		
				Frusemide	C		
				Lacidipine	C		
				Fluoxetine	C		
				Aspirin	C		
				Nitroglycerin	C		
				Beclomethasone			
				Dipropion	C		
				Ranitidine			

Hydrochloride C  
 Amitriptyline C  
 Sennosides C

Date:03/23/01ISR Number: 3688984-0Report Type:Expedited (15-DaCompany Report #B0088230A

Age:18 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 50 Initial or Prolonged TABLET/SINGLE DOSE/ORAL 24 TABLET /SINGLE DOSE/ ORAL UNKNOWN	Acidosis Blood Pressure Decreased Coma Hyperhidrosis Hypotension Intentional Misuse Irritability Pyrexia Tachycardia Vomiting	Foreign Health Professional	Bupropion Hydrochloride   Anadin Extra Tablet (Anadin Extra)   Ethanol L(Alcohol)	PS    SS   SS	Glaxo Wellcome Inc	ORAL    ORAL

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Freedom Of Information (FOI) Report

Date:03/23/01ISR Number: 3688986-4Report Type:Expedited (15-DaCompany Report #B0094744A  
 Age:18 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 7.5 G /ORAL	Acidosis Convulsion	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged 7.5 G /ORAL	Hypotension		Anadin((Anadin)	SS		ORAL
Required Intervention to Prevent Permanent Impairment/Damage	Intentional Misuse Tachycardia					

Date:03/23/01ISR Number: 3688988-8Report Type:Expedited (15-DaCompany Report #B0091168A  
 Age:18 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 150 MG/ORAL	Acidosis Convulsion	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged Required UNKNOWN	Hyperhidrosis Hypotension		Paracetamol (Acetaminophen)	SS		
Intervention to UNKNOWN Prevent Permanent Impairment/Damage	Irritability  Overdose Pyrexia Tachycardia		Ethanol (Alcohol)	SS		

Date:03/23/01ISR Number: 3689074-3Report Type:Expedited (15-DaCompany Report #B0101536A  
 Age:37 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Cerebrovascular Accident Medication Error	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:03/23/01ISR Number: 3689077-9Report Type:Expedited (15-DaCompany Report #B0095941A  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL	4 WK	Malaise Tonsillitis	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Weight Decreased White Blood Cell Count Decreased	Professional Company Representative				

Date:03/23/01ISR Number: 3689470-4Report Type:Expedited (15-DaCompany Report #A0141624A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Brain Damage	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
		Convulsion Thinking Abnormal					

Date:03/23/01ISR Number: 3689485-6Report Type:Expedited (15-DaCompany Report #A0136852A  
Age:53 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Corneal Erosion Eye Pain

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Keratitis Keratoconjunctivitis Sicca	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Sensory Loss	Professional	Salbutamol Sulphate	C		
				Montelukast Sodium	C		
				Aspirin	C		
				Paroxetine Hydrochloride	C		

Date:03/23/01ISR Number: 3689486-8Report Type:Expedited (15-DaCompany Report #A0131385A  
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Back Pain	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG / Hospitalization - TWICE PER DAY Initial or Prolonged / ORAL Required Intervention to Prevent Permanent Impairment/Damage		Confusional State	Professional				
		Dermatitis		Alprazolam	C		
		Eosinophilia					
		Headache					
		Liver Function Test Abnormal					
		Multi-Organ Failure					
		Neutrophilia					
		Petechiae					
		Pyrexia					
		Stevens-Johnson Syndrome					
		Thrombocytopenia					
		Toxic Epidermal Necrolysis					
		Viral Infection					

Date:03/23/01ISR Number: 3689989-6Report Type:Expedited (15-DaCompany Report #B0100785A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aggression	Foreign	Bupropion			

ORAL	7	DAY	Crying	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
			Depression	Professional				
			Diarrhoea					
			Diplopia					
			Disturbance In Attention					
			Dizziness					
			Eye Pain					
			Fatigue					
			Headache					
			Nausea					
			Suicidal Ideation					
			Vomiting					

Date:03/23/01ISR Number: 3689990-2Report Type:Expedited (15-DaCompany Report #B0101027A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Drug Interaction	Foreign	Bupropion			
Hospitalization -		Grand Mal Convulsion		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (PER							
Initial or Prolonged							
DAY) ORAL							
				Gamolenic Acid			
				(Gamolenic Acid)	SS		ORAL
120 MG (TWICE							



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Freedom Of Information (FOI) Report

PER DAY) ORAL

Diclofenac C  
Co-Comadol C

Date:03/23/01ISR Number: 3689992-6Report Type:Expedited (15-DaCompany Report #B0100789A  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Anxiety Arthralgia	Foreign Study	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL		Hyperhidrosis	Health				
		Nausea Peripheral Coldness Tremor	Professional				

Date:03/23/01ISR Number: 3689993-8Report Type:Expedited (15-DaCompany Report #B0088388A  
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG ORAL	3 WK	Circulatory Collapse Ecchymosis	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Prevent Permanent Impairment/Damage		Petit Mal Epilepsy	Professional	Lithium Salt	C		

Date:03/23/01ISR Number: 3689997-5Report Type:Expedited (15-DaCompany Report #B0101343A  
Age:68 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG/ TWICE PER DAY/ ORAL		Myocardial Infarction	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Diclofenac	C		

Date:03/23/01ISR Number: 3689998-7Report Type:Expedited (15-DaCompany Report #B0101342A  
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Dyspnoea					
PER DAY/ ORAL		Lower Respiratory Tract Infection		Ramipril	C		
				Amlodipine	C		
				Isosorbide	C		
				Simvastatin	C		
				Combivent	C		

Date:03/23/01ISR Number: 3690000-1Report Type:Expedited (15-DaCompany Report #B0101422A  
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Therapeutic Response	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Unexpected					
DAY/ ORAL		Vitreous Haemorrhage		Amitriptyline	C		
				Isosorbide	C		
				Co-Codamol	C		
				Cerivastatin	C		
				Clopidogrel	C		
				Frusemide	C		
				Lacidipine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Fluoxetine	C
Aspirin	C
Ranitidine	
Hydrochloride	C
Sennosides	C
Salbutamol Sulphate	C
Naproxen	C
Trimethoprim	C
Nicorandil	C

Date:03/23/01ISR Number: 3690001-3Report Type:Expedited (15-DaCompany Report #B0101366A  
 Age:38 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ AS	Anaphylactic Reaction	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged DIRECTED/ ORAL	Blood Pressure Increased Chills					
500 MG/ THREE TIMES PER DAY	Heart Rate Increased Pruritus Urticaria		Amoxicillin Capsule (Amoxicillin)	SS		
			Budesonide Xylometazoline	C C		

Date:03/23/01ISR Number: 3690002-5Report Type:Expedited (15-DaCompany Report #B0101365A  
 Age:48 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG	Blood Pressure Increased	Foreign	Zyban	PS	Glaxo Wellcome Inc	
Initial or Prolonged	Embolism Facial Palsy Haemoconcentration	Consumer				

Date:03/23/01ISR Number: 3690003-7Report Type:Expedited (15-DaCompany Report #B0101341A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG/ PER	Atrial Fibrillation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Hospitalization -	DAY/ ORAL	Cardiac Failure					
Initial or Prolonged		Congestive		Salbutamol Sulphate	C		
		Dizziness		Beclomethasone			
		Headache		Dipropion.	C		

Date:03/23/01ISR Number: 3690004-9Report Type:Expedited (15-DaCompany Report #B0101340A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG/ AS	Atrial Fibrillation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged	DIRECTED/	Drug Interaction					
ORAL				Thyroxine Sodium (Formulation Unknown) (Levothyroxine Sodium)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/23/01ISR Number: 3690006-2Report Type:Expedited (15-DaCompany Report #B0101331A  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coronary Artery Disease Myocardial Infarction	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL							

Date:03/23/01ISR Number: 3690007-4Report Type:Expedited (15-DaCompany Report #B0101420A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL							

Date:03/23/01ISR Number: 3690008-6Report Type:Expedited (15-DaCompany Report #B0101413A  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Grand Mal Convulsion Status Epilepticus	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL							
				Tramadol Hydrochloride	C		
				Ethanol	C		

Date:03/23/01ISR Number: 3690009-8Report Type:Expedited (15-DaCompany Report #B0101358A  
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death	Hypertension	Foreign	Bupropion		
	Myocardial Ischaemia		Hydrochloride	PS	Glaxo Wellcome Inc
150 MG (TWICE					
PER DAY)			Valsartan	C	
			Bendrofluazide	C	

Date:03/23/01ISR Number: 3690010-4Report Type:Expedited (15-DaCompany Report #B0101353A  
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebrovascular Accident	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE							
PER DAY) ORAL				Aspirin	C		
				Salbutamol Sulphate	C		
				Fluticasone			
				Propionate	C		
				Diclofenac	C		
				Salmeterol Xinafoate	C		
				Oxitropium Bromide	C		
				Imipramine	C		
				Frusemide	C		

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Freedom Of Information (FOI) Report

Date:03/23/01ISR Number: 3690011-6Report Type:Expedited (15-DaCompany Report #B0101349A  
 Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG ORAL Prevent Permanent Impairment/Damage (AS DIRECT)		Drug Interaction	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Warfarin Sodium (Warfarin Sodium)	SS		ORAL
ORAL							

Date:03/23/01ISR Number: 3690012-8Report Type:Expedited (15-DaCompany Report #B0101345A  
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG (PER DAY) ORAL		Respiratory Disorder Sudden Death	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Combivent	C		

Date:03/23/01ISR Number: 3690013-XReport Type:Expedited (15-DaCompany Report #B0099526A  
 Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - Initial or Prolonged		Astrocytoma Body Temperature Increased Coma Dysphagia Ganglioneuroma Infection Memory Impairment Urinary Incontinence Vomiting	Foreign Health Professional	Bupropion Hydrochloride Fluoxetine Hydrochloride	PS  C	Glaxo Wellcome Inc	

Date:03/23/01ISR Number: 3690014-1Report Type:Expedited (15-DaCompany Report #D0014384A  
Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Sedation	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL			Professional				

Date:03/23/01ISR Number: 3690015-3Report Type:Expedited (15-DaCompany Report #B0099522A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 100 MG (TWICE Disability PER DAY) ORAL		Aggression Attention  Deficit/Hyperactivity  Disorder Encephalitis Meningitis Aseptic	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/26/01ISR Number: 3688834-2Report Type:Expedited (15-DaCompany Report #A0140463A

Age:27 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	10 DAY	Arrhythmia		Zyban	PS	Glaxo Wellcome	ORAL
		Euphoric Mood Feeling Abnormal Hypersensitivity Myocardial Infarction Palpitations Psychomotor Hyperactivity					

Date:03/26/01ISR Number: 3689555-2Report Type:Direct

Company Report #

Age:42 YR Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Angioneurotic Oedema Pruritus Urticaria		Zyban	PS		

Date:03/26/01ISR Number: 3689561-8Report Type:Direct

Company Report #

Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	3 PER DAY	Circulatory Collapse Coma		Gammahydroxybutyrate (Ghb)	PS	Wellbutrin	
				Wellbutrin	SS		

Date:03/26/01ISR Number: 3689879-9Report Type:Expedited (15-DaCompany Report #B0100198A

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG	Dyspnoea Hyperventilation	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Professional

Medroxyprogesterone

Ace. C

Amitriptyline Hcl C

Fluoxetine

Hydrochloride C

Pizotifen Malate C

Orlistat C

Date:03/26/01ISR Number: 3689881-7Report Type:Expedited (15-DaCompany Report #B0101350A

Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
PER DAY ORAL							
				Isosorbide	C		
				Nifedipine	C		
				Alverine Citrate	C		
				Aspirin	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/26/01ISR Number: 3689884-2Report Type:Expedited (15-DaCompany Report #B0101558A  
 Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL				Lormetazepam	C		
				Thyroxine Sodium	C		
				Gliclazide	C		

Date:03/26/01ISR Number: 3689886-6Report Type:Expedited (15-DaCompany Report #B0101569A  
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG Prevent Permanent VARIABLE DOSE Impairment/Damage ORAL		Anxiety Asthma Hallucination Insomnia Panic Attack	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Fluticasone Propionate	C		
				Salbutamol Sulphate	C		
				Trinordiol	C		

Date:03/26/01ISR Number: 3689888-XReport Type:Expedited (15-DaCompany Report #B0100196A  
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG TWICE PER DAY		Blood Pressure Decreased Cough Dehydration Dyspnoea Fatigue Hypersensitivity Oedema Peripheral		Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Pharyngolaryngeal Pain  
Pyrexia  
Rash Pruritic  
Urticaria

Date:03/26/01ISR Number: 3690157-2Report Type:Expedited (15-DaCompany Report #A0142820A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Insomnia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Lung Neoplasm Malignant	Consumer				
PER DAY /		Pruritus					
ORAL							

Date:03/26/01ISR Number: 3690158-4Report Type:Expedited (15-DaCompany Report #A0101552A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Myocardial Infarction	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL		Sudden Death					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/26/01ISR Number: 3690159-6Report Type:Expedited (15-DaCompany Report #B0101671A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Distension	Foreign	Zyban	PS	Glaxo Wellcome Inc	
150 MG /		Nephritis	Health				
		Renal Disorder	Professional				
		Swelling					

Date:03/27/01ISR Number: 3689661-2Report Type:Expedited (15-DaCompany Report #B0100993A

Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Asthma		Zyban	PS	Glaxo Wellcome	ORAL
150MG See		Chronic Obstructive					
Initial or Prolonged	4 DAY	Airways Disease		Salmeterol	C	Glaxo Wellcome	
text		Exacerbated					
Other				Salbutamol	C	Glaxo Wellcome	
50MCG Twice							
per day							
4U As				Salbutamol	C	Glaxo Wellcome	
required							
				Becotide	C	Glaxo Wellcome	
200MCG Twice							
per day							

Date:03/27/01ISR Number: 3689665-XReport Type:Expedited (15-DaCompany Report #B0101861A

Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Depression		Zyban	PS	Glaxo Wellcome	
150MG Twice		Drug Hypersensitivity					
Initial or Prolonged	19 DAY	Dysphagia					
per day							

Eyelid Oedema  
 Face Oedema  
 Gastritis  
 Insomnia  
 Muscle Spasms  
 Pharyngeal Oedema  
 Pruritus  
 Urticaria

Date:03/27/01ISR Number: 3689666-1Report Type:Expedited (15-DaCompany Report #B0101874A

Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	11 DAY	Chronic Obstructive Airways Disease Exacerbated		Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY (INHALATION)	2PUFF Twice			Salmeterol	C	Glaxo Wellcome	
per day RESPIRATORY (INHALATION)	2PUFF Four			Combivent	C		
times per day 20MG per day 400MG Twice per day				Frusemide	C		ORAL
				Theophylline	C		ORAL
				Didronel Pmo	C		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/27/01ISR Number: 3689667-3Report Type:Expedited (15-DaCompany Report #B0101875A

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Delirium		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	2 DAY	Dizziness		Lansoprazole	C		ORAL
15MG per day		Hyperaesthesia		Diazepam	C		ORAL
5MG per day		Nausea		Metoclopramide	C		ORAL
10MG per day		Tongue Oedema		Dicyclomine	C		ORAL
20MG per day				Beclomethasone	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)	200MCG per						
day				Salmeterol	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)	25MCG per day			Montelukast	C		ORAL
10MG per day				Combivent	C		
RESPIRATORY							
(INHALATION)							

Date:03/27/01ISR Number: 3689668-5Report Type:Expedited (15-DaCompany Report #B0101979A

Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Dizziness		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	7 DAY	Feeling Drunk					
Other		Insomnia					
		Vision Blurred					

Date:03/27/01ISR Number: 3689669-7Report Type:Expedited (15-DaCompany Report #B0101980A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Twice		Multiple Sclerosis		Zyban	PS	Glaxo Wellcome	ORAL
per day	24	DAY		Irbesartan	C		ORAL
150MG Per day							

Date:03/27/01ISR Number: 3689670-3Report Type:Expedited (15-DaCompany Report #B0101992A  
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 75MG per day		Psoriasis		Zyban Venlafaxine	PS C	Glaxo Wellcome	ORAL ORAL
12.5MG per day				Captopril	C	Glaxo Wellcome	ORAL
				Co-Proxamol Senna	C C		ORAL ORAL

Date:03/27/01ISR Number: 3689678-8Report Type:Expedited (15-DaCompany Report #B0100993A  
Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged text	4	Asthma Chronic Obstructive Airways Disease		Zyban	PS	Glaxo Wellcome	ORAL
Other 50MCG Twice		Exacerbated		Salmeterol	C	Glaxo Wellcome	
per day				Salbutamol	C	Glaxo Wellcome	
4U As required				Becotide	C	Glaxo Wellcome	
200MCG Twice per day							





FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/27/01ISR Number: 3689682-XReport Type:Expedited (15-DaCompany Report #B0101861A  
 Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Depression		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged per day	19 DAY	Gastritis Hypersensitivity Insomnia Muscle Spasms					

Date:03/27/01ISR Number: 3689683-1Report Type:Expedited (15-DaCompany Report #B0101874A  
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Chronic Obstructive		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	11 DAY	Airways Disease Exacerbated		Salmeterol	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)	2PUFF Twice			Combivent	C		
per day				Frusemide	C		ORAL
RESPIRATORY (INHALATION)	2PUFF Four			Theophylline	C		ORAL
times per day				Didronel Pmo	C		ORAL
20MG per day							
400MG Twice							
per day							

Date:03/27/01ISR Number: 3689684-3Report Type:Expedited (15-DaCompany Report #B0101875A  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Delirium		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	2 DAY	Dizziness		Lansoprazole	C		ORAL
15MG per day		Hyperaesthesia		Diazepam	C		ORAL
5MG per day		Nausea		Metoclopramide	C		ORAL
10MG per day		Tongue Oedema		Dicyclomine	C		ORAL
20MG per day				Beclomethasone	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)	200MCG per						
day				Salmeterol	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)	25MCG per day			Montelukast	C		ORAL
10MG per day				Combivent	C		
RESPIRATORY							
(INHALATION)							

Date:03/27/01ISR Number: 3689685-5Report Type:Expedited (15-DaCompany Report #B0101979A  
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Dizziness		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	7 DAY	Feeling Drunk					
Other		Insomnia					
		Vision Blurred					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/27/01ISR Number: 3689686-7Report Type:Expedited (15-DaCompany Report #B0101980A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Difficulty In Walking		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	24	DAY	Joint Stiffness				
150MG Per day			Multiple Sclerosis	Irbesartan	C		ORAL

Date:03/27/01ISR Number: 3689687-9Report Type:Expedited (15-DaCompany Report #B0101992A  
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Psoriasis		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged				Venlafaxine	C		ORAL
75MG per day							
12.5MG per				Captopril	C	Glaxo Wellcome	ORAL
day							
				Co-Proxamol	C		ORAL
				Senna	C		ORAL

Date:03/27/01ISR Number: 3690482-5Report Type:Expedited (15-DaCompany Report #A0142629A  
Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Akathisia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							
DAY ORAL			Depressed Level Of				
			Professional				
			Consciousness	Multivieamin	C		
			Dissociative Disorder	Calcium Salt	C		
			Restlessness				
			Tremor				

Date:03/27/01ISR Number: 3690667-8Report Type:Direct  
Age:15 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged X 46 TABS Required ORAL		Convulsion Intentional Misuse		Wellbutrin 150mg Glaxo-Wellcome	PS	Glaxo-Wellcome	ORAL
Intervention to Prevent Permanent X 50 TABS Impairment/Damage ORAL		Vomiting		Neurontin 300mg Parke-Davis	SS	Parke-Davis	ORAL
				Celexa	C		

Date:03/27/01ISR Number: 3691039-2Report Type:Expedited (15-DaCompany Report #B0099781A  
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 4 DAY Initial or Prolonged		Haemorrhoids Malaise	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	

Date:03/27/01ISR Number: 3691770-9Report Type:Expedited (15-DaCompany Report #A0142629A  
Age: Gender:Female I/FU:I

Outcome	PT
Other	Akathisia Feeling Abnormal Mental Impairment

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL		Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
			Multivitamin Calcium Salt	C		

Date:03/28/01ISR Number: 3692427-0Report Type:Expedited (15-DaCompany Report #A0140463A  
Age:27 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	10 DAY	Arrhythmia Cardiac Arrest	Foreign Literature	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL		Euphoric Mood Feeling Abnormal Hypersensitivity Myocardial Infarction Palpitations Psychomotor Hyperactivity	Consumer				

Date:03/29/01ISR Number: 3691244-5Report Type:Expedited (15-DaCompany Report #B0101602A  
Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Amnesia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice Hospitalization - per day 9 DAY Initial or Prolonged 1G per day Other		Apnoea Cardio-Respiratory Arrest		Vitamin C	C		ORAL
		Cyanosis Eye Movement Disorder Grand Mal Convulsion Hypoaesthesia		Multivitamins	C		ORAL

Date:03/29/01ISR Number: 3691583-8Report Type:Expedited (15-DaCompany Report #A0142204A  
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Intracranial Aneurysm	Health	Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
14 MON			Professional				

Date:03/29/01ISR Number: 3691593-0Report Type:Expedited (15-DaCompany Report #A0143409A  
Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG Twice		Anaesthesia		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day		Insomnia					
		Sleep Apnoea Syndrome					
		Weight Increased					

Date:03/29/01ISR Number: 3691605-4Report Type:Expedited (15-DaCompany Report #B0099220A  
Age:41 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Agitation Confusional State

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	50 DAY	Grand Mal Convulsion Hyperglycaemia Lactic Acidosis	Foreign	Zyban	PS	Glaxo Wellcome	ORAL
		Leukocytosis	Health Professional	Aspirin	C		
		Thrombocythaemia		Alcohol	C		

Date:03/29/01ISR Number: 3691606-6Report Type:Expedited (15-DaCompany Report #B0099526A  
Age:52 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Astrocytoma	Foreign	Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged			Coma	Health Professional	Prozac	C		
			Dysphagia					
			Glioma					
			Infection					
			Nervous System Disorder					
			Urinary Incontinence					
			Vomiting					

Date:03/29/01ISR Number: 3691615-7Report Type:Expedited (15-DaCompany Report #B0101048A  
Age:66 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150MG Per day	3 DAY	Asthma	Foreign	Zyban	PS	Glaxo Wellcome	ORAL
Disability	1TAB per day		Cyanosis	Health Professional	Maxsoten	C		
Other	100MG per day		Malaise		Ciprofibrate	C		
	150UG per day				Allopurinol	C	Glaxo Wellcome	
	50UG Twice				Salmeterol	C	Glaxo Wellcome	
	per day				Fluticasone	C	Glaxo Wellcome	
	500UG Twice							



per day

Duovent C

2PUFF Three

times per day

Methylprednisolone C

16MG Per day

Date:03/29/01ISR Number: 3691617-0Report Type:Expedited (15-DaCompany Report #B0101365A

Age:48 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 10 DAY	Blood Disorder	Foreign	Zyban	PS	Glaxo Wellcome	
Initial or Prolonged	Blood Pressure Increased Embolism Facial Palsy	Health Professional				

Date:03/29/01ISR Number: 3691635-2Report Type:Expedited (15-DaCompany Report #B0102032A

Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Depression
Initial or Prolonged	Dermatitis Dysgeusia Fatigue Feeling Hot And Cold Headache Injection Site Haemorrhage

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
9 DAY		Nocturia Pruritus Skin Warm Urticaria		Zyban	PS	Glaxo Wellcome	

Date:03/29/01ISR Number: 3691636-4Report Type:Expedited (15-DaCompany Report #B0102048A  
Age:52 YR Gender:Male I/FU:I

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	34 DAY	PT Angina Pectoris Chest Discomfort		Zyban	PS	Glaxo Wellcome	ORAL
5MG per day				Bendrofluazide	C	Glaxo Wellcome	ORAL

Date:03/29/01ISR Number: 3691637-6Report Type:Expedited (15-DaCompany Report #B0102049A  
Age:69 YR Gender:Male I/FU:I

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		PT Blood Viscosity Increased Dermatitis Joint Swelling Muscular Weakness Night Sweats Serum Sickness		Zyban	PS	Glaxo Wellcome	ORAL

Date:03/29/01ISR Number: 3691638-8Report Type:Expedited (15-DaCompany Report #B0102062A  
Age: Gender:Male I/FU:I

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
150MG Per day Initial or Prolonged	17 DAY	PT Blood Creatine Phosphokinase Increased		Zyban	PS	Glaxo Wellcome	ORAL

Dyspnoea

Date:03/29/01ISR Number: 3691639-XReport Type:Expedited (15-DaCompany Report #B0102063A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	37 DAY		Subarachnoid Haemorrhage	Zyban	PS	Glaxo Wellcome	ORAL

Date:03/29/01ISR Number: 3691640-6Report Type:Expedited (15-DaCompany Report #B0102228A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Twice		Erythema Multiforme	Zyban	PS	Glaxo Wellcome	ORAL
Other	per day	29 DAY	Psoriasis				

Date:03/29/01ISR Number: 3691641-8Report Type:Expedited (15-DaCompany Report #B0102229A  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Twice		Asthma	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	per day	5 DAY		Terbutaline	C		

RESPIRATORY  
(INHALATION)

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/01ISR Number: 3691642-XReport Type:Expedited (15-DaCompany Report #B0102251A

Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG As		Circulatory Collapse	Zyban	PS	Glaxo Wellcome	
directed			Malaise				
			Ventricular Fibrillation	Homeopathic Drug	C		

Date:03/30/01ISR Number: 3692352-5Report Type:Expedited (15-DaCompany Report #254325

Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	108 DAY		Grand Mal Convulsion	Accutane Capsules	PS	Roche	
	123 DAY			Wellbutrin	SS		
				Prozac	C		

Date:03/30/01ISR Number: 3692576-7Report Type:Expedited (15-DaCompany Report #B0087305A

Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Alcohol Interaction	Zyban	PS	Glaxo Wellcome	ORAL
			Completed Suicide	Alcohol	I		
			Drowning				
			Medication Error				

Date:03/30/01ISR Number: 3692580-9Report Type:Expedited (15-DaCompany Report #B0099756A

Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Alcohol Interaction	Zyban	PS	Glaxo Wellcome	ORAL
			Drowning	Alcohol	SS		
			Medication Error				

Date:03/30/01ISR Number: 3693047-4Report Type:Direct  
Age:34 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diplopia Dizziness		Comtrex Maximum Bristol-Myers Squibb	PS	Bristol-Myers Squibb	ORAL
2 TABS Q		Nausea					
12HR PO		Vision Blurred		Wellbutrin Sr 150mg Glaxo	SS	Glaxo	ORAL
150MG QAM							
PO							

Date:03/31/01ISR Number: 3693549-0Report Type:Expedited (15-DaCompany Report #B0101979A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Feeling Drunk	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Disability Required		Insomnia					
150 MG /PER		Vision Blurred					
Intervention to DAY/ORAL							
Prevent Permanent Impairment/Damage							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/31/01ISR Number: 3693550-7Report Type:Expedited (15-DaCompany Report #B0101992A  
Age:72 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG /ORAL	Psoriasis	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
			Venlafaxine Hydrochloride	C		
			Captopril	C		
			Co-Proxamol	C		
			Sennosides	C		

Date:03/31/01ISR Number: 3693551-9Report Type:Expedited (15-DaCompany Report #B0101861A  
Age:31 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged UNKNOWN PER DAY	Depression Dysphagia Eyelid Oedema Face Oedema Gastritis Hypersensitivity Insomnia Muscle Spasms Pharyngeal Oedema Pruritus Urticaria	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
	150 MG/TWICE					

Date:03/31/01ISR Number: 3693552-0Report Type:Expedited (15-DaCompany Report #B0101875A  
Age:48 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to 150 MG/PER Prevent Permanent DAY/ORAL	Angioneurotic Oedema Delirium Dizziness	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Impairment/Damage

Hyperaesthesia

Tongue Oedema

Lansoprazole

C

Diazepam

C

Metoclopramide

C

Dicyclomine

C

Beclomethasone

Dipropion

C

Salmeterol Xinafoate

C

Montelukast Sodium

C

Combivent

C

Date:03/31/01ISR Number: 3693553-2Report Type:Expedited (15-DaCompany Report #B0101980A

Age:49 YR Gender:Male

I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Difficulty In Walking Joint Stiffness	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE PER DAY/ORAL		Multiple Sclerosis		Irbesartan	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/31/01ISR Number: 3693554-4Report Type:Expedited (15-DaCompany Report #B0100993A  
 Age:57 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/SEE	Asthma Chronic Obstructive	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Required TEXT/ORAL Intervention to Prevent Permanent Impairment/Damage	Airways Disease Exacerbated	Professional	Salmeterol Xinafoate Salbutamol Sulphate Beclomethasone Dipropion	C C C		

Date:03/31/01ISR Number: 3693561-1Report Type:Expedited (15-DaCompany Report #B0101874A  
 Age:72 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG, TWICE	Chronic Obstructive Airways Disease Exacerbated	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS	Glaxo Wellcome Inc	ORAL
PER DAY, ORAL			Salmeterol Xinafoate Combivent Frusemide Theophylline Disodium Etidronate	C C C C C		

Date:04/02/01ISR Number: 3693386-7Report Type:Expedited (15-DaCompany Report #A0141624A  
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Brain Damage Convulsion Thinking Abnormal		Wellbutrin	PS	Glaxo Wellcome	ORAL



Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Arrhythmia		Zyban	PS	Glaxo Wellcome	ORAL
10.5G per day 18 DAY						
Hospitalization -	Blood Alcohol Increased		Dafalgan	C		
10TAB Per day						
Initial or Prolonged	Brain Oedema		Alcohol	C		
Other	Cardiac Arrest					
	Coagulation Factor V					
	Level Decreased					
	Coma					
	Condition Aggravated					
	Convulsion					
	Electroencephalogram					
	Abnormal					
	Encephalopathy					
	Hallucination					
	Hepatic Failure					
	Intentional Misuse					
	Prothrombin Level					
	Decreased					
	Suicide Attempt					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/02/01ISR Number: 3693398-3Report Type:Expedited (15-DaCompany Report #B0100368A  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anxiety Dermatitis Difficulty In Walking Dyspnoea Lip Disorder Swelling		Zyban	PS	Glaxo Wellcome	

Date:04/02/01ISR Number: 3693402-2Report Type:Expedited (15-DaCompany Report #B0102234A  
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Twice Other per day	22 DAY	Bronchospasm Insomnia		Zyban	PS	Glaxo Wellcome	ORAL
3MG per day				Warfarin	C	Glaxo Wellcome	ORAL
40MG per day				Frusemide	C		ORAL
5MG per day				Lisinopril	C		ORAL
10MG per day				Simvastatin	C		ORAL

Date:04/02/01ISR Number: 3694389-9Report Type:Expedited (15-DaCompany Report #254325  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 40 MG 2 PER DAY; ORAL		Grand Mal Convulsion Loss Of Consciousness	Health Professional	Accutane	PS	Hlr Technology	ORAL
25 MG 2 PER				Wellbutrin (Bupropion Hydrochloride)	SS		ORAL

DAY; ORAL

Prozac (Fluoxetine  
Hydrochloride) C

Date:04/02/01ISR Number: 3694567-9Report Type:Expedited (15-DaCompany Report #B0102048A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Angina Pectoris	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Intervention to Prevent Permanent Impairment/Damage				Bendrofluazide	C		

Date:04/02/01ISR Number: 3694568-0Report Type:Expedited (15-DaCompany Report #A0143409A  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hypoaesthesia Insomnia	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE		Sleep Apnoea Syndrome					
PER DAY/ORAL		Weight Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/02/01ISR Number: 3694569-2Report Type:Expedited (15-DaCompany Report #B0102228A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required 150MG /TWICE Intervention to PER DAY/ORAL Prevent Permanent Impairment/Damage		Erythema Multiforme Psoriasis	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:04/02/01ISR Number: 3694570-9Report Type:Expedited (15-DaCompany Report #B0102229A  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/TWICE PER DAY/ORAL		Asthma	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Terbutaline	C		

Date:04/02/01ISR Number: 3694571-0Report Type:Expedited (15-DaCompany Report #B0102251A  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening UNKNOWN DIRECTRED	150 MG / AS	Circulatory Collapse Condition Aggravated Ventricular Fibrillation	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
				Homeopathic Medication	C		

Date:04/02/01ISR Number: 3694572-2Report Type:Expedited (15-DaCompany Report #B0102063A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Headache Subarachnoid Haemorrhage	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /ORAL							

Date:04/02/01ISR Number: 3694573-4Report Type:Expedited (15-DaCompany Report #B0102049A  
Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG / Prevent Permanent TWICE PER DAY Impairment/Damage / ORAL		Blood Viscosity Increased Dermatitis  Joint Swelling  Muscular Weakness  Night Sweats Serum Sickness	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:04/02/01ISR Number: 3694574-6Report Type:Expedited (15-DaCompany Report #B0102062A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG /PER  DAY/ ORAL		Blood Creatine Phosphokinase Increased  Dyspnoea	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/02/01ISR Number: 3694575-8Report Type:Expedited (15-DaCompany Report #B0102032A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged UNKNOWN	150 MG	9 DAY	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
				Fatigue Feeling Hot And Cold Headache Injection Site Haemorrhage Rash Pruritic Urticaria			

Date:04/02/01ISR Number: 3694755-1Report Type:Expedited (15-DaCompany Report #A0142204A

Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL			Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
			Health Professional				

Date:04/02/01ISR Number: 3694757-5Report Type:Expedited (15-DaCompany Report #B0101048A

Age:66 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / PER Initial or Prolonged DAY / ORAL			Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Disability Required Intervention to Prevent Permanent Impairment/Damage			Health Professional	Maxsoten Ciprofibrate Allopurinol Salmeterol Xinafoate Fluticasone Propionate Duovent Methylprednisolone	C C C C C C C C		
				Asthma Bronchospasm Cyanosis Dyspnoea Malaise			

Date:04/02/01ISR Number: 3694758-7Report Type:Expedited (15-DaCompany Report #B0101365A  
Age:48 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG Initial or Prolonged	Blood Pressure Increased Coagulopathy Embolism Facial Palsy	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	

Date:04/02/01ISR Number: 3694759-9Report Type:Expedited (15-DaCompany Report #B0099526A  
Age:52 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death AS DIRECTED / Hospitalization - ORAL Initial or Prolonged	Coma Dysphagia Eating Disorder Gliosis Infection Memory Impairment Urinary Incontinence Vomiting	Foreign Health Professional	Zyban Fluoxetine Hydrochloride	PS C	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/02/01ISR Number: 3694761-7Report Type:Expedited (15-DaCompany Report #B0099220A  
Age:41 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / PER Initial or Prolonged DAY / ORAL	Agitation	Foreign Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
	Grand Mal Convulsion	Professional	Aspirin	C		
	Hyperglycaemia		Ethanol	C		
	Lactic Acidosis					
	Leukocytosis					
	Nervousness					
	Salivary Hypersecretion					
	Thrombocythaemia					
	Urinary Incontinence					

Date:04/02/01ISR Number: 3694825-8Report Type:Expedited (15-DaCompany Report #B0101602A  
Age:40 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150 MG/TWICE Hospitalization - PER DAY/ORAL Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage	Amnesia		Zyban	PS	Glaxo Wellcome Inc	ORAL
	Balance Disorder					
	Cardio-Respiratory Arrest		Ascorbic Acid	C		
	Cyanosis		Multivitamin	C		
	Dizziness					
	Eye Rolling					
	Fall					
	Grand Mal Convulsion					
	Hypoaesthesia					
	Screaming					

Date:04/03/01ISR Number: 3694452-2Report Type:Expedited (15-DaCompany Report #A0143055A  
Age:64 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 100MG Per day YR	Arthralgia		Wellbutrin	PS	Glaxo Wellcome	ORAL



150MG In the		Dizziness		Wellbutrin Sr	SS	Glaxo Wellcome	ORAL
morning	YR	Ear Discomfort					
		Hearing Impaired		Hormones	C		
		Hyperhidrosis					
		Hypersensitivity					
		Pharyngeal Oedema					
		Tinnitus					

Date:04/03/01ISR Number: 3694459-5Report Type:Expedited (15-DaCompany Report #B0100069A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	5 DAY	Anxiety	Foreign	Zyban	PS	Glaxo Wellcome	
Initial or Prolonged		Chest Pain	Health	Sertraline	C		
		Gastrooesophageal Reflux	Professional				
		Disease					
		Hyperventilation					
		Malaise					
		Nausea					
		Pallor					
		Tremor					
		Vomiting					
		Weight Decreased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/03/01ISR Number: 3694461-3Report Type:Expedited (15-DaCompany Report #B0100785A  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression	Foreign	Zyban	PS	Glaxo Wellcome	ORAL
8 DAY		Crying	Health				
		Depression	Professional				
		Diarrhoea					
		Diplopia					
		Disturbance In Attention					
		Dizziness					
		Eye Pain					
		Fatigue					
		Headache					
		Retching					
		Suicidal Ideation					
		Vomiting					

Date:04/03/01ISR Number: 3694465-0Report Type:Expedited (15-DaCompany Report #B0102365A  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Circulatory Collapse		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Convulsion					
Hospitalization -	18 DAY	Loss Of Consciousness		Salbutamol	C	Glaxo Wellcome	
per day		Metabolic Acidosis					
Initial or Prolonged		Respiratory Acidosis		Medroxyprogesterone	C		ORAL
RESPIRATORY				Oestrone	C		ORAL
(INHALATION)							
5MG Per day							
.625MG Per							
day							

Date:04/03/01ISR Number: 3694468-6Report Type:Expedited (15-DaCompany Report #B0102451A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Tremor Vision Blurred		Zyban	PS	Glaxo Wellcome	

Date:04/03/01ISR Number: 3694469-8Report Type:Expedited (15-DaCompany Report #B0102453A  
 Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	6 WK	Convulsion  Haemorrhage Salivary Hypersecretion Sudden Death		Zyban	PS	Glaxo Wellcome	

Date:04/03/01ISR Number: 3694470-4Report Type:Expedited (15-DaCompany Report #B0102456A  
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day Initial or Prolonged 300MG Per day		Peripheral Ischaemia		Zyban	PS	Glaxo Wellcome	ORAL
TOPICAL				Quinine	C		ORAL
TOPICAL	2UNIT per day			Latanoprost	C		
				Carteolol	C		
2.5MG per day				Calcium Carbonate	C		ORAL
1MG Per day				Bendrofluazide	C	Glaxo Wellcome	ORAL
				Sunerven	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/03/01ISR Number: 3694471-6Report Type:Expedited (15-DaCompany Report #B0102461A  
Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	8 DAY	Dyspnoea					
200MG Per day		Heart Rate Irregular		Diltiazem	C		ORAL
		Pallor					

Date:04/03/01ISR Number: 3694472-8Report Type:Expedited (15-DaCompany Report #B0102462A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice		Bronchospasm		Zyban	PS	Glaxo Wellcome	ORAL
per day	20 DAY	Face Oedema					
300MG Per day		Pruritus		Allopurinol	C	Glaxo Wellcome	ORAL
		Urticaria					

Date:04/03/01ISR Number: 3696703-7Report Type:Expedited (15-DaCompany Report #B0087305A  
Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death ORAL		Alcohol Interaction	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Blood Alcohol Increased	Literature	Ethanol (Formulation			
		Completed Suicide	Health	Unknown) (Alcohol)	SS		
		Drowning	Professional				
		Drug Abuser					
		Drug Level Above					
		Therapeutic					

Date:04/03/01ISR Number: 3696706-2Report Type:Expedited (15-DaCompany Report #B0099756A  
Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alcohol Interaction Drowning	Foreign Literature	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL				Ethanol (Formulation Unknown) (Alcohol)	SS		

Date:04/04/01ISR Number: 3695150-1Report Type:Expedited (15-DaCompany Report #A0143570A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice							
Hospitalization - per day 9 DAY		Overdose					
Initial or Prolonged 40MG Per day				Celexa	C		ORAL
				Buspar	C		ORAL
15MG Twice							
per day							
250MG Per day				Lamisil	C		ORAL
				Zyrtec	C	Glaxo Wellcome	ORAL
10MG Per day				Prevacid	C		ORAL
30MG Twice							
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/04/01ISR Number: 3695152-5Report Type:Expedited (15-DaCompany Report #A0143899A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:04/04/01ISR Number: 3695161-6Report Type:Expedited (15-DaCompany Report #B0102052A

Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day 8 DAY	Dizziness Eye Pain Headache Joint Stiffness Paraesthesia Tinnitus Tremor		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/04/01ISR Number: 3695165-3Report Type:Expedited (15-DaCompany Report #B0102458A

Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day 41 DAY	Asthma Lower Respiratory Tract Infection Sudden Death		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/05/01ISR Number: 3698516-9Report Type:Direct

Company Report #

Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Headache Vomiting		Saint Johns Wort / 300mg Wellbutrin 150mg	PS SS		

Date:04/05/01ISR Number: 3698808-3Report Type:Direct  
Age:29 YR Gender:Male I/FU:I

Company Report #USP 53877

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Zyban	PS	Catalytica Pharmaceuticals/Glax o-Wellcome	

Date:04/05/01ISR Number: 3698868-XReport Type:Expedited (15-DaCompany Report #B0102234A  
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Required		Bronchospasm Insomnia	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE Intervention to PER DAY/ ORAL Prevent Permanent Impairment/Damage				Warfarin Sodium Frusemide Lisinopril Simvastatin	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/05/01ISR Number: 3698872-1Report Type:Expedited (15-DaCompany Report #B0100368A  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG		Anxiety Asthenia	Foreign Literature	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
		Difficulty In Walking Dyspnoea Lip Disorder Oxygen Saturation Decreased Rash Generalised Swelling	Consumer				

Date:04/05/01ISR Number: 3699658-4Report Type:Expedited (15-DaCompany Report #A0141624A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Brain Damage	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
		Convulsion Decreased Activity Thinking Abnormal					

Date:04/05/01ISR Number: 3699746-2Report Type:Expedited (15-DaCompany Report #B0095016A  
Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG / ORAL		Arrhythmia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage		Brain Oedema Cardiac Arrest Coagulation Factor V Level Decreased Coma Convulsion Electroencephalogram Abnormal Encephalopathy	Health Professional	Paracetamol Ethanol	C C		



Hallucination  
Hepatic Failure  
Intentional Misuse  
Prothrombin Level  
Decreased  
Suicide Attempt

Date:04/06/01ISR Number: 3700116-9Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to 1 FOR THEN Prevent Permanent BID ORAL Impairment/Damage		Appetite Disorder Dizziness Dyspepsia Fatigue Headache Insomnia Malaise Nausea Palpitations Restlessness		Wellbutrin Sr / 150mg	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/06/01ISR Number: 3700201-1Report Type:Expedited (15-DaCompany Report #A0143055A  
 Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100 MG / PER DAY / ORAL		Arthralgia Deafness Dizziness Hyperhidrosis Pain In Extremity Pharyngeal Oedema	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
150 MG / IN THE MORNING / ORAL		Tinnitus		Wellbutrin Tablet - Controlled Release (Bupropion Hydrochloride)	SS		ORAL
				Hormones	C		

Date:04/06/01ISR Number: 3700334-XReport Type:Expedited (15-DaCompany Report #B0102451A  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150 MG		Tremor Vision Blurred	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:04/06/01ISR Number: 3700370-3Report Type:Expedited (15-DaCompany Report #B0102453A  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG	6 WK	Convulsion Haemorrhage Salivary Hypersecretion Sudden Death	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG PER DAY ORAL		Peripheral Ischaemia	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Quinine	C		
				Latanoprost	C		
				Carteolol	C		
				Calcium Carbonate	C		
				Bendrofluazide	C		
				Sunerven	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/TWICE PER DAY/ORAL		Arthralgia Dyspnoea Heart Rate Irregular Pallor	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Diltiazem	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/06/01ISR Number: 3700420-4Report Type:Expedited (15-DaCompany Report #B0102462A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Bronchospasm	Foreign	Bupropion			
Intervention to		Face Oedema	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
Prevent Permanent		Pruritus	Professional				
PER DAY ORAL							
Impairment/Damage		Urticaria		Allopurinol	C		

Date:04/06/01ISR Number: 3700496-4Report Type:Expedited (15-DaCompany Report #B0102365A  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Circulatory Collapse	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
Hospitalization -		Convulsion					
TWICE PER DAY							
Initial or Prolonged		Loss Of Consciousness					
/ ORAL							
		Metabolic Acidosis		Salbutamol Sulphate	C		
		Respiratory Acidosis		Medroxyprogesterone	C		
				Oestrone	C		

Date:04/06/01ISR Number: 3700499-XReport Type:Expedited (15-DaCompany Report #B0100785A  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aggression	Foreign	Bupropion			
ORAL		Crying	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Depression	Professional				
		Depression Suicidal					
		Diarrhoea					
		Diplopia					
		Disturbance In Attention					
		Dizziness					
		Eye Pain					
		Fatigue					

Headache  
Nausea  
Retching  
Vomiting

Date:04/06/01ISR Number: 3700500-3Report Type:Expedited (15-DaCompany Report #B0100069A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG		Abdominal Pain Chest Pain	Foreign Literature	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
		Gastrooesophageal Reflux Disease Hyperventilation Malaise Tremor Vomiting	Health Professional	Sertraline	C		

Date:04/06/01ISR Number: 3700868-8Report Type:Periodic Company Report #990926.01  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2.5 MG QD,		Cough	Health Professional	Indapamide	PS	Mylan Pharmaceuticals Inc	ORAL

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ORAL		Allopurinol	SS	
		Beconase	SS	
OCCLUSIVE				
DRESSING	2 PUFFS UD,			
ORAL		Cozaar	SS	ORAL
50 MG QD.				
ORAL		Glucophage	SS	ORAL
1GRAM				
TID, ORAL		Lipitor	SS	ORAL
10 MG, QD,				
ORAL		Rezulin	SS	ORAL
200MG, TID,				
ORAL		Trental	SS	ORAL
400MG TID,				
ORAL		Welbutrin	SS	ORAL
150MG, QD,				
ORAL		Glipizide	SS	ORAL
10MG, BID,				
OROAL		Quinine	SS	ORAL
325MG, TID,				
ORAL				

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	1TAB Twice	Face Crushing		Wellbutrin	PS	Glaxo Wellcome	ORAL
Hospitalization -	per day	Grand Mal Convulsion					
Initial or Prolonged	250MG Per day	Memory Impairment		Antabuse	C		ORAL
Other	20MG Per day	Peripheral Ischaemia		Prozac	C		ORAL
		Skin Disorder					

Date:04/09/01ISR Number: 3699940-0Report Type:Expedited (15-DaCompany Report #A0143021A  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG See	Confusional State		Wellbutrin	PS	Glaxo Wellcome	ORAL
Hospitalization -	text	Drug Toxicity					
Initial or Prolonged	100MG Three	Fall		Amantadine	SS		ORAL
times per day	8 MON	Limb Injury					
		Schizophrenia		Prozac	C		
		Tremor		Loxapine	C		
				Seroquel	C		
				Risperdal	C		

Date:04/09/01ISR Number: 3699941-2Report Type:Expedited (15-DaCompany Report #A0143253A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day DAY	Pancreatic Disorder		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged							

Date:04/09/01ISR Number: 3699996-5Report Type:Expedited (15-DaCompany Report #B0102580A  
Age:40 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Chest Pain
Initial or Prolonged	Dyspnoea





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Freedom Of Information (FOI) Report

		Triple Vessel Bypass Graft				
Dose	Duration		Report Source	Product	Role	Manufacturer
150MG Twice per day	6 WK			Zyban	PS	Glaxo Wellcome
						Route
						ORAL

Date:04/09/01ISR Number: 3699998-9Report Type:Expedited (15-DaCompany Report #B0102924A  
 Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG As Initial or Prolonged directed	11 DAY	Agitation Anxiety		Zyban	PS	Glaxo Wellcome	
RESPIRATORY (INHALATION)		Blood Pressure Increased		Steroids	C		
		Convulsion					
		Depressed Level Of Consciousness Diplopia Feeling Abnormal Haemorrhagic Stroke Headache Hemianopia Memory Impairment Respiratory Disorder Urinary Incontinence Visual Acuity Reduced		Cyproterone Acetate	C		

Date:04/09/01ISR Number: 3700304-1Report Type:Expedited (15-DaCompany Report #B0091784A  
 Age:65 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice per day		Abnormal Dreams		Zyban	PS	Glaxo Wellcome	
		Chronic Obstructive					

250MG Four  
 times per day 10 DAY  
 Airways Disease  
 Exacerbated  
 Drug Interaction  
 Emphysema  
 Hallucination  
 Lower Respiratory Tract  
 Infection  
 Lymphadenopathy

Erythromycin SS Glaxo Wellcome

Date:04/09/01ISR Number: 3700305-3Report Type:Expedited (15-DaCompany Report #B0094748A  
 Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 32 DAY		Erythema		Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY (INHALATION)	40MCG Three times per day			Ipratropium	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)	500MCG Twice per day			Fluticasone	C	Glaxo Wellcome	
2TAB As required				Norethisterone Acetate + Oestradiol + Oestriol	C		ORAL
				Dextropropoxyphene + Paracetamol	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/09/01ISR Number: 3700311-9Report Type:Expedited (15-DaCompany Report #B0100686A

Age:55 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Bone Disorder		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Delirium		Ssri	C		
	Post Procedural		Zaleplon	C		
	Complication		Diazepam	C		
5G Twice per						
day						
			Sevoflurane	C		
			Propofol	C		
			Fentanyl	C		
			Rocuronium	C		
			Atropine	C		
			Bupivacaine	C		

Date:04/09/01ISR Number: 3700315-6Report Type:Expedited (15-DaCompany Report #B0102788A

Age:42 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Disability	Psoriasis		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice						
Other	Rash Pustular					
per day						

Date:04/09/01ISR Number: 3700316-8Report Type:Expedited (15-DaCompany Report #B0102951A

Age:45 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Other	Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice						
per day	Depression					
4						
DAY	Palpitations		Perindopril	C		ORAL
4MG Per day						
			Nicorandil	C		ORAL
20MG Twice						

per day		Lansoprazole	C		ORAL
15MG Per day		Isosorbide	C		ORAL
30MG Per day		Felodipine	C		
UNKNOWN	10MG Per day	Atorvastatin	C		
UNKNOWN	30MG Per day	Gaviscon	C		
UNKNOWN		Aspirin	C		
UNKNOWN	75MG Per day	Beclomethasone	C	Glaxo Wellcome	
RESPIRATORY					
(INHALATION)	50MCG Twice				
per day		Salbutamol	C	Glaxo Wellcome	
RESPIRATORY					
(INHALATION)					

Date:04/09/01ISR Number: 3700318-1Report Type:Expedited (15-DaCompany Report #B0103067A  
Age:60 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Per day 3 DAY Initial or Prolonged	Cardiac Failure		Quomem	PS	Glaxo Wellcome	ORAL

Date:04/09/01ISR Number: 3700954-2Report Type:Expedited (15-DaCompany Report #A0143570A  
Age:33 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death 150 MG, TWICE Hospitalization - PER DAY, ORAL 9 DAY Initial or Prolonged	Completed Suicide  Overdose	Health  Professional	Wellbutrin Sr  Citalopram	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrobromide	C
Buspiurone	
Hydrochloride	C
Terbinafine	C
Cetirizine	
Hydrochloride	C
Lansoprazole	C

Date:04/09/01ISR Number: 3700968-2Report Type:Expedited (15-DaCompany Report #A0143253A  
 Age:39 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Pancreatic Disorder	Health Professional	Bupropion Hydrochloride - Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG, PER DAY, ORAL						

Date:04/09/01ISR Number: 3700972-4Report Type:Expedited (15-DaCompany Report #A0143021A  
 Age:53 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death Hospitalization - Initial or Prolonged	Confusional State Drug Toxicity Fall Hallucination, Auditory	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS	Glaxo Wellcome Inc	ORAL
150 MG, SEE TEXT, ORAL	Injury					
100 MG, THREE TIMES PER DAY, ORAL	Schizophrenia Surgical Procedure Repeated Tremor		Amantadine Tablet (Amantadine Hydrochloride)	SS		ORAL
	8 MON					

Fluoxetine	
Hydrochloride	C
Loxapine	C
Quetiapine Fumarate	C

Risperidone

C

Date:04/09/01ISR Number: 3701006-8Report Type:Expedited (15-DaCompany Report #A0143899A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL							

Date:04/09/01ISR Number: 3701201-8Report Type:Expedited (15-DaCompany Report #A0141931A  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Angiopathy	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
1 TABLET / Hospitalization - TWICE PER DAY		Electroencephalogram	Professional				
Initial or Prolonged / ORAL		Abnormal					
Required Intervention to Prevent Permanent Impairment/Damage		Grand Mal Convulsion		Disulfiram	C		
		Memory Impairment		Fluoxetine			
		Skin Injury		Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/09/01ISR Number: 3701485-6Report Type:Expedited (15-DaCompany Report #B0102052A  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG / PER DAY/ ORAL		Blood Pressure Diastolic Increased	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Prevent Permanent Impairment/Damage		Dizziness Eye Pain Headache Joint Stiffness Paraesthesia Tinnitus Tremor					

Date:04/09/01ISR Number: 3701486-8Report Type:Expedited (15-DaCompany Report #B0102458A  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG/ PER DAY/ORAL		Asthma Lower Respiratory Tract Infection Sudden Death	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:04/09/01ISR Number: 3701489-3Report Type:Expedited (15-DaCompany Report #B0102580A  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/TWICE PER DAY/ORAL		Chest Pain Dyspnoea Triple Vessel Bypass Graft	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:04/09/01ISR Number: 3701495-9Report Type:Expedited (15-DaCompany Report #B0102924A  
Age:68 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Agitation	Foreign	Bupropion			
Initial or Prolonged	Anxiety	Health	Hydrochloride	PS	Glaxo Wellcome Inc	
UNKNOWN	150 MG/AS					
	Blood Pressure Increased	Professional				
DIRECTED/						
	Convulsion		Steroid	C		
	Diplopia		Cyproterone Acetate	C		
	Dyspnoea					
	Feeling Abnormal					
	Haemorrhagic Stroke					
	Headache					
	Hemianopia					
	Hypertension					
	Memory Impairment					
	Urinary Incontinence					
	Visual Acuity Reduced					

Date:04/10/01ISR Number: 3701093-7Report Type:Expedited (15-DaCompany Report #A0144245A  
Age:45 YR Gender:Male I/FU:F

Outcome	PT
Other	Choroidal Detachment
	Conjunctival Hyperaemia
	Eye Disorder



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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Eye Pain Intraocular Pressure Increased					
		Malaise		Zyban Cozaar	PS C	Glaxo Wellcome	ORAL

Date:04/10/01ISR Number: 3701094-9Report Type:Expedited (15-DaCompany Report #A0144419A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Neuropathy Peripheral		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL

Date:04/10/01ISR Number: 3701105-0Report Type:Expedited (15-DaCompany Report #B0101861A  
Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	19 DAY	Depression Drug Hypersensitivity	Health Professional	Zyban	PS	Glaxo Wellcome	
		Dysphagia Dyspnoea Eyelid Oedema Face Oedema Gastritis Insomnia Laryngeal Oedema Muscle Spasms Pharyngeal Oedema Pruritus Urticaria					

Date:04/10/01ISR Number: 3701106-2Report Type:Expedited (15-DaCompany Report #B0102451A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Disability  
 300MG Twice  
 per day

Asthenia  
 Disturbance In Attention  
 Dry Mouth  
 Overdose  
 Paraesthesia  
 Tremor  
 Vision Blurred

Health  
 Professional

Zyban  
 PS  
 Glaxo Wellcome

Date:04/10/01  
 Age:29 YR  
 Gender:Female  
 I/FU:I

ISR Number: 3703146-6  
 Report Type:Direct  
 Company Report #

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 BID PO		Agitation		Zyban(R), Bupropriion	PS		ORAL
		Depression Hallucination Tremor		Demulin 1/35	C		

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Freedom Of Information (FOI) Report

Date:04/11/01ISR Number: 3702378-0Report Type:Expedited (15-DaCompany Report #A0139749A

Age:33 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Accidental Overdose		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Astrocytoma		Wellbutrin Sr	SS	Glaxo Wellcome	ORAL
150MG Twice						
	Body Temperature					
per day						
	Increased		Klonopin	SS		
6 MON						
	Grand Mal Convulsion		Luvox	SS		
6 MON						
	Loss Of Consciousness		Triphasil	C		
	Ocular Hyperaemia		Aspirin	C		
			Biotin	C		
			Vitamin C	C		
			Zinc	C		
			Mvi	C		

Date:04/11/01ISR Number: 3702388-3Report Type:Expedited (15-DaCompany Report #A0144603A

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Other	Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice						
	Asthma					
per day						
	Bronchospasm					
	Chest Pain					
	Dermatitis					
	Face Oedema					
	Insomnia					
	Myocardial Infarction					
	Pruritus					
	Urticaria					

Date:04/11/01ISR Number: 3702391-3Report Type:Expedited (15-DaCompany Report #A0144702A

Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Dyspnoea		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	8 DAY	Feeling Jittery Lethargy Oedema Peripheral Rash Erythematous Throat Tightness		Ortho Novum 777	C		

Date:04/11/01ISR Number: 3702397-4Report Type:Expedited (15-DaCompany Report #B0098194A  
Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Per day	16 DAY	Bronchopneumonia		Zyban	PS	Glaxo Wellcome	ORAL
		Ecchymosis Gastritis Erosive Hepatic Congestion Hypotension Intracranial Pressure Increased Loss Of Consciousness Pleural Haemorrhage Pulmonary Congestion Respiratory Distress Syncope Vision Blurred		Chloroquine Diphenhydramine	C C		

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Freedom Of Information (FOI) Report

Date:04/11/01ISR Number: 3702413-XReport Type:Expedited (15-DaCompany Report #B0103161A

Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Blister		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	25 DAY	Dermatitis		Doxepin	C		ORAL
10MG Per day		Psoriasis		Norimin	C		ORAL
				Acitretin	C		
25MG Two times per week				Trandolapril	C		ORAL
2MG Per day				Panadeine Forte	C		

Date:04/11/01ISR Number: 3702415-3Report Type:Expedited (15-DaCompany Report #B0103201A

Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Per day	13 DAY	Lichen Planus		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/11/01ISR Number: 3702416-5Report Type:Expedited (15-DaCompany Report #B0103209A

Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Twice		Dizziness		Zyban	PS	Glaxo Wellcome	ORAL
per day	2 DAY	Influenza Like Illness					
135MG Per day		Myalgia		Mebeverine	C		ORAL

Date:04/11/01ISR Number: 3702417-7Report Type:Expedited (15-DaCompany Report #B0103210A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day	30 DAY	Joint Swelling		Co-Dydramol	C	Glaxo Wellcome	ORAL
		Medication Error		Omeprazole	C		ORAL
20MG Per day		Serum Sickness					
		Urticaria					

Date:04/11/01ISR Number: 3702419-0Report Type:Expedited (15-DaCompany Report #B0103211A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Antinuclear Antibody		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Positive					
per day	32 DAY	Arthralgia					
		Autoimmune Disorder					
		Mouth Ulceration					
		Rash Pruritic					

Date:04/11/01ISR Number: 3702422-0Report Type:Expedited (15-DaCompany Report #B0103453A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Optic Disc Disorder		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Retinopathy					
Other							
per day	29 DAY			Premique	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/11/01ISR Number: 3702423-2Report Type:Expedited (15-DaCompany Report #B0103470A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hemiplegia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Single							
dose	1	DAY					
		Nausea					
		Palpitations					

Date:04/11/01ISR Number: 3702424-4Report Type:Expedited (15-DaCompany Report #B0103473A

Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Oedema Peripheral		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Initial or Prolonged		Purpura					
per day	45	DAY					

Date:04/11/01ISR Number: 3702426-8Report Type:Expedited (15-DaCompany Report #B0103477A

Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Headache		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Hemiparesis		Gaviscon	C		ORAL
10ML Twice							
per day		Nystagmus					
				Omeprazole	C		

Date:04/11/01ISR Number: 3702427-XReport Type:Expedited (15-DaCompany Report #B0103481A

Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							

per day 56 DAY

50MG Per day

75MG Per day

SUBLINGUAL

150MG Twice

per day

Atenolol C ORAL

Aspirin C ORAL

Glyceryl Trinitrate C Glaxo Wellcome

Ranitidine C Glaxo Wellcome ORAL

Date:04/11/01ISR Number: 3702428-1Report Type:Expedited (15-DaCompany Report #B0103576A

Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety Atrial Fibrillation Sleep Disorder		Zyban	PS	Glaxo Wellcome	

Date:04/11/01ISR Number: 3702429-3Report Type:Expedited (15-DaCompany Report #B0103584A

Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -			Haemorrhagic Stroke	Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day 2 WK							
Initial or Prolonged		Hemiparesis		Carvedilol	C	Glaxo Wellcome	ORAL
25MG Per day							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/11/01ISR Number: 3702430-XReport Type:Expedited (15-DaCompany Report #D0013784A  
 Age:67 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	Atrial Fibrillation		Zyban	PS	Glaxo Wellcome	ORAL
Life-Threatening Hospitalization -		Bundle Branch Block Right Cardiac Failure		Dynacil	C		ORAL
				Digimerck	C		ORAL
Initial or Prolonged Other		Nervous System Disorder Pleural Effusion Pyrexia Torsade De Pointes Ventricular Fibrillation		Allopurinol	C	Glaxo Wellcome	ORAL

Date:04/12/01ISR Number: 3703312-XReport Type:Expedited (15-DaCompany Report #A0139738A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	2 YR	Blindness		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
		Optic Nerve Disorder		Serzone	C		

Date:04/12/01ISR Number: 3703322-2Report Type:Expedited (15-DaCompany Report #A0144832A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Brief Psychotic Disorder		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Without Marked Stressors Complex Partial Seizures Confusional State Hallucination, Visual Memory Impairment Persecutory Delusion		Alcohol	SS		

Date:04/12/01ISR Number: 3703326-XReport Type:Expedited (15-DaCompany Report #B0100605A  
 Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	22TAB Single	Aggression		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - dose		Convulsion					
Initial or Prolonged	14TAB Single	Drug Interaction		Dothiepin	SS		ORAL
		Intentional Misuse					
		Sedation					

Date:04/12/01ISR Number: 3703333-7Report Type:Expedited (15-DaCompany Report #B0103253A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 10 DAY		Anxiety		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged		Dry Throat Dysgeusia Eye Disorder					

Date:04/12/01ISR Number: 3703334-9Report Type:Expedited (15-DaCompany Report #B0103454A  
Age:51 YR Gender:Female I/FU:I

Outcome	PT
Other	Anaphylactic Shock Irritable Bowel Syndrome

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Proctitis Rash Pruritic				
Dose	Duration		Report Source	Product	Role	Manufacturer
150MG Twice per day	26 DAY			Zyban	PS	Glaxo Wellcome
				Premique Cycle	C	

Date:04/12/01ISR Number: 3703335-0Report Type:Expedited (15-DaCompany Report #B0103456A  
Age: Gender:Unknown I/FU:I

Outcome	PT		Report Source	Product	Role	Manufacturer
Dose	Duration					
Hospitalization - Initial or Prolonged		Serotonin Syndrome Tremor		Zyban Sertraline	PS C	Glaxo Wellcome

Date:04/12/01ISR Number: 3703336-2Report Type:Expedited (15-DaCompany Report #B0103457A  
Age:26 YR Gender:Female I/FU:F

Outcome	PT		Report Source	Product	Role	Manufacturer
Dose	Duration					
Hospitalization - 150MG Twice Initial or Prolonged per day	13 DAY	Hallucination Tachycardia		Zyban	PS	Glaxo Wellcome

Date:04/12/01ISR Number: 3703340-4Report Type:Expedited (15-DaCompany Report #B0103553A  
Age:45 YR Gender:Female I/FU:I

Outcome	PT		Report Source	Product	Role	Manufacturer
Dose	Duration					
Life-Threatening 150MG Per day		Cerebrovascular Accident		Zyban	PS	Glaxo Wellcome
Other 2.5MG per day		Feeling Abnormal		Bendrofluazide	C	Glaxo Wellcome
100MG per day		Hypertension		Atenolol	C	
				Ethynodiol	C	

Date:04/12/01ISR Number: 3703342-8Report Type:Expedited (15-DaCompany Report #B0103580A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Twice per day	18 DAY	Angina Pectoris Arrhythmia		Zyban  Irbesartan Alprazolam	PS  C C	Glaxo Wellcome	ORAL  ORAL ORAL

Date:04/12/01ISR Number: 3703343-XReport Type:Expedited (15-DaCompany Report #B0103725A  
Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Zyban	PS	Glaxo Wellcome	

Date:04/12/01ISR Number: 3703344-1Report Type:Expedited (15-DaCompany Report #B0103731A  
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day Initial or Prolonged	19 DAY	Diarrhoea Difficulty In Walking Joint Stiffness Oedema Peripheral Urticaria		Zyban	PS	Glaxo Wellcome	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/12/01ISR Number: 3703345-3Report Type:Expedited (15-DaCompany Report #B0103735A

Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Vaginal Haemorrhage		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day							

Date:04/12/01ISR Number: 3704094-8Report Type:Expedited (15-DaCompany Report #A0144419A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL							
		Neuropathy Peripheral	Professional Company Representative				

Date:04/12/01ISR Number: 3704295-9Report Type:Expedited (15-DaCompany Report #A0144245A

Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Choroidal Detachment	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL							
		Intraocular Pressure Increased	Professional	Losartan Potassium	C		
Malaise							

Date:04/12/01ISR Number: 3704308-4Report Type:Expedited (15-DaCompany Report #B0101861A

Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Depression	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
150 MG/TWICE							
		Dyspnoea	Professional				
PER DAY							

Eyelid Oedema  
Face Oedema  
Gastritis  
Hypersensitivity  
Insomnia  
Muscle Spasms  
Oral Intake Reduced  
Pharyngeal Oedema  
Pruritus  
Urticaria

Date:04/12/01ISR Number: 3704309-6Report Type:Expedited (15-DaCompany Report #B0102451A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability		Asthenia Disturbance In Attention	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
300 MG/TWICE PER DAY		Dry Mouth Overdose Paraesthesia Tremor Vision Blurred	Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/12/01ISR Number: 3704311-4Report Type:Expedited (15-DaCompany Report #B0103067A  
Age:60 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 Initial or Prolonged MG/PER/ORAL	Cardiac Failure	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:04/12/01ISR Number: 3704312-6Report Type:Expedited (15-DaCompany Report #B0102788A  
Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability 150 MG/TWICE Required PER DAY/ORAL Intervention to Prevent Permanent Impairment/Damage	Psoriasis	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:04/12/01ISR Number: 3704313-8Report Type:Expedited (15-DaCompany Report #B0100686A  
Age:55 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ ORAL Initial or Prolonged	Delirium Post Procedural Complication	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Ssri	C		
			Zaleplon	C		
			Diazepam	C		
			Sevoflurane	C		
			Propofol	C		
			Fentanyl	C		
			Rocuronium Bromide	C		
			Atropine	C		
			Bupivacaine	C		

Date:04/12/01ISR Number: 3704314-XReport Type:Expedited (15-DaCompany Report #B0102951A  
Age:45 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 150 MG/TWICE	Angina Pectoris	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Intervention to PER DAY/ORAL	Depression					
Prevent Permanent Impairment/Damage	Palpitations		Perindopril	C		
			Nicorandil	C		
			Lansoprazole	C		
			Isosorbide	C		
			Felodipine	C		
			Atorvastatn Calcium	C		
			Gaviscon	C		
			Aspirin	C		
			Beclomethasone			
			Dipropion.	C		
			Salbutamol Sulphate	C		

Date:04/12/01ISR Number: 3704317-5Report Type:Expedited (15-DaCompany Report #B0091784A  
Age:65 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Abnormal Dreams Bronchospasm Chronic Obstructive

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Airways Disease Exacerbated Drug Interaction Dyspnoea	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Emphysema	Health				
250 MG/FOUR TIMES PER DAY 10 DAY		Hallucination Insomnia Lower Respiratory Tract Infection Lymphadenopathy Productive Cough	Professional	Erythromycin (Formulation Unknown) (Erythromycin)	SS		

Date:04/12/01ISR Number: 3704321-7Report Type:Expedited (15-DaCompany Report #B0094748A  
Age:52 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150 MG ORAL			Erythema	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
					Ipratropium Bromide Fluticasone Propionate Norethi.+Oestradiol+ Oestr Dextroprop. + Paracetamol	C C C C		

Date:04/12/01ISR Number: 3714095-1Report Type:Periodic Company Report #2001SUS0121  
Age:40 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
PO 1 TAB BID PO			Dry Mouth Dyspepsia Dysphonia Pharyngitis	Consumer	Sustiva Wellbutrin (Bupropion Hcl) Prilosec	PS SS	Dupont Pharmaceuticals Co	ORAL ORAL

(Omeprazole)

C

Date:04/13/01ISR Number: 3704016-XReport Type:Expedited (15-DaCompany Report #B0103385A  
Age:69 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 2 WK Initial or Prolonged		Lower Respiratory Tract Infection	Zyban	PS	Glaxo Wellcome	ORAL

Date:04/13/01ISR Number: 3704380-1Report Type:Direct Company Report #  
Age:69 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1 PO BID		Confusional State Disorientation	Wellbutrin Sr (150mg)	PS		ORAL
1 PO BID		Hallucination	Detrol 2mg	SS		ORAL
		Paranoia	Sinemet	C		
			Metamucil	C		
			Prilosec	C		
			Asa	C		
			Plavix	C		
			Trazodone	C		
			Fibercon	C		
			Pericolace	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/13/01  
 Age: Gender:Male I/FU:I  
 Report Type:Direct

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1 TAB TWICE A Initial or Prolonged DAY ORAL	Arthralgia Oedema Pain In Extremity Pruritus Urticaria		Wellbutrin Darvocet Vioxx	PS C C		ORAL

Date:04/16/01  
 Age:33 YR Gender:Female I/FU:F  
 Report Type:Expedited (15-Da  
 Company Report #A0139749A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL  150 MG/ TWICE PER DAY/ ORAL	Abnormal Behaviour Astrocytoma Body Temperature Increased Brain Neoplasm Malignant Coma Convulsion Eye Movement Disorder Grand Mal Convulsion Loss Of Consciousness Medication Error Ocular Hyperaemia	Consumer	Bupropion Hydrochloride Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride) Clonazepam (Formulation Unknown) Fluvoxamine Maleate (Formulation Unknown) (Fluvoxamine Maleate) Triphasil Aspirin Biotin Ascorbic Acid Zinc Salt M.V.I.	PS SS SS C C C C C C	Glaxo Wellcome Inc	ORAL ORAL

Date:04/16/01ISR Number: 3705982-9Report Type:Expedited (15-DaCompany Report #A0144702A  
Age:33 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG TWICE PER DAY ORAL	Asthenia Dermatitis Dry Mouth Dyspnoea Erythema Feeling Jittery Lethargy Oedema Peripheral Throat Tightness	Consumer	Bupropion Hydrochloride  Ortho-Novum	PS  C	Glaxo Wellcome Inc	ORAL

Date:04/16/01ISR Number: 3705992-1Report Type:Expedited (15-DaCompany Report #PHBS2001US03455  
Age:26 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 800 MG/DAY 450 MG/DAY	Drug Interaction Drug Level Below Therapeutic	Literature Health Professional	Tegretol  Bupropion (Amfebutamone)  Lithium	PS  SS C	Novartis Pharmaceuticals Corp	

Freedom Of Information (FOI) Report

Trifluoperazine  
(Trifluoperazine) C

Date:04/16/01ISR Number: 3706039-3Report Type:Expedited (15-DaCompany Report #B0098194A  
Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG PER DAY ORAL	Acute Respiratory	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Distress Syndrome	Literature				
		Blood Pressure Decreased	Health	Chloroquine	C		
		Bronchopneumonia	Professional	Diphenhydramine	C		
		Conjunctival Hyperaemia					
		Deep Vein Thrombosis					
		Ecchymosis					
		Feeling Abnormal					
		Gastritis Erosive					
		Headache					
		Hepatic Congestion					
		Intracranial Pressure					
		Increased					
		Loss Of Consciousness					
		Mucous Membrane Disorder					
		Pleural Haemorrhage					
		Pulmonary Congestion					
		Purulence					
		Sudden Death					
		Swelling					
		Vision Blurred					

Date:04/16/01ISR Number: 3706057-5Report Type:Expedited (15-DaCompany Report #B0100605A  
Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Aggression	Foreign	Bupropion			
Hospitalization -		Convulsion	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
22 TABLET /							
Initial or Prolonged		Drug Interaction	Professional				
SINGLE DOSE /							

ORAL Intentional Misuse  
 Sedation Dothiepin Tablet (Dothiepin) SS ORAL  
 14 TABLET /  
 SINGLE DOSE /  
 ORAL

Date:04/16/01ISR Number: 3706069-1Report Type:Expedited (15-DaCompany Report #A0139738A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blindness Unilateral	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL	2 YR	Optic Neuritis	Health	Nefazodone			
		Photopsia	Professional	Hydrochloride	C		
		Scotoma					

Date:04/16/01ISR Number: 3706088-5Report Type:Expedited (15-DaCompany Report #D0013784A  
 Age:67 YR Gender:Male I/FU:F

Outcome  
 Death  
 Life-Threatening

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Freedom Of Information (FOI) Report

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required							
Dose	Duration						
Intervention to 150 MG PER		Arrhythmia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Prevent Permanent DAY ORAL		Bundle Branch Block Right					
Impairment/Damage		Cardiac Failure		Fosinopril Sodium	C		
		Cardiomegaly		Digitoxin	C		
		Pleural Effusion		Allopurinol	C		
		Torsade De Pointes					
		Ventricular Fibrillation					

Date:04/16/01ISR Number: 3706093-9Report Type:Expedited (15-DaCompany Report #B0103731A  
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG /PER		Diarrhoea	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
DAY/ORAL		Joint Stiffness					
		Oedema					
		Urticaria					

Date:04/16/01ISR Number: 3706095-2Report Type:Expedited (15-DaCompany Report #B0103735A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Vaginal Haemorrhage	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Intervention to 150 MG/PER							
Prevent Permanent DAY/ORA							
Impairment/Damage							

Date:04/16/01ISR Number: 3706099-XReport Type:Expedited (15-DaCompany Report #B0103253A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG	Anxiety Dry Throat	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
UNKNOWN		Dysgeusia Feeling Abnormal Photophobia					

Date:04/16/01ISR Number: 3706100-3Report Type:Expedited (15-DaCompany Report #B0103456A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG	Serotonin Syndrome	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
UNKNOWN			Professional Company Representative	Sertraline	C		

Date:04/16/01ISR Number: 3706101-5Report Type:Expedited (15-DaCompany Report #B0103209A  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG /TWICE PER DAY/ORAL	Dizziness Influenza Like Illness Myalgia	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Mebeverine C

Date:04/16/01ISR Number: 3706102-7Report Type:Expedited (15-DaCompany Report #B0103453A  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required 150 MG /TWICE Intervention to PER DAY/ORAL Prevent Permanent Impairment/Damage		Papilloedema Retinopathy	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Premique	C		

Date:04/16/01ISR Number: 3706103-9Report Type:Expedited (15-DaCompany Report #B0103470A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG Prevent Permanent /SINGLE Impairment/Damage DOSE/ORAL		Gait Disturbance Hemiplegia Nausea Palpitations	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:04/16/01ISR Number: 3706104-0Report Type:Expedited (15-DaCompany Report #B0103473A  
 Age:26 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG /TWICE PER DAY/ORAL		Oedema Peripheral Purpura	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:04/16/01ISR Number: 3706105-2Report Type:Expedited (15-DaCompany Report #B0103477A  
Age:45 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG/ORAL	Haemorrhagic Stroke Hemiparesis Nystagmus	Foreign	Bupropion Hydrochloride Gaviscon Omeprazole	PS C C	Glaxo Wellcome Inc	ORAL

Date:04/16/01ISR Number: 3706106-4Report Type:Expedited (15-DaCompany Report #B0103481A  
Age:62 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required Intervention to 150 MG /TWICE Prevent Permanent PER DAY/ORAL Impairment/Damage	Angina Pectoris		Bupropion Hydrochloride Atenolol Aspirin Nitroglycerin Ranitidine Hydrochloride	PS C C C C	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/16/01ISR Number: 3706107-6Report Type:Expedited (15-DaCompany Report #B0103576A  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Foreign	Bupropion			
		Atrial Fibrillation	Consumer	Hydrochloride	PS	Glaxo Wellcome Inc	
UNKNOWN	150 MG	Insomnia					

Date:04/16/01ISR Number: 3706109-XReport Type:Expedited (15-DaCompany Report #B0103584A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Haemorrhagic Stroke	Foreign	Bupropion			
Initial or Prolonged		Hemiparesis		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
300 MG /PER							
DAY/ ORAL	2 WK			Carvedilol	C		

Date:04/16/01ISR Number: 3706119-2Report Type:Expedited (15-DaCompany Report #B0103201A  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Lichen Planus	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER							
DAY ORAL							

Date:04/16/01ISR Number: 3706121-0Report Type:Expedited (15-DaCompany Report #B0103161A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Psoriasis	Foreign	Bupropion			
Initial or Prolonged		Rash Generalised	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							

PER DAY ORAL

Professional

Doxepin	C
Normin	C
Acitretin	C
Trandolapril	C
Panadeine Forte	C

Date:04/16/01ISR Number: 3706122-2Report Type:Expedited (15-DaCompany Report #A0144603A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Asthma	Health				
PER DAY ORAL		Bronchospasm	Professional				
		Burning Sensation					
		Chest Pain					
		Dermatitis					
		Dyspnoea					
		Face Oedema					
		Insomnia					
		Joint Stiffness					
		Oedema Peripheral					
		Pruritus					
		Urticaria					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/16/01ISR Number: 3706239-2Report Type:Expedited (15-DaCompany Report #B0103725A  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG UNK	Death	Foreign	Zyban	PS	Glaxo Wellcome Inc	
			Health Professional				

Date:04/16/01ISR Number: 3706241-0Report Type:Expedited (15-DaCompany Report #B0103553A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG PER	Feeling Abnormal	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Required	DAY ORAL	Hypertension					
Intervention to Prevent Permanent Impairment/Damage				Bendrofluazide	C		
				Atenolol	C		
				Ethynodiol	C		

Date:04/16/01ISR Number: 3706242-2Report Type:Expedited (15-DaCompany Report #B0103454A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required	150 MG TWICE	Proctitis	Foreign	Bupropion			
Intervention to Prevent Permanent Impairment/Damage	PER DAY ORAL	Pruritus	Health Professional	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Urticaria	Professional				
			Other	Premique Cycle	C		

Date:04/16/01ISR Number: 3706243-4Report Type:Expedited (15-DaCompany Report #B0103580A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Life-Threatening	Angina Pectoris	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE						
	Arrhythmia					
PER DAY ORAL						
			Irbesartan	C		
			Alprazolam	C		

Date:04/16/01ISR Number: 3706262-8Report Type:Expedited (15-DaCompany Report #A0144832A  
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Brief Psychotic Disorder	Foreign	Bupropion			
Initial or Prolonged	Without Marked Stressors	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
SINGLE						
DOSE/ORAL	Cognitive Disorder	Professional				
	Complex Partial Seizures		Ethanol (Alcohol)	SS		
	Confusional State					
	Hallucination, Visual					
	Memory Impairment					
	Persecutory Delusion					

Date:04/16/01ISR Number: 3706263-XReport Type:Expedited (15-DaCompany Report #B0103457A  
 Age:26 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Asthenia	Foreign	Bupropion			
Initial or Prolonged	Drug Toxicity	Consumer	Hydrochloride	PS	Glaxo Wellcome Inc	
150 MG/TWICE						
PER DAY	Hallucination					
	Tachycardia					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/16/01ISR Number: 3709305-0Report Type:Expedited (15-DaCompany Report #B0103210A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Arthralgia	Foreign	Bupropion			
Intervention to		Feeling Hot		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
300 MG /PER							
Prevent Permanent		Joint Swelling					
DAY/ORAL							
Impairment/Damage		Serum Sickness		Co-Dydramol	C		
		Urticaria		Omeprazole	C		

Date:04/16/01ISR Number: 3710210-4Report Type:Expedited (15-DaCompany Report #B0103211A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Antinuclear Antibody	Foreign	Bupropion			
Intervention to		Positive		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
Prevent Permanent		Arthralgia					
TWICE PER DAY							
Impairment/Damage		Autoimmune Disorder					
/ ORAL							
		Mouth Ulceration					
		Rash Pruritic					

Date:04/17/01ISR Number: 3705492-9Report Type:Expedited (15-DaCompany Report #A0144855A  
Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Papilloedema		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
2TAB Per day							
		Photosensitivity Reaction		Pepcid	C		
		Retinal Disorder		Nasalide	C		
		Vision Blurred		Covera	C		
		Visual Field Defect		Calan	C		
		Vitreous Floaters		Tylenol With Codeine	C		
				Estrace	C		
				Librium	C		
				Benadryl	C		

Date:04/17/01ISR Number: 3707000-5Report Type:Expedited (15-DaCompany Report #B0103385A

Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL	2 WK	Dizziness	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Lower Respiratory Tract Infection Nausea	Health Professional				

Date:04/17/01ISR Number: 3707225-9Report Type:Expedited (15-DaCompany Report #S01-USA-00683-01

Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 60 MG QD PO		Complications Of Maternal Exposure To Therapeutic	Health Professional	Celexa	PS	Forest Laboratories Inc	ORAL
200 MG BID PO		Drugs Intra-Uterine Death Stillbirth		Wellbutrin - Slow Release (Bupropion Hydrochloride)	SS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/18/01ISR Number: 3706392-0Report Type:Expedited (15-DaCompany Report #A0145045A  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	400MG Per day	Complications Of Maternal		Wellbutrin	PS	Glaxo Wellcome	ORAL
UNKNOWN	60MG Per day	Exposure To Therapeutic Drugs Stillbirth		Celexa	SS		

Date:04/19/01ISR Number: 3707389-7Report Type:Expedited (15-DaCompany Report #B0097263A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Amnesia Constipation Depression Diarrhoea Disturbance In Attention Dysphonia Headache Insomnia Joint Stiffness Lethargy Mouth Haemorrhage Mouth Ulceration Pain Palpitations Panic Attack Personality Change Psychiatric Symptom Rectal Tenesmus Sedation Social Avoidant Behaviour Tongue Oedema Vision Blurred Weight Increased		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/19/01ISR Number: 3707406-4Report Type:Expedited (15-DaCompany Report #B0103442A  
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day							
5MG Per day				Nitrazepam	C		ORAL

Date:04/19/01ISR Number: 3713272-3Report Type:Periodic Company Report #US007954  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased	Consumer	Provigil	PS	Cephalon Inc	ORAL
200 MG QAM		Heart Rate Increased					
ORAL				Provigil	SS		ORAL
100 MG QD							
ORAL				Zyban	SS		
150 MG BID				Zyban	SS		
				Avonex "Biogen"	C		
				Micronor	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/20/01ISR Number: 3708056-6Report Type:Expedited (15-DaCompany Report #A0131125A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly 150MG Twice per day	5 WK	Aortic Valve Disease Aortic Valve Incompetence		Bupropion	PS	Glaxo Wellcome	ORAL
UNKNOWN		Caesarean Section Cardiac Valve Disease Complications Of Maternal Exposure To Therapeutic Drugs Neonatal Disorder Twin Pregnancy		Multivitamins Septra	C C	Glaxo Wellcome	

Date:04/20/01ISR Number: 3708065-7Report Type:Expedited (15-DaCompany Report #A0145057A

Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death UNKNOWN		Coma Completed Suicide Epistaxis Face Oedema Mouth Haemorrhage Pulse Absent		Wellbutrin Unisom	PS SS	Glaxo Wellcome	ORAL

Date:04/20/01ISR Number: 3708069-4Report Type:Expedited (15-DaCompany Report #B0091583A

Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Per day		Circulatory Collapse Respiratory Disorder Sudden Death		Bupropion Hydrochloride Alcohol	PS C	Glaxo Wellcome	ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	16 DAY	Bronchopneumonia	Zyban	PS	Glaxo Wellcome	ORAL
			Convulsion	Medroxy Progesterone			
			Drug Hypersensitivity	Acetate	C		
			Ecchymosis	Chloroquine	C		
			Epistaxis	Diphenhydramine	C		
			Gastritis Erosive				
			Head Injury				
			Hearing Impaired				
			Hepatic Congestion				
			Hepatitis				
			Intracranial Pressure				
			Increased				
			Pleural Haemorrhage				
			Pulmonary Congestion				
			Pulse Absent				
			Respiratory Arrest				
			Syncope				
			Vision Blurred				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/20/01ISR Number: 3708072-4Report Type:Expedited (15-DaCompany Report #B0100488A  
 Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Single Initial or Prolonged dose	1 DAY	Anaphylactic Reaction		Zyban	PS	Glaxo Wellcome	
1 DAY		Dermatitis		Tinidazole	SS		
		Drug Interaction					
		Myocardial Infarction Pyrexia Swelling					

Date:04/20/01ISR Number: 3708079-7Report Type:Expedited (15-DaCompany Report #B0104098A  
 Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Depression Hallucination		Zyban	PS	Glaxo Wellcome	

Date:04/20/01ISR Number: 3708692-7Report Type:Expedited (15-DaCompany Report #A0144855A  
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2 TABLET / PER DAY / ORAL		Papilloedema Photosensitivity Reaction Retinal Disorder	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
		Vision Blurred		Famotidine	C		
		Visual Field Defect		Flunisolide	C		
		Vitreous Floaters		Verapamil Hydrochloride	C		
				Tylenol With Codeine	C		
				Oestradiol	C		
				Librium	C		
				Diphenhydramine Hcl	C		

Date:04/23/01ISR Number: 3708623-XReport Type:Expedited (15-DaCompany Report #A0145297A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erythema Nodosum		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Oedema Peripheral					
per day							

Date:04/23/01ISR Number: 3708624-1Report Type:Expedited (15-DaCompany Report #A0145351A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Bupropion	PS	Glaxo Wellcome	ORAL
150MG Per day		Pregnancy					

Date:04/23/01ISR Number: 3708627-7Report Type:Expedited (15-DaCompany Report #B0102453A  
Age:37 YR Gender:Male I/FU:F

Outcome	PT
Death	Convulsion
	Hypoglycaemia
	Mouth Haemorrhage
	Respiratory Arrest
	Salivary Hypersecretion

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sudden Death

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	6 WK		Zyban	PS	Glaxo Wellcome	

Date:04/23/01ISR Number: 3709034-3Report Type:Direct  
 Age: Gender:Male I/FU:I Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage		Hypertension Medication Error Nervousness Paranoia Sleep Disorder		Covera 180mg Wellbutrin 100mg	PS SS		

Date:04/23/01ISR Number: 3709035-5Report Type:Direct  
 Age:62 YR Gender:Male I/FU:I Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage	1 BID ORAL	Arthralgia Fatigue Gastrooesophageal Reflux Disease Head Injury		Wellbutrin 75mg Glaxo Quinapril Naicin	PS C C	Glaxo	ORAL

Date:04/23/01ISR Number: 3709107-5Report Type:Expedited (15-DaCompany Report #2001009504-1  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Anxiety	Consumer	Paxil	PS	Smithkline Beecham	

Initial or Prolonged	Condition Aggravated	Pharmaceuticals	ORAL
20 MILLIGRAMS			
	Lethargy		
1.0 DAILY			
ORAL			
		Wellbutrin Sr	
		(Bupropion)	SS
100			ORAL
MILLIGRAMS			
1.0 DAILY			
ORAL	9 DAY		
		Wellbutrin Sr	
		(Bupropion)	SS
200			ORAL
MILLIGRAMS			
ORAL			

Date:04/23/01ISR Number: 3709146-4Report Type:Expedited (15-DaCompany Report #A0145045A  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Arrest	Study	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
400 MG/PER		Complications Of Maternal	Health				
DAY/ORAL		Exposure To Therapeutic	Professional	Citalopram			
		Drugs		Hydrobromide			
		Foetal Disorder		(Citalopram			
		Stillbirth		Hydrobromide)	SS		
60 MG/PER DAY							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/23/01  
 Age:58 YR  
 Gender:Male  
 I/FU:I

Report Type:Periodic  
 Company Report #PHEH2000US03164

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hallucination	Health Professional	Restoril	PS	Novartis Pharmaceuticals Corp	
				Bupropion (Amfebutamone)	SS		
				Alprazolam "Nm Pharma" (Alprazolam)	SS		
				Zolpidem Tartrate (Zolpidem Tartrate)	SS		

Date:04/24/01  
 Age:  
 Gender:Unknown  
 I/FU:F

Report Type:Expedited (15-Da  
 Company Report #B0093206A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/24/01  
 Age:  
 Gender:Unknown  
 I/FU:I

Report Type:Expedited (15-Da  
 Company Report #B0104336A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Zyban	PS	Glaxo Wellcome	

Date:04/24/01  
 Age:44 YR  
 Gender:Male  
 I/FU:F

Report Type:Expedited (15-Da  
 Company Report #B0104339A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Erythema Multiforme  Infection In An  Immunocompromised Host Stevens-Johnson Syndrome		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/24/01ISR Number: 3709078-1Report Type:Expedited (15-DaCompany Report #B0104342A  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day							
				Alcohol	C		
				Triazolam	C		
1TAB Per day							

Date:04/24/01ISR Number: 3710170-6Report Type:Expedited (15-DaCompany Report #A0131125A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Congenital Anomaly		Aortic Valve Disease	Study	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ORAL		Aortic Valve Incompetence	Health				
		Caesarean Section	Professional	Multivitamin	C		
		Complications Of Maternal		Co-Trimoxazole	C		
		Exposure To Therapeutic					
		Drugs					
		Twin Pregnancy					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/24/01ISR Number: 3710197-4Report Type:Expedited (15-DaCompany Report #A0145057A

Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coma	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
UNK / UNK /		Completed Suicide					
ORAL		Cyanosis		Doxylamine Succinate			
		Epistaxis		(Formulation			
		Face Oedema		Unknown) (Doxylamine			
		Mouth Haemorrhage		Succinate)	SS		
		Pulse Absent					

Date:04/24/01ISR Number: 3710613-8Report Type:Expedited (15-DaCompany Report #B0091583A

Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Circulatory Collapse	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Sudden Death	Literature				
DAY / ORAL				Ethanol	C		

Date:04/24/01ISR Number: 3710614-XReport Type:Expedited (15-DaCompany Report #B0098194A

Age:21 YR Gender:Female I/FU:F

Outcome	PT
Death	Bronchopneumonia
	Convulsion
	Cyanosis
	Dizziness
	Ecchymosis
	Encephalitis
	Epistaxis
	Fall
	Fatigue
	Feeling Abnormal
	Gastritis Erosive
	Head Injury
	Headache

Hearing Impaired  
Hepatic Congestion  
Hepatitis  
Intracranial Pressure  
Increased  
Livedo Reticularis  
Loss Of Consciousness  
Lymphadenopathy  
Meningitis  
Nasal Congestion  
Ocular Hyperaemia  
Pallor  
Pleural Haemorrhage  
Pulmonary Congestion  
Pulmonary Oedema  
Pulse Absent  
Respiratory Arrest  
Respiratory Distress  
Rhinitis  
Sedation  
Sudden Death  
Syncope  
Trismus

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Vision Blurred Vomiting				
Dose	Duration		Report Source	Product	Role	Manufacturer
150 MG / PER DAY / ORAL			Foreign Literature	Zyban	PS	Glaxo Wellcome Inc
			Health Professional	Medroxyprogesterobne Ace. Chloroquine Diphenhydramine	C C C	Route ORAL

Date:04/24/01ISR Number: 3710778-8Report Type:Expedited (15-DaCompany Report #B0104098A  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150 MG		Depression	Foreign	Zyban	PS	Glaxo Wellcome Inc	
		Hallucination Nervousness	Consumer				

Date:04/24/01ISR Number: 3710780-6Report Type:Expedited (15-DaCompany Report #B0100488A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / Initial or Prolonged SINGLE DOSE /		Anaphylactic Reaction	Foreign	Zyban	PS	Glaxo Wellcome Inc	
		Chest Pain	Health				
		Dermatitis Myocardial Infarction Pyrexia Swelling	Professional	Tinidazole (Formulation Unknown) (Tinidazole)	SS		
SINGLE DOSE							

Date:04/24/01ISR Number: 3710787-9Report Type:Expedited (15-DaCompany Report #A107917  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 100.00 MG		Anger Depression	Foreign Health	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
Prevent Permanent TOTAL:DAILY:O Impairment/Damage RAL		Drug Interaction Irritability	Professional Other				
300.00 MG		Suicidal Ideation Thinking Abnormal		Bupropion	SS		ORAL
TOTAL: BID:ORA							
L				Atorvastatin	C		

Date: 04/24/01  
 ISR Number: 3710830-7  
 Report Type: Expedited (15-DaCompany Report #B0097263A)  
 Age: 39 YR  
 Gender: Female  
 I/FU: F

Outcome	PT
Life-Threatening	Amnesia Constipation Depressed Mood Diarrhoea Disturbance In Attention Dysphonia Headache Insomnia Joint Stiffness Lethargy Mouth Haemorrhage

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Mouth Ulceration Pain Palpitations	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Panic Attack	Health Professional				
		Personality Change Psychiatric Symptom Rectal Tenesmus Sedation Social Avoidant Behaviour Tongue Oedema Vision Blurred Weight Increased					

Date:04/25/01ISR Number: 3710049-XReport Type:Expedited (15-DaCompany Report #A0143021A  
Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Confusional State	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
300MG per day Hospitalization - 100MG Three Initial or Prolonged times per day		Drug Toxicity	Professional	Amantadine	SS		ORAL
	YR	Extrapyramidal Disorder					
		Fall		Prozac	C		
		Hallucination, Auditory		Loxapine	C		
		Insomnia		Seroquel	C		
		Limb Injury		Risperdal	C		
		Memory Impairment					
		Paranoia					
		Schizophrenia					
		Tremor					

Date:04/25/01ISR Number: 3710054-3Report Type:Expedited (15-DaCompany Report #A0145483A  
Age:10 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Growth Retardation		Wellbutrin	PS	Glaxo Wellcome	ORAL
400MG per day	2 YR	Hand Fracture		No Concurrent			

Insomnia  
Irritability

Medications

C

Date:04/25/01ISR Number: 3710056-7Report Type:Expedited (15-DaCompany Report #A0145632A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Surgery Tooth Disorder		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/25/01ISR Number: 3710063-4Report Type:Expedited (15-DaCompany Report #B0103456A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Drug Interaction Serotonin Syndrome	Foreign Health	Zyban	PS	Glaxo Wellcome	ORAL
75MG Per day			Professional	Effexor Xr	SS		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/25/01ISR Number: 3710072-5Report Type:Expedited (15-DaCompany Report #B0104340A  
Age:51 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300MG Per day	Cholelithiasis		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged 2U Per day	Pancreatitis Acute		Combivir	C	Glaxo Wellcome	
400MG Per day			Nevirapine	C		
1TAB Per day			Co-Trimoxazole	C	Glaxo Wellcome	
3U Per day			Carbasalate	C		
			Alcohol	C		

Date:04/25/01ISR Number: 3710076-2Report Type:Expedited (15-DaCompany Report #B0104482A  
Age:59 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	Alopecia Erythema		Zyban	PS	Glaxo Wellcome	ORAL
	Flushing		Cyanocobalamin	C	Glaxo Wellcome	
	Oedema Peripheral		Kliogest	C		
	Pruritus					
	Psoriasis					
	Rash Erythematous					
	Rash Generalised					
	Rash Pruritic					
	Skin Exfoliation					

Date:04/25/01ISR Number: 3710077-4Report Type:Expedited (15-DaCompany Report #B0104484A  
Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 300MG Per day	Cardiac Arrest		Zyban	PS	Glaxo Wellcome	

200MG per day	Shock	Amiodarone	C	
50MG per day	Ventricular Fibrillation	Captopril	C	Glaxo Wellcome
250MG per day	Ventricular Tachycardia	Warfarin	C	Glaxo Wellcome
		Lanoxin	C	Glaxo Wellcome
		Influenza Vaccination	C	

INTRAVENOUS

Date:04/25/01ISR Number: 3710078-6Report Type:Expedited (15-DaCompany Report #B0104493A  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 4 DAY Initial or Prolonged RESPIRATORY		Asthma		Zyban	PS	Glaxo Wellcome	ORAL
(INHALATION)		Bronchospasm		Salbutamol	C	Glaxo Wellcome	
RESPIRATORY				Beclomethasone	C	Glaxo Wellcome	
(INHALATION)							

Date:04/25/01ISR Number: 3710079-8Report Type:Expedited (15-DaCompany Report #B0104495A  
Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 300MG Per day 22 DAY		Dermatitis Bullous		Zyban	PS	Glaxo Wellcome	ORAL
		Myocardial Infarction		Cefaclor	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/25/01ISR Number: 3710080-4Report Type:Expedited (15-DaCompany Report #B0104497A  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Hospitalization -		Blood Pressure Increased					
per day	36 DAY						
Initial or Prolonged		Chest Pain		Bendrofluazide	C	Glaxo Wellcome	ORAL
2.5MG per day	54 MON						
Disability							

Date:04/25/01ISR Number: 3711364-6Report Type:Expedited (15-DaCompany Report #B0103442A  
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Dermatitis	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL		Sudden Death					
				Nitrazepam	C		

Date:04/26/01ISR Number: 3710947-7Report Type:Expedited (15-DaCompany Report #A0133478A  
Age:17 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Coma		Wellbutrin	PS	Glaxo Wellcome	ORAL
38TAB Single							
Hospitalization -		Dialysis					
dose							
Initial or Prolonged		Drug Level Above		Tegretol	SS		ORAL
54TAB Single							
Other		Therapeutic					
dose							
12TAB Single		Ileus Paralytic		Trazodone	SS		ORAL
dose		Overdose					
38TAB Single		Oxygen Saturation		Seroquel	SS		ORAL

dose  
 1600MG Single  
 dose

Decreased  
 Sedation  
 Advil  
 SS  
 ORAL

Date:04/26/01ISR Number: 3710953-2Report Type:Expedited (15-DaCompany Report #B0102510A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased Convulsion Dizziness Dry Mouth Dysgeusia Fatigue Weight Decreased		Zyban Celecoxib	PS C	Glaxo Wellcome	

Date:04/26/01ISR Number: 3710957-XReport Type:Expedited (15-DaCompany Report #B0104518A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - UNKNOWN		Angina Pectoris		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged		Dizziness		Atenolol	C		
25MG Per day		Lower Respiratory Tract Infection		Aspirin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/26/01ISR Number: 3711532-3Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Feeling Abnormal		Wellbutrn-Sr 100 Mg	PS		
100 MG				Zovirax 200 Mg	SS		ORAL
200 MG Q4H							
ORAL							

Date:04/26/01ISR Number: 3711895-9Report Type:Expedited (15-DaCompany Report #A0145351A  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Study	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Complications Of Maternal	Health				
DAY / ORAL		Exposure To Therapeutic	Professional				
		Drugs					
		Pregnancy					

Date:04/26/01ISR Number: 3712047-9Report Type:Expedited (15-DaCompany Report #A0145297A  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erythema Nodosum	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Joint Swelling	Consumer				
TWICE PER DAY							
/ ORAL							

Date:04/26/01ISR Number: 3712092-3Report Type:Expedited (15-DaCompany Report #B0102453A  
 Age:37 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG/TWICE		Convulsion	Foreign	Zyban	PS	Glaxo Wellcome Inc	
	PER DAY	6 WK	Diabetes Mellitus	Health				
			Hypoglycaemia	Professional				
			Respiratory Arrest					
			Sudden Death					

Date:04/27/01ISR Number: 3711700-0Report Type:Expedited (15-DaCompany Report #A0138130A  
Age:56 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Twice		Agitation		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	per day	4 DAY	Anger					
	100MG At		Angina Pectoris		Nefazodone	C		ORAL
	night		Blood Pressure Increased					
	1TAB At night		Cyanosis		Risperidone	C		ORAL
			Speech Disorder					

Date:04/27/01ISR Number: 3711702-4Report Type:Expedited (15-DaCompany Report #A0142161A  
Age:54 YR Gender:Female I/FU:F

Outcome	PT
Life-Threatening	Activated Partial Thromboplastin Time Shortened Blood Potassium Decreased Culture Urine Positive Decreased Activity

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Ecchymosis Epistaxis Gram Stain Negative Haematocrit Decreased Haemoglobin Decreased	Zyban	PS	Glaxo Wellcome	ORAL
.5MG Twice per day		Idiopathic Thrombocytopenic Purpura	Rivotril	C		ORAL
		Lymphocyte Count Increased Monocyte Count Increased Musculoskeletal Stiffness Neutrophil Count Increased Purpura Rectal Haemorrhage Red Blood Cell Count Decreased Specific Gravity Urine Abnormal White Blood Cell Count Increased	Asa	C		

Date:04/27/01ISR Number: 3711707-3Report Type:Expedited (15-DaCompany Report #A0144646A  
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Twice per day	6 DAY	Dermatitis		Zyban	PS	Glaxo Wellcome	ORAL
25MG Per day		Dizziness Hyperhidrosis		Hydrochlorothiazide	C		ORAL
10MG Per day		Middle Insomnia		Zestril	C		ORAL
25MG Per day		Nausea		Vioxx	C		ORAL
.625MG Per		Tremor		Premarin	C		ORAL

day

Date:04/27/01ISR Number: 3711709-7Report Type:Expedited (15-DaCompany Report #A0145297A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erythema Nodosum		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	88	DAY					
YR				Synthroid	C	Glaxo Wellcome	ORAL
				Pravachol	C		ORAL

Date:04/27/01ISR Number: 3713522-3Report Type:Expedited (15-DaCompany Report #A0145483A  
Age:10 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Growth Retardation	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
SEE TEXT							
/ORAL		Hand Fracture	Professional				
		Insomnia					
		Irritability					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/27/01ISR Number: 3713528-4Report Type:Expedited (15-DaCompany Report #A0143021A  
 Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Condition Aggravated	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL							
Hospitalization - Initial or Prolonged		Confusional State Drug Toxicity Extrapyramidal Disorder	Professional	Amantadine Tablet (Amantadine Hydrochloride)	SS		ORAL
SEE TEXT/ORAL							
		Fall		Fluoxetine			
		Hallucination, Auditory		Hydrochloride	C		
		Insomnia		Loxapine	C		
		Limb Injury		Quetiapine Fumarate	C		
		Memory Impairment		Risperidone	C		
		Paranoia					
		Schizophrenia					
		Tremor					

Date:04/27/01ISR Number: 3713600-9Report Type:Expedited (15-DaCompany Report #B0103456A  
 Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Agitation Drug Interaction	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL							
		Dyskinesia	Professional				
		Flushing	Company Representative	Velafaxine Hydrochloride (Venlafaxine Hydrochloride)	SS		ORAL
75 MG (PER DAY) ORAL		Heart Rate Increased Movement Disorder Serotonin Syndrome					
		Tremor					

Date:04/27/01ISR Number: 3713603-4Report Type:Expedited (15-DaCompany Report #B0093206A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death	Completed Suicide	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL		Professional Company Representative				

Date:04/27/01ISR Number: 3713608-3Report Type:Expedited (15-DaCompany Report #B0104336A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
150 MG			Professional				

Date:04/27/01ISR Number: 3713609-5Report Type:Expedited (15-DaCompany Report #B0104339A  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Erythema Multiforme Immunosuppression	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL		Stevens-Johnson Syndrome	Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/27/01ISR Number: 3713610-1Report Type:Expedited (15-DaCompany Report #B0104342A  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Circulatory Collapse	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE		Convulsion					
PER DAY) ORAL		Nasopharyngitis	Professional				
				Ethanol	C		
				Triazolam	C		

Date:04/27/01ISR Number: 3713612-5Report Type:Expedited (15-DaCompany Report #B0104482A  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Alopecia	Foreign Health	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS	Glaxo Wellcome Inc	ORAL
Intervention to Prevent Permanent		Oedema Peripheral Pruritus	Professional				
150 MG, TWICE							
Impairment/Damage		Psoriasis					
PER DAY, ORAL		Rash Generalised		Cyanocobalamin Kliogest	C		
					C		

Date:04/27/01ISR Number: 3713614-9Report Type:Expedited (15-DaCompany Report #B0104495A  
Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Blister Dermatitis	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS	Glaxo Wellcome Inc	ORAL
300 MG, PER		Lower Respiratory Tract					
DAY, ORAL		Infection					
		Myocardial Infarction		Cefaclor	C		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Tooth Repair	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS	Glaxo Wellcome Inc	ORAL
ORAL							

Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300 MG/PER	Abdominal Pain Blood Pressure Increased Cholelithiasis	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
DAY		Pallor Pancreatitis Acute Vomiting		Combivir Nevirapine Co-Trimoxazole Carbaspirin Calcium Ethanol	C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/27/01ISR Number: 3713672-1Report Type:Expedited (15-DaCompany Report #B0104497A  
Age:57 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 150 MG/TWICE Initial or Prolonged PER DAY/ORAL Disability	Angina Pectoris Blood Pressure Increased Chest Pain	Foreign	Bupropion Hydrochloride  Bendrofluazide	PS  C	Glaxo Wellcome Inc	ORAL

Date:04/27/01ISR Number: 3713673-3Report Type:Expedited (15-DaCompany Report #B0104493A  
Age:46 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG/PER DAY/ORAL	Asthma	Foreign	Bupropion Hydrochloride  Salbutamol Sulphate Beclomethasone Dipropion	PS  C C	Glaxo Wellcome Inc	ORAL

Date:04/27/01ISR Number: 3713674-5Report Type:Expedited (15-DaCompany Report #B0104484A  
Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death 300 MG/PER DAY	Cardiac Arrest Shock Ventricular Fibrillation Ventricular Tachycardia	Foreign	Bupropion Hydrochloride  Amiodarone Captopril Warfarin Sodium Digoxin Influenza Vaccine	PS  C C C C C	Glaxo Wellcome Inc	

Date:04/30/01ISR Number: 3713068-2Report Type:Expedited (15-DaCompany Report #B0100732A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Anxiety Palpitations Sleep Disorder Due To General Medical Condition, Insomnia Type Tremor Vein Pain		Zyban	PS	Glaxo Wellcome	

Date:04/30/01ISR Number: 3713865-3Report Type:Expedited (15-DaCompany Report #A0133478A  
Age:17 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 38 TABLET, Hospitalization - SINGLE DOSE, Initial or Prolonged ORAL Required Intervention to 54 Prevent Permanent TABLET/SINGLE Impairment/Damage DOSE/ORAL 12 TABLET/SINGLE		Dialysis  Overdose	Health  Professional	Wellbutrin   Carbamazepine Tablet (Carbamazepine)  Trazodone Tablet	PS   SS  SS	Glaxo Wellcome Inc	ORAL   ORAL  ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

DOSE/ORAL

38

Quetiapine Fumarate  
Tablet (Quetiapine  
Fumarate)

SS

ORAL

TABLET/SINGLE

DOSE/ORAL

1600

Ibuprofen Tablet  
(Ibuprofen)

SS

ORAL

MG/SINGLE

DOSE/ORAL

Date:04/30/01ISR Number: 3715077-6Report Type:Expedited (15-DaCompany Report #B0104518A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Angina Pectoris Asthenia Chest Pain Cough Dizziness Nausea Neck Pain	Foreign Consumer	Zyban Atenolol Aspirin	PS C C	Glaxo Wellcome Inc	

Date:04/30/01ISR Number: 3715080-6Report Type:Expedited (15-DaCompany Report #B0102510A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG		Blood Pressure Increased Convulsion Dizziness Dry Mouth Dysgeusia Fatigue Weight Decreased	Foreign Health Professional	Zyban Celecoxib	PS C	Glaxo Wellcome Inc	

Date:05/01/01ISR Number: 3713767-2Report Type:Expedited (15-DaCompany Report #A0145759A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
100MG Per day	2	YR	Blood Pressure Increased	Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
			Lethargy				
			Myocardial Ischaemia				
			Sedation				
			Ventricular Hypertrophy				

Date:05/01/01ISR Number: 3713772-6Report Type:Expedited (15-DaCompany Report #B0104997A  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability							
300MG Per day			Diabetes Mellitus	Zyban	PS	Glaxo Wellcome	ORAL
Other							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/01/01ISR Number: 3713773-8Report Type:Expedited (15-DaCompany Report #B0105005A  
Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
150MG							
Hospitalization -		Soft Tissue Injury					
Variable dose	8 DAY						
Initial or Prolonged		Urinary Tract Infection		Dianette	C		ORAL
Other				Salbutamol	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)							

Date:05/01/01ISR Number: 3713775-1Report Type:Expedited (15-DaCompany Report #B0105091A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Inflammation		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	17 DAY						
RESPIRATORY		Swelling		Beclomethasone	C	Glaxo Wellcome	
(INHALATION)	200MCG Twice						
per day							
RESPIRATORY				Salbutamol	C	Glaxo Wellcome	
(INHALATION)							

Date:05/01/01ISR Number: 3713776-3Report Type:Expedited (15-DaCompany Report #B0105093A  
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Atrial Fibrillation		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	31 DAY	Gastroenteritis					

200MG Twice	Helicobacter	Dipyridamole	C	ORAL
per day				
75MG per day		Aspirin	C	ORAL
10MG per day		Amlodipine	C	ORAL
		Omeprazole	C	ORAL

Date:05/01/01ISR Number: 3714642-XReport Type:Expedited (15-DaCompany Report #A0138130A  
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/ TWICE		Agitation Anger	Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY /		Angina Pectoris					
ORAL	4 DAY	Blood Pressure Increased					
		Cyanosis Speech Disorder		Nefazodone Hydrochloride Risperidone	C C		

Date:05/01/01ISR Number: 3715176-9Report Type:Expedited (15-DaCompany Report #A0145297A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG (TWICE)		Erythema Nodosum Pain In Extremity	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY) ORAL				Thyroxine Sodium Pravastatin Sodium	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/01/01ISR Number: 3715194-0Report Type:Expedited (15-DaCompany Report #A0142161A  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG / TWICE PER DAY / ORAL	Activated Partial Thromboplastin Time  Shortened  Bacterial Infection  Blood Potassium Decreased Ecchymosis Epistaxis Haematocrit Decreased Haemoglobin Decreased Idiopathic Thrombocytopenic Purpura Lymphocyte Count Increased Monocyte Count Increased Musculoskeletal Stiffness Neutrophil Count Increased Rectal Haemorrhage Red Blood Cell Count Decreased Specific Gravity Urine Abnormal White Blood Cell Count Increased	Foreign Health  Professional	Bupropion Hydrochloride   Clonazepam Aspirin	PS    C C	Glaxo Wellcome Inc	ORAL

Date:05/01/01ISR Number: 3715202-7Report Type:Expedited (15-DaCompany Report #A0144646A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG / TWICE PER DAY / ORAL	Dermatitis Dizziness  Hyperhidrosis  Middle Insomnia	Foreign Health  Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Nausea  
Tremor

Hydrochlorothiazide C  
Lisinopril C  
Rofecoxib C  
Conjugated Estrogens C

Date:05/02/01ISR Number: 3714664-9Report Type:Expedited (15-DaCompany Report #A0139332A

Age:7 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Teeth Brittle		Wellbutrin	PS	Glaxo Wellcome	ORAL
6 MON				No Concurrent Medications	C		

Date:05/02/01ISR Number: 3714671-6Report Type:Expedited (15-DaCompany Report #B0099812A

Age: Gender:Female I/FU:F

Outcome	PT
Other	Anaphylactic Shock Blood Pressure Increased Dizziness Dry Mouth Dry Skin

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Food Allergy Headache Malaise					
		Nausea Pallor Palpitations Paraesthesia Pharyngeal Oedema Sneezing Tongue Oedema Vertigo Vision Blurred		Zyban Quinapril Trisequens Ketoconazole	PS C C C	Glaxo Wellcome	

Date:05/02/01ISR Number: 3714677-7Report Type:Expedited (15-DaCompany Report #B0102612A  
Age:45 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other WK			Metrorrhagia		Zyban	PS	Glaxo Wellcome	
					Levonorgestrel	C		
					Diazepam	C		ORAL

INTRA-UTERINE 4 YR  
10MG Per day

Date:05/02/01ISR Number: 3714679-0Report Type:Expedited (15-DaCompany Report #B0104340A  
Age:51 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day Initial or Prolonged 400MG Per day 750MG per day			Cholelithiasis Pancreatitis Acute		Zyban Nevirapine Combivir Co-Trimoxazole Carbasalate Alcohol	PS SS SS C C C	Glaxo Wellcome Glaxo Wellcome	

3U Per day

Date:05/02/01ISR Number: 3714684-4Report Type:Expedited (15-DaCompany Report #B0105086A  
Age:38 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150MG As	Dry Mouth		Zyntabac	PS	Glaxo Wellcome	ORAL
Initial or Prolonged directed	Dyspnoea					
	Laryngeal Oedema Throat Irritation					

Date:05/02/01ISR Number: 3714685-6Report Type:Expedited (15-DaCompany Report #B0105236A  
Age:51 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening 150MG As	Arrhythmia		Zyban	PS	Glaxo Wellcome	
Hospitalization - directed 7 DAY	Myocardial Infarction					
Initial or Prolonged						

Date:05/02/01ISR Number: 3714687-XReport Type:Expedited (15-DaCompany Report #D0013944A  
Age:37 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Anxiety
Initial or Prolonged	Bronchospasm
	Chest Pain

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG See text	6 DAY	Convulsion Dizziness Dyspnoea Paralysis Tachycardia	Zyban	PS	Glaxo Wellcome	ORAL

Date:05/03/01ISR Number: 3716074-7Report Type:Expedited (15-DaCompany Report #A0130681A  
Age:21 YR Gender:Male I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 2 MON Initial or Prolonged Other	2 MON	Completed Suicide Fall Grand Mal Convulsion Humerus Fracture Intentional Misuse	Wellbutrin Alcohol No Concurrent Medications Marijuana	PS C C C	Glaxo Wellcome	ORAL

Date:05/03/01ISR Number: 3716083-8Report Type:Expedited (15-DaCompany Report #B0099476A  
Age:60 YR Gender:Male I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 5 WK Initial or Prolonged 10G per day	5 WK	Constipation Decreased Appetite Diverticulum Erythema Hallucination Insomnia	Zyban Temazepam	PS C	Glaxo Wellcome	ORAL

Date:05/03/01ISR Number: 3716084-XReport Type:Expedited (15-DaCompany Report #B0101144A  
Age: Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Circulatory Collapse	Zyban	PS	Glaxo Wellcome	

50MG At night	Dyskinesia	Dothiepin	C
10MG At night	Formication	Temazepam	C
	Heart Rate Increased	Oestradiol	C
	Hyperhidrosis		
	Muscle Contractions		
	Involuntary		
	Nausea		
	Night Sweats		
	Palpitations		
	Panic Attack		
	Tremor		

Date:05/03/01ISR Number: 3716094-2Report Type:Expedited (15-DaCompany Report #B0105094A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Epilepsy		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -		Loss Of Consciousness					
Initial or Prolonged		Overdose					
Other							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/03/01ISR Number: 3716098-XReport Type:Expedited (15-DaCompany Report #B0105940A

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Duration Aphasia Monoplegia Tunnel Vision		Zyban	PS	Glaxo Wellcome	

Date:05/03/01ISR Number: 3717044-5Report Type:Expedited (15-DaCompany Report #B0100732A

Age:38 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150 MG	Duration Anxiety	Foreign	Zyban	PS	Glaxo Wellcome Inc	
	Palpitations Tremor Vein Pain	Health Professional				

Date:05/04/01ISR Number: 3716877-9Report Type:Expedited (15-DaCompany Report #A0146512A

Age:59 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice Initial or Prolonged per day	Duration Anxiety Circulatory Collapse		Wellbutrin	PS	Glaxo Wellcome	OTHER
	Drug Level Above Therapeutic Parkinson'S Disease Tremor		Paxil Depakote Celexa	SS SS SS		

Date:05/04/01ISR Number: 3716885-8Report Type:Expedited (15-DaCompany Report #B0105221A

Age:55 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Duration Anaphylactoid Reaction Angioneurotic Oedema		Zyban	PS	Glaxo Wellcome	ORAL

Anxiety  
Circulatory Collapse  
Flushing  
Irritability  
Irritable Bowel Syndrome  
Rash Erythematous  
Swelling  
Urticaria

Date:05/04/01ISR Number: 3717519-9Report Type:Direct  
Age:39 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - 1 TABLET Initial or Prolonged TWICE DAILY Disability ORAL Other Required Intervention to Prevent Permanent Impairment/Damage	Cerebrovascular Accident Embolism Hemiplegia		Wellbutrin Sr 150mg Glaxo Wellcome	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/04/01ISR Number: 3718211-7Report Type:Expedited (15-DaCompany Report #B0104997A  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 300 MG/PER Required DAY/ORAL Intervention to Prevent Permanent Impairment/Damage		Diabetes Mellitus	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:05/04/01ISR Number: 3718213-0Report Type:Expedited (15-DaCompany Report #B0105091A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 150 MG/PER Intervention to DAY/ORAL Prevent Permanent Impairment/Damage		Inflammation  Swelling	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Beclomethasone Dipropion Salbutamol Sulphate	 C C		

Date:05/04/01ISR Number: 3718216-6Report Type:Expedited (15-DaCompany Report #B0105093A  
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 150 MG/TWICE Intervention to PER DAY/ORAL Prevent Permanent Impairment/Damage		Atrial Fibrillation  Gastroenteritis  Helicobacter	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Dipyridamole Aspirin Amlodipine Omeprazole	 C C C C		

Date:05/04/01ISR Number: 3718217-8Report Type:Expedited (15-DaCompany Report #B0105005A  
Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Convulsion	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG/VARIABL							
Hospitalization -		Head Injury					
E DOSE/ORAL							
Initial or Prolonged		Urinary Tract Infection		Dianette	C		
Required				Salbutamol Sulphate	C		
Intervention to							
Prevent Permanent							
Impairment/Damage							

Date:05/04/01ISR Number: 3718413-XReport Type:Expedited (15-DaCompany Report #A0145759A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
100 MG / PER							
		Lethargy					
DAY / ORAL							
		Peripheral Ischaemia					
		Sedation					
		Ventricular Hypertrophy					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/04/01ISR Number: 3719166-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #PRIUSA2000008733

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company	Ultram	PS	Rw Johnson Pharmaceutical Research Institute	ORAL
ORAL			Representative	Bupropion Hydrochloride (Amfebutamone Hydrochloride)	SS		

Date:05/04/01ISR Number: 3720085-5Report Type:Expedited (15-DaCompany Report #NSADSS2001012819  
Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Foreign Study Health Professional	Remicade (Infliximab, Recombinant)	PS		
INTRAVENOUS	5 MG/KG, 1 IN						
1 TIME(S), IV				Wellbutrin (Amfebutamone Hydrochloride)	SS		
				Pentasa	C		
				Losec	C		
				Buscopan	C		
				Demerol	C		
				Maxeran	C		
				Vitamin B12	C		
				Entocort	C		
				Imuran	C		

Date:05/07/01ISR Number: 3717303-6Report Type:Expedited (15-DaCompany Report #B0096539A  
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Discomfort		Bupropion			

150MG Per day 2 DAY  
Nausea

Eyelid Oedema

Hydrochloride

PS

Glaxo Wellcome

ORAL

Date:05/07/01ISR Number: 3717304-8Report Type:Expedited (15-DaCompany Report #B0098194A  
Age:21 YR Gender:Female I/FU:F

Outcome

PT

Death

Benign Lung Neoplasm  
Dizziness  
Drug Hypersensitivity  
Ecchymosis  
Encephalitis  
Epilepsy  
Epistaxis  
Fall  
Fatigue  
Feeling Abnormal  
Gastritis Erosive  
Head Injury  
Headache  
Hearing Impaired  
Hepatic Congestion

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	16 DAY	Hepatitis Intracranial Pressure Increased	Zyban	PS	Glaxo Wellcome	ORAL
		Livedo Reticularis				
		Loss Of Consciousness	Chloroquine	SS		
		Lymphadenopathy	Diphenhydramine	SS		
		Meningitis	Medroxy Progesterone Acetate	C		
		Ocular Hyperaemia				
		Pallor				
		Pleural Haemorrhage				
		Pulmonary Congestion				
		Pulmonary Oedema				
		Pulse Absent				
		Respiratory Arrest				
		Rhinitis				
		Sudden Death				
		Viral Infection				
		Vision Blurred				
		Vomiting				

Date:05/07/01ISR Number: 3717305-XReport Type:Expedited (15-DaCompany Report #B0100897A  
Age:44 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Death		PT				
150MG As		Arrhythmia	Zyban	PS	Glaxo Wellcome	
directed	17 DAY	Cardiac Arrest				
		Coma	Paracetamol	C		
		Hypertension				
		Hypotension				
		Respiratory Failure				
		Subarachnoid Haemorrhage				

Date:05/07/01ISR Number: 3717306-1Report Type:Expedited (15-DaCompany Report #B0105726A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Death		PT				
		Anger	Zyban	PS	Glaxo Wellcome	

Completed Suicide  
Irritability  
Psychomotor Hyperactivity

Date:05/07/01ISR Number: 3718428-1Report Type:Expedited (15-DaCompany Report #B0099476A  
Age:60 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - ORAL Initial or Prolonged	Abdominal Pain Constipation Decreased Appetite Diverticulum Erythema Hallucination Insomnia	Foreign Health Professional	Zyban Temazepam	PS C	Glaxo Wellcome Inc	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/07/01ISR Number: 3718431-1Report Type:Expedited (15-DaCompany Report #B0101144A

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150 MG	Duration Circulatory Collapse	Foreign	Zyban	PS	Glaxo Wellcome Inc	
	Formication	Health	Dothiepin	C		
	Heart Rate Increased	Professional	Temazepam	C		
	Hyperhidrosis		Oestradiol	C		
	Muscle Contractions Involuntary Night Sweats Palpitations Panic Attack Tremor					

Date:05/07/01ISR Number: 3719009-6Report Type:Expedited (15-DaCompany Report #B0105094A

Age:39 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150 MG / ORAL	Duration Epilepsy	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage	Loss Of Consciousness Overdose	Health Professional				

Date:05/07/01ISR Number: 3719010-2Report Type:Expedited (15-DaCompany Report #B0105940A

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Duration Aphasia Movement Disorder Tunnel Vision	Foreign Consumer	Zyban	PS	Glaxo Wellcome Inc	

Date:05/07/01ISR Number: 3719021-7Report Type:Expedited (15-DaCompany Report #A0139332A  
Age:7 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 150 MG / ORAL		Teeth Brittle	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Intervention to Prevent Permanent Impairment/Damage		Tooth Disorder	Professional Company Representative				

Date:05/07/01ISR Number: 3719175-2Report Type:Expedited (15-DaCompany Report #B0105086A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / AS		Accident	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged DIRECTED		Dry Mouth	Health				
/ORAL		Dyspnoea	Professional				
		Laryngeal Oedema Throat Irritation					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/07/01ISR Number: 3719177-6Report Type:Expedited (15-DaCompany Report #B0105236A  
Age:51 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150 MG/AS Hospitalization - DIRECTED/ Initial or Prolonged	Arrhythmia Myocardial Infarction	Foreign	Zyban	PS	Glaxo Wellcome Inc	

Date:05/07/01ISR Number: 3719180-6Report Type:Expedited (15-DaCompany Report #B0102612A  
Age:45 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required Intervention to Prevent Permanent Impairment/Damage	Metrorrhagia	Foreign Health Professional	Bupropion Hydrochloride Levonorgestrel Diazepam	PS C C	Glaxo Wellcome Inc	

Date:05/07/01ISR Number: 3719183-1Report Type:Expedited (15-DaCompany Report #B0099812A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG	Anaphylactic Shock	Foreign	Zyban	PS	Glaxo Wellcome Inc	
	Dehydration	Health	Quinapril	C		
	Dizziness	Professional	Trisequens	C		
	Dry Mouth		Ketoconazole	C		
	Dry Skin					
	Feeling Abnormal					
	Food Allergy					
	Headache					
	Hypersensitivity					
	Hypertension					
	Malaise					
	Nausea					
	Pallor					
	Palpitations					
	Paraesthesia					
	Pharyngeal Oedema					

Tongue Oedema  
Vertigo  
Vision Blurred

Date:05/07/01ISR Number: 3719214-9Report Type:Expedited (15-DaCompany Report #B0104340A  
Age:51 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG/ PER	Abdominal Pain Abdominal Tenderness	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
DAY/	Blood Amylase Increased					
400 MG/ PER	Cholelithiasis Pallor Pancreatitis Acute		Nevirapine (Formulation Unknown)	SS		
DAY	Vomiting					
			Combivir (Formulation Unknown) (Combivir)	SS		
			Co-Trimoxazole	C		
			Carbaspirin Calcium	C		
			Ethanol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/07/01ISR Number: 3719217-4Report Type:Expedited (15-DaCompany Report #D0013944A

Age:37 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/ SEE	Anxiety Bronchospasm	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TEXT/ ORAL	Chest Pain	Professional				
	Convulsion Dizziness Dyspnoea Paralysis					

Date:05/07/01ISR Number: 3719244-7Report Type:Expedited (15-DaCompany Report #A0130681A

Age:21 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / Initial or Prolonged PER DAY / Other ORAL	Fall Grand Mal Convulsion	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
	Humerus Fracture	Company				
	Intentional Misuse Suicide Attempt	Representative	Ethanol No Concurrent Medication Cannabis	C C C		

Date:05/08/01ISR Number: 3718282-8Report Type:Expedited (15-DaCompany Report #A0145045A

Age:31 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200MG Twice per day 60MG Per day	Complications Of Maternal Exposure To Therapeutic Drugs		Wellbutrin Celexa	PS SS	Glaxo Wellcome	ORAL ORAL
	Stillbirth					

Date:05/08/01ISR Number: 3718287-7Report Type:Expedited (15-DaCompany Report #A0146305A

Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
150MG Per day	10 DAY			Synthroid	C	Glaxo Wellcome	

Date:05/08/01ISR Number: 3718884-9Report Type:Direct

Company Report #

Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis		Zyban	PS		
				Fosinopril Na	C		
				Simvastatin	C		
				Ibuprofen	C		
				Diphenhydramine Hcl	C		
				Triamcinolone			
				Acetonide	C		
				Nicotine	C		
				Quit Smart Bupropion			
				Sr	C		
				Hydrocortisone	C		
				Furosemide	C		
				Potassium Chloride	C		
				Beclomethasone	C		

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Freedom Of Information (FOI) Report

Date:05/08/01ISR Number: 3719692-5Report Type:Expedited (15-DaCompany Report #A0146512A  
Age:59 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY / PERIPHERAL LINE	Anxiety Circulatory Collapse Difficulty In Walking Parkinson'S Disease Tremor	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	
			Paroxetine Hydrochloride (Formulation Unknown)	SS		

Date:05/08/01ISR Number: 3719867-5Report Type:Expedited (15-DaCompany Report #B0105221A  
Age:55 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Abdominal Pain Upper Anaphylactic Reaction Angioneurotic Oedema Anxiety Circulatory Collapse Diarrhoea Flushing Irritability Irritable Bowel Syndrome Rash Erythematous Urticaria	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/10/01ISR Number: 3720063-6Report Type:Expedited (15-DaCompany Report #B0093206A  
Age:35 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 150MG Twice	Arrhythmia		Zyban	PS	Glaxo Wellcome	ORAL

Coronary Artery Disease

per day 47 DAY

20MG per day

1TAB At night

UNKNOWN 70MG At night

10ML Four

times per day

409ML per day

Pantoprazole	C	ORAL
Chlormethiazole	C	ORAL
Lofepramine	C	
Gaviscon	C	ORAL
Lactulose	C	
Desipramine	C	
Tramadol	C	
Ecstasy	C	

Date:05/10/01ISR Number: 3720071-5Report Type:Expedited (15-DaCompany Report #B0106124A  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hearing Impaired		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	5 DAY	Photopsia					

Date:05/10/01ISR Number: 3720072-7Report Type:Expedited (15-DaCompany Report #B0106152A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Nail Disorder		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	29 DAY	Skin Exfoliation					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/10/01ISR Number: 3720073-9Report Type:Expedited (15-DaCompany Report #B0106347A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Furuncle		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day							

Date:05/10/01ISR Number: 3720074-0Report Type:Expedited (15-DaCompany Report #B0106364A

Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eye Pain		Zyban	PS	Glaxo Wellcome	
4 DAY							
1TAB Per day		Eyelid Oedema		Levonorgestrel	C		ORAL
		Headache					

Date:05/10/01ISR Number: 3720075-2Report Type:Expedited (15-DaCompany Report #B0106366A

Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Asthma		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	7 DAY						
RESPIRATORY		Chest Discomfort		Salbutamol	C	Glaxo Wellcome	
(INHALATION)	8PUFF As	Hypoventilation					
required							
RESPIRATORY				Budesonide	C		
(INHALATION)	400MCG Twice						
per day							
RESPIRATORY				Ipratropium	C	Glaxo Wellcome	

(INHALATION) 2PUFF Twice

per day

Migraleve Pink C ORAL

Date:05/10/01ISR Number: 3720076-4Report Type:Expedited (15-DaCompany Report #B0106367A

Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Twice		Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - per day	21 DAY	Dermatitis					
Initial or Prolonged 1TAB Per day		Dyspnoea		Microgynon	C		ORAL
Disability		Visual Disturbance					

Date:05/10/01ISR Number: 3721582-9Report Type:Expedited (15-DaCompany Report #B0105726A

Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG		Aggression	Foreign	Zyban	PS	Glaxo Wellcome Inc	
		Completed Suicide Irritability Psychomotor Hyperactivity	Consumer				

Date:05/10/01ISR Number: 3721612-4Report Type:Expedited (15-DaCompany Report #B0096539A

Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/PER DAY/ORAL		Discomfort Eyelid Oedema	Foreign Study	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Nausea	Health				
		Oesophageal Disorder	Professional				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/10/01ISR Number: 3721614-8Report Type:Expedited (15-DaCompany Report #B0100897A

Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia	Foreign	Bupropion			
150 MG/AS		Atrioventricular Block	Health	Hydrochloride	PS	Glaxo Wellcome Inc	
DIRECTED		Coma	Professional				
		Headache		Paracetamol	C		
		Hypertension					
		Hypotension					
		Musculoskeletal Stiffness					
		Myalgia					
		Respiratory Failure					
		Subarachnoid Haemorrhage					

Date:05/10/01ISR Number: 3721616-1Report Type:Expedited (15-DaCompany Report #B0098194A

Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Benign Lung Neoplasm	Foreign	Bupropion			
150 MG/PER		Convulsion	Literature	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
DAY/ORAL		Cyanosis	Health				
		Dizziness	Professional	Chloroquine			
		Dysgeusia		(Chloroquine)	SS		
		Ecchymosis		Diphenhydramine			
		Embolism		(Diphenhydramine)	SS		
		Encephalitis		Medroxyprogesterone			
		Epilepsy		Ace	C		
		Epistaxis					
		Fall					
		Fatigue					
		Feeling Abnormal					
		Gastritis Erosive					
		Headache					
		Hearing Impaired					
		Hepatic Congestion					
		Hepatitis					
		Inflammation					
		Injury					

Intracranial Pressure  
Increased  
Loss Of Consciousness  
Lung Disorder  
Lymphadenopathy  
Nasal Congestion  
Ocular Hyperaemia  
Pallor  
Petechiae  
Pleural Haemorrhage  
Pulmonary Congestion  
Pulmonary Oedema  
Pulse Absent  
Respiratory Arrest  
Respiratory Distress  
Skin Discolouration  
Sudden Death  
Syncope  
Vision Blurred  
Vomiting

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/11/01ISR Number: 3720855-3Report Type:Expedited (15-DaCompany Report #A0144832A  
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Single Initial or Prolonged dose	7 DAY	Complex Partial Seizures		Zyban	PS	Glaxo Wellcome	ORAL
		Confusional State Hallucination, Visual Psychotic Disorder		Alcohol	SS		

Date:05/11/01ISR Number: 3720862-0Report Type:Expedited (15-DaCompany Report #B0101512A  
Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Per day Other	6 WK	Diabetes Mellitus		Zyban	PS	Glaxo Wellcome	ORAL
				Hypoglycaemic Medication	C		ORAL
				Viagra	C		ORAL
				Lipostat	C		
				Tritace	C		
				Aspirin	C		
				Gliclazide	C		ORAL
80MG per day							

Date:05/11/01ISR Number: 3720866-8Report Type:Expedited (15-DaCompany Report #B0105221A  
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Other		Abdominal Pain Amnesia Anaphylactic Reaction Angioneurotic Oedema Anxiety Circulatory Collapse Diarrhoea Disturbance In Attention Dry Skin		Zyban	PS	Glaxo Wellcome	ORAL

Flushing  
Irritability  
Rash Erythematous  
Rash Pruritic  
Swelling  
Urticaria

Date:05/11/01ISR Number: 3720871-1Report Type:Expedited (15-DaCompany Report #B0106121A  
Age:42 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Agitation
Initial or Prolonged	Anxiety
Other	Asthenia
	Bronchospasm
	Chest Discomfort
	Depression
	Dermatitis
	Disturbance In Attention
	Dry Mouth
	Headache
	Hyperhidrosis
	Insomnia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Tinnitus				
		Tremor				
		Visual Disturbance	Report Source	Product	Role	Manufacturer
Dose	Duration			Zyban	PS	Glaxo Wellcome
150MG	Per day					ORAL

Date:05/11/01ISR Number: 3720872-3Report Type:Expedited (15-DaCompany Report #B0106128A  
 Age:50 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer
Outcome	Duration			Zyban	PS	Glaxo Wellcome
Dose		Blister				ORAL
Other		Joint Stiffness				
150MG	Twice	Oedema				
per day	15 DAY	Pain In Extremity				
		Proteinuria				
		Rash Erythematous				

Date:05/11/01ISR Number: 3720873-5Report Type:Expedited (15-DaCompany Report #B0106260A  
 Age:37 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer
Outcome	Duration			Zyban	PS	Glaxo Wellcome
Dose		Agitation				ORAL
Other		Depression				
150MG	Twice	Insomnia				
per day	15 DAY	Paranoia				
		Suicidal Ideation				

Date:05/11/01ISR Number: 3720874-7Report Type:Expedited (15-DaCompany Report #B0106345A  
 Age:60 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer
Outcome	Duration			Zyban	PS	Glaxo Wellcome
Dose		Facial Palsy				ORAL
Other						
150MG						

Hypertension

Variable dose 41 DAY

Date:05/11/01ISR Number: 3720875-9Report Type:Expedited (15-DaCompany Report #B0106348A

Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	15 DAY			Zyban	PS	Glaxo Wellcome	
Hospitalization - Initial or Prolonged		Circulatory Collapse Loss Of Consciousness Myocardial Infarction Ventricular Fibrillation					

Date:05/11/01ISR Number: 3721857-3Report Type:Expedited (15-DaCompany Report #A0145045A

Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	200 MG / TWICE PER DAY / ORAL	Complications Of Maternal Exposure To Therapeutic Drugs Intra-Uterine Death Stillbirth	Study Health Professional Other	Wellbutrin Sr   Citalopram Hydrobromide Tablet (Citalopram Hydrobromide)	PS   SS	Glaxo Wellcome Inc	ORAL

OROPHARINGEAL 60 MG /

PER DAY /

ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/11/01ISR Number: 3721864-0Report Type:Expedited (15-DaCompany Report #A0146305A

Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Study	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER		Complications Of Maternal	Consumer				
DAY/ ORAL		Exposure To Therapeutic Drugs Pregnancy		Thyroxine Sodium	C		

Date:05/11/01ISR Number: 3721948-7Report Type:Expedited (15-DaCompany Report #10819928

Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dermatitis	Health	Paraplatin	PS	Bristol Myers Squibb	
Initial or Prolonged		Dyspnoea	Professional			Co Pharmaceutical	
Other		Feeling Abnormal				Research Institute	
INTRAVENOUS	640	Flushing					
MILLIGRAM,		Hypersensitivity					
1/1 CYCLE IV		Petit Mal Epilepsy		Wellbutrin (Bupropion Hcl)	SS		

Date:05/14/01ISR Number: 3721874-3Report Type:Expedited (15-DaCompany Report #B0104334A

Age:22 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dysuria		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Myocardial Infarction					
Initial or Prolonged							
per day	23 DAY						
Other		Oedema Peripheral		Adrenaline	SS		
INTRAVENOUS		Urticaria		Citalopram	C		

Date:05/14/01ISR Number: 3721876-7Report Type:Expedited (15-DaCompany Report #B0106581A  
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abnormal Behaviour Completed Suicide Road Traffic Accident		Zyban	PS	Glaxo Wellcome	ORAL

Date:05/14/01ISR Number: 3721878-0Report Type:Expedited (15-DaCompany Report #B0106828A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Brain Oedema		Zyban	PS	Glaxo Wellcome	
Hospitalization - Initial or Prolonged		Cerebral Infarction Dysphagia Hemiparesis		Contraceptive	C		ORAL

Date:05/14/01ISR Number: 3721879-2Report Type:Expedited (15-DaCompany Report #B0106830A  
Age:65 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Abdominal Discomfort Diarrhoea Inflammation Nausea Nephritis

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vomiting

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1 WK			Zyban	PS	Glaxo Wellcome	

Date:05/14/01ISR Number: 3721883-4Report Type:Expedited (15-DaCompany Report #B0107035A  
Age:19 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Zyban	PS	Glaxo Wellcome	ORAL
2 MON		Depression Dissociation Disturbance In Attention Feeling Abnormal Headache Palpitations		Chloroquine	C		

Date:05/14/01ISR Number: 3721884-6Report Type:Expedited (15-DaCompany Report #B0107040A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Choking Cyanosis Loss Of Consciousness		Zyban Nitrazepam	PS C	Glaxo Wellcome	ORAL ORAL

Date:05/14/01ISR Number: 3723935-1Report Type:Expedited (15-DaCompany Report #S01-USA-00846-01  
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Circulatory Collapse Parkinson'S Disease Tremor	Consumer	Celexa	PS	Forest Laboratories Inc	
				Wellbutrin(Amfebutam one Hydrochloride)	SS		
				Depakote(Valproate Semidsodium)	SS		

Paxil (Paroxetine  
Hydrochloride) SS

Date:05/14/01ISR Number: 3724057-6Report Type:Expedited (15-DaCompany Report #B0106367A  
Age:16 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY Disability / ORAL	Arthralgia Dermatitis Dizziness Dyspnoea Visual Disturbance	Foreign	Bupropion Hydrochloride  Microgynon	PS  C	Glaxo Wellcome Inc	ORAL

Date:05/14/01ISR Number: 3724065-5Report Type:Expedited (15-DaCompany Report #B0106152A  
Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required 150 MG/ PER Intervention to DAY/ ORAL Prevent Permanent Impairment/Damage	Dermatitis Exfoliative Nail Disorder	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/14/01ISR Number: 3724066-7Report Type:Expedited (15-DaCompany Report #B0106347A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Furuncle	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
Intervention to							
PER DAY/ ORAL							
Prevent Permanent							
Impairment/Damage							

Date:05/14/01ISR Number: 3724067-9Report Type:Expedited (15-DaCompany Report #B0106364A

Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Eye Pain	Foreign	Zyban	PS	Glaxo Wellcome Inc	
150 MG/ UNK							
Intervention to							
Prevent Permanent							
Impairment/Damage							
Eyelid Oedema							
Headache							
Levonorgestrel							
C							

Date:05/14/01ISR Number: 3724069-2Report Type:Expedited (15-DaCompany Report #B0106124A

Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Hearing Impaired	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							
Intervention to							
DAY/ ORAL							
Prevent Permanent							
Impairment/Damage							
Migraine							
Photopsia							

Date:05/14/01ISR Number: 3724070-9Report Type:Expedited (15-DaCompany Report #B0106366A

Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Disability	Asthma	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER						
DAY/ ORAL						
			Salbutamol Sulphate	C		
			Budesonide	C		
			Ipratropium Bromide	C		
			Migraleve (Pink			
			Tablets)	C		

Date:05/14/01ISR Number: 3724079-5Report Type:Expedited (15-DaCompany Report #B0093206A  
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Coronary Artery Disease	Health				
PER DAY/ ORAL			Professional	Pantoprazole	C		
			Company	Chlormethiazole	C		
			Representative	Lofepamine	C		
				Gaviscon	C		
				Lactulose	C		
				Desipramine	C		
				Tramadol			
				Hydrochloride	C		
				Methylenedioxymetham			
				phet.	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/14/01  
 ISR Number: 3726066-X  
 Report Type:Periodic  
 Age:37 YR Gender:Male I/FU:I

Company Report #RS001648-USA

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
20 MG, PER		Blood Glucose Increased	Consumer	Aciphex	PS	Eisai Inc	ORAL
ORAL		Libido Decreased					
PER ORAL		Muscle Rigidity Pollakiuria		Zyban (Amfetbutamone Hydrochloride)	SS		ORAL
				Tagamet (Cimetidine)	C		
				Pepcid (Famotidine)	C		
				Maalox (Maalox)	C		
				Tums (Tums)	C		
				Other Medications (Unspecified)	C		
				Cephalexin (Cefalexin)	C		

Date:05/15/01  
 ISR Number: 3722915-X  
 Report Type:Expedited (15-Da  
 Company Report #A0142161A  
 Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening 150MG Twice		Abnormal Behaviour		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - per day		Activated Partial					
Initial or Prolonged .5MG Twice		Thromboplastin Time		Rivotril	C		ORAL
Disability per day		Shortened					
		Asthenia		Asa	C		
		Bacillus Infection					
		Bacterial Infection					
		Blister					
		Blood Cholesterol Increased					
		Blood Potassium Decreased					
		Decreased Activity					
		Ecchymosis					
		Fatigue					
		Haematocrit Decreased					

Haemoglobin Decreased  
Idiopathic  
Thrombocytopenic Purpura  
Infection  
Lymphocyte Count  
Increased  
Monocyte Count Increased  
Musculoskeletal Stiffness  
Nervous System Disorder  
Neutrophil Count  
Increased  
Oedema Peripheral  
Pain  
Platelet Count Decreased  
Rectal Haemorrhage  
Red Blood Cell Count  
Decreased  
Specific Gravity Urine  
Abnormal  
Spleen Disorder  
Thrombocytopenia  
Visual Disturbance  
Weight Increased  
White Blood Cell Count  
Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/15/01ISR Number: 3722930-6Report Type:Expedited (15-DaCompany Report #B0106346A

Age:27 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Per day Initial or Prolonged	Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
	Back Pain		Cilest	C		ORAL
	Depression		Senna	C		ORAL
	Disturbance In Attention					
	Dizziness					
	Headache					
	Indifference					
	Nausea					
	Neck Pain					
	Optic Discs Blurred					
	Sedation					
	Staring					

Date:05/15/01ISR Number: 3724222-8Report Type:Direct

Age:34 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required TOBACCO	Erythema		Zyban 150 Mg Skb	PS	Skb	
Intervention to CESSATION	Urticaria					
Prevent Permanent Impairment/Damage			Nicoderm	C		

Date:05/15/01ISR Number: 3724223-XReport Type:Direct

Age:20 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 150 MG BID	Rash Pruritic		Zyban 150mg Skb	PS	Skb	ORAL
Intervention to ORAL						
Prevent Permanent Impairment/Damage			Nicoderm	C		
			Colace	C		
			Loovral	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Diverticulum Oesophageal		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	124 DAY	Dry Mouth					
Disability 75MG Twice		Gastric Polyps		Effexor Xr	SS		ORAL
Other per day	3 YR	Gingival Pain					
		Hypokalaemia		Nicotine	SS		
		Hyponatraemia		Trazodone	C		ORAL
50MG At night		Pituitary Tumour		Klonopin	C		ORAL
.5MG As		Post Procedural					
required		Complication		Dextrostat	C		ORAL
10MG Twice		Salivary Hypersecretion					
per day	700 DAY	Speech Disorder		Synthroid	C	Glaxo Wellcome	ORAL
		Stress					
		Tardive Dyskinesia					
		Visual Field Defect					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/16/01ISR Number: 3723640-1Report Type:Expedited (15-DaCompany Report #B0099812A

Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain		Zyban	PS	Glaxo Wellcome	
150MG Twice		Accident					
per day		Asthenia		Quinapril	C		
		Blood Pressure Increased		Trisequens	C		
		Dehydration		Ketoconazole	C		
		Diplopia		Alcohol	C		
		Dizziness					
		Dry Mouth					
		Dry Skin					
		Feeling Abnormal					
		Food Allergy					
		Fungal Infection					
		Headache					
		Hypersensitivity					
		Malaise					
		Nausea					
		Pallor					
		Palpitations					
		Paraesthesia					
		Pharyngeal Oedema					
		Sneezing					
		Tongue Oedema					
		Vertigo					
		Vision Blurred					

Date:05/16/01ISR Number: 3723641-3Report Type:Expedited (15-DaCompany Report #B0104518A

Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Asthenia		Zyban	PS	Glaxo Wellcome	
UNKNOWN	150MG	Per day					
Hospitalization -		Chest Pain		Atenolol	C		ORAL
50MG Per day							
Initial or Prolonged		Cough		Aspirin	C		ORAL
75MG In the							
Disability		Dizziness					
morning							

200MG Twice	Lower Respiratory Tract	Dipyridamole	C	ORAL
per day	Infection			
10MG Per day	Nausea	Atorvastatin	C	ORAL
1.25MG Per	Neck Pain	Premarin	C	ORAL
day				

Date:05/16/01ISR Number: 3723644-9Report Type:Expedited (15-DaCompany Report #B0107247A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hypoaesthesia Paralysis		Zyban	PS	Glaxo Wellcome	

Date:05/16/01ISR Number: 3723645-0Report Type:Expedited (15-DaCompany Report #B0107381A  
Age:40 YR Gender:Male I/FU:F

Outcome	PT
Death	Amnesia Balance Disorder Confusional State Ecchymosis Excoriation

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		Fall Head Injury Hypoaesthesia					
		Overdose		Zyban	PS	Glaxo Wellcome	
		Periorbital Haematoma		Fluoxetine	C		
		Speech Disorder		Alcohol	C		
		Subdural Haematoma					
		Toxicologic Test Abnormal					

Date:05/16/01ISR Number: 3724981-4Report Type:Expedited (15-DaCompany Report #2001-BP-01469  
Age:65 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 0.4MG/1 Initial or Prolonged CAPSULE / QD/ PO	350 MG / PO	7 YR	Cardio-Respiratory Arrest Convulsion Tachycardia	Health Professional	Flomax	PS	Boehringer Ingelheim Pharmaceuticals Inc	ORAL
					Wellbutrin	SS		ORAL
					Lipitor	C		
					Lorazepam	C		
					Viagra	C		
					Baby Aspirin	C		
					Msm	C		
					Co Q10	C		
					Multivitamins	C		
					Fish Oil	C		
					Pentatonic Acid	C		

Date:05/17/01ISR Number: 3724257-5Report Type:Expedited (15-DaCompany Report #B0107423A  
Age:49 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice			Abdominal Distension		Zyban	PS	Glaxo Wellcome	ORAL

per day  
Depression  
Drug Withdrawal Syndrome

Date:05/17/01ISR Number: 3724393-3Report Type:Direct Company Report #  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Polyarthrititis		Zyban	PS		ORAL
1	BID PO			Nicoderm Patches	C		

Date:05/17/01ISR Number: 3725080-8Report Type:Expedited (15-DaCompany Report #B0107035A  
Age:19 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide Disturbance In Attention	Foreign Literature	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL		Feeling Abnormal Headache Palpitations		Chloroquine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/17/01ISR Number: 3725082-1Report Type:Expedited (15-DaCompany Report #B0104334A

Age:22 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG TWICE Required PER DAY ORAL	Dermatitis Dysuria Myocardial Infarction	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Intervention to Prevent Permanent INTRA VENOUS Impairment/Damage	Swelling Urticaria INTRA VENOUS	Company Representative	Adrenaline (Epinephrine) Citalopram	SS C		

Date:05/17/01ISR Number: 3725084-5Report Type:Expedited (15-DaCompany Report #B0106830A

Age:65 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged UNKNOWN	Abdominal Discomfort C-Reactive Protein Increased Chromaturia Diarrhoea Inflammation Nausea Nephritis Vomiting White Blood Cell Count Increased	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:05/17/01ISR Number: 3725086-9Report Type:Expedited (15-DaCompany Report #B0106581A

Age:61 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death ORAL	Abnormal Behaviour Road Traffic Accident	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/17/01ISR Number: 3725088-2Report Type:Expedited (15-DaCompany Report #B0106828A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Brain Oedema	Foreign	Bupropion			
Hospitalization -		Cerebral Infarction		Hydrochloride	PS	Glaxo Wellcome Inc	
UNKNOWN	150 MG						
Initial or Prolonged		Dysphagia					
UNKNOWN		Hemiparesis		Bith Controlpills	C		

Date:05/17/01ISR Number: 3725090-0Report Type:Expedited (15-DaCompany Report #B0107040A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Choking	Foreign	Bupropion			
		Loss Of Consciousness	Consumer	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL		Skin Discolouration		Nitrazepam	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/18/01ISR Number: 3724928-0Report Type:Expedited (15-DaCompany Report #B0092571A

Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Overdose		Zyban	PS	Glaxo Wellcome	ORAL
1TAB As							
Hospitalization -		Stress					
directed	34 DAY						
Initial or Prolonged		Suicide Attempt		Paracetamol	SS		
Other				Alcohol	SS		

Date:05/18/01ISR Number: 3724932-2Report Type:Expedited (15-DaCompany Report #B0099532A

Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Poor Peripheral		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	6 MON						
60MG In the		Circulation		Isib	C		ORAL
morning		Retinal Detachment					
		Vitreous Haemorrhage		Co-Codamol	C		
400MCG At				Cerivastatin	C		
night							
75MG Per day				Clopidogrel	C		
40MG In the				Frusamide	C		
morning							
4MG In the				Lacidipine	C	Glaxo Wellcome	
morning							
20MG In the				Fluoxetine	C		
morning							
75MG Per day				Aspirin	C		
				Gtn	C	Glaxo Wellcome	

RESPIRATORY		Beclomethasone	C	Glaxo Wellcome	
(INHALATION)	2PUFF Twice				
per day					
150MG Twice		Ranitidine	C	Glaxo Wellcome	
per day					
20MG At night		Amitriptyline	C		
7.5G At night		Senna	C		
RESPIRATORY		Salbutamol	C	Glaxo Wellcome	
(INHALATION)					
1000MG per		Naproxen	C		ORAL
day					
400MG per day		Trimethoprim	C	Glaxo Wellcome	ORAL
		Nicorandil	C		ORAL
		Cephradine	C	Glaxo Wellcome	
		Metronidazole	C		

Date:05/18/01ISR Number: 3724934-6Report Type:Expedited (15-DaCompany Report #B0101422A  
Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day	92 DAY	Eye Haemorrhage	Zyban	PS	Glaxo Wellcome	ORAL
			Therapeutic Response	Amitriptyline	C		ORAL
			Unexpected	Isosorbide	C		ORAL
			Vitreous Disorder	Co-Codamol	C		ORAL
				Cerivastatin	C		ORAL
				Clopidogrel	C		ORAL
				Frusemide	C		ORAL
				Lacidipine	C	Glaxo Wellcome	ORAL

20MG per day

75MG per day

300MG per day

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Fluoxetine	C		ORAL
Aspirin	C		ORAL
Ranitidine	C	Glaxo Wellcome	ORAL

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7.5MG per day	Senna	C		ORAL
RESPIRATORY	Salbutamol	C	Glaxo Wellcome	
(INHALATION)				
1000MG per	Naproxen	C		ORAL
day				
400MG per day	Trimethoprim	C	Glaxo Wellcome	ORAL
	Nicorandil	C		ORAL

Date:05/18/01ISR Number: 3725865-8Report Type:Expedited (15-DaCompany Report #A0142161A  
 Age:54 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Activated Partial	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /						
Hospitalization -	Thromboplastin Time	Health				
TWICE PER DAY						
Initial or Prolonged	Shortened	Professional				
/ ORAL						
Disability	Bacteriuria		Clonazepam	C		
	Blister		Aspirin	C		
	Blood Cholesterol					
	Increased					
	Blood Potassium Decreased					
	Ecchymosis					
	Epistaxis					
	Eye Disorder					
	Fatigue					
	Haematocrit Decreased					
	Haemoglobin Decreased					
	Idiopathic					
	Thrombocytopenic Purpura					
	Laboratory Test Abnormal					
	Lymphocyte Count					
	Increased					
	Mental Disorder					
	Monocyte Count Increased					
	Musculoskeletal Stiffness					
	Nervous System Disorder					

Neutrophil Count  
Increased  
Oedema Peripheral  
Pain  
Purpura  
Rectal Haemorrhage  
Red Blood Cell Count  
Decreased  
Specific Gravity Urine  
Abnormal  
Spleen Disorder  
Thrombocytopenia  
Weight Increased  
White Blood Cell Count  
Increased

Date:05/18/01ISR Number: 3725897-XReport Type:Expedited (15-DaCompany Report #B0106346A  
Age:27 YR Gender:Female I/FU:I

Outcome PT  
Hospitalization - Arthralgia  
Initial or Prolonged Back Pain  
Coordination Abnormal  
Depression

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Disturbance In Attention Dizziness Electroencephalogram Abnormal Feeling Abnormal Headache Nausea Neck Pain Papilloedema Sedation	Foreign	Bupropion Hydrochloride  Cileste Sennosides	PS  C C	Glaxo Wellcome Inc	ORAL

Date:05/18/01  
 Age:46 YR  
 Gender:Male  
 I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG ORAL	MON	Aortic Atherosclerosis Cardiomegaly Cardiovascular Disorder Coma Convulsion Coronary Artery Disease Drug Level Above Therapeutic Ecchymosis Emphysema Excoriation Hepatic Congestion Hepatic Fibrosis Hepatic Steatosis Histology Normal Hypertension Overdose Pulmonary Congestion Pulmonary Fibrosis Pulmonary Oedema Ventricular Hypertrophy Vomiting	Health  Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 50.00 MG		Akinesia	Consumer	Zoloft	PS	Pfizer	
Prevent Permanent TOTAL:DAILY: Impairment/Damage 300.00 MG		Blood Pressure Increased				Pharmaceuticals Inc	
TOTAL:BID:ORA		Drug Ineffective					
L		Dysphonia		Wellbutrin	SS		ORAL
		Ecchymosis					
		Epistaxis					
300.00 MG		Haemorrhagic Stroke		Coumadin	SS		
TOTAL:BID:ORA		Headache		Effexor Sr	SS		ORAL
L		Heart Rate Decreased					
		Mood Swings					
		Overdose		Toprol	C		
		Parkinson'S Disease		Zebeta	C		
		Staring		Lanoxin	C		
		Syncope		Lasix	C		
		Thyroid Function Test Abnormal		Synthroid	C		
				Vitamin E	C		
				Estrogen	C		

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Freedom Of Information (FOI) Report

Date:05/21/01ISR Number: 3725777-XReport Type:Expedited (15-DaCompany Report #A0139670A

Age:74 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day 2 MON	Cardiac Failure		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	INTRA VENOUS 1G Per day	Congestive		Rocephin	C		
	20MG Per day	Convulsion		Pepcid	C		
	12.5MG Twice	Loss Of Consciousness		Kcl	C		
	per day			Captopril	C	Glaxo Wellcome	
	500MG Per day			Depakote	C		
	10MG Twice			Buspar	C		
	per day						
	7MG At night			Nicoderm Patch	C		
	75MG Per day			Seroquel	C		
	12.5MG Per			Lasix	C		
	day			Lanoxin	C	Glaxo Wellcome	
	325MG Per day			Albuterol	C	Glaxo Wellcome	
	12.5MG Three			Spiro nolactone	C		
	times per day						
RESPIRATORY				Aspirin	C		
(INHALATION)				Antivert	C		
RESPIRATORY				Artificial Tears	C		
				Proventil	C	Glaxo Wellcome	
				Atrovent	C	Glaxo Wellcome	



( INHALATION)

Date:05/21/01ISR Number: 3725781-1Report Type:Expedited (15-DaCompany Report #B0100682A  
Age:32 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice Initial or Prolonged per day	Agitation		Zyban	PS	Glaxo Wellcome	
50MG Twice per day	Blindness					
	Blood Glucose Decreased		Coffee	C		ORAL
	Circulatory Collapse		Naprosyn	C		ORAL
	Dizziness					
	Headache					
	Hypoaesthesia					
	Photophobia					
	Vision Blurred					

Date:05/21/01ISR Number: 3725784-7Report Type:Expedited (15-DaCompany Report #B0107510A  
Age:53 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Per day 16 DAY Initial or Prolonged	Asthma		Zyban	PS	Glaxo Wellcome	ORAL
50MG As required			Aminophylline	C		ORAL
			Prednisolone	C		ORAL
50MG As required			Diclofenac	C		ORAL

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Freedom Of Information (FOI) Report

Date:05/21/01ISR Number: 3725785-9Report Type:Expedited (15-DaCompany Report #B0107658A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	7 DAY	Agitation Disturbance In Attention Feeling Abnormal Speech Disorder Tinnitus Visual Disturbance		Zyban	PS	Glaxo Wellcome	ORAL

Date:05/21/01ISR Number: 3725787-2Report Type:Expedited (15-DaCompany Report #B0107823A  
Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Unknown 1 Initial or Prolonged	1 WK	Decreased Appetite Depression Overdose		Zyban	PS	Glaxo Wellcome	ORAL

Date:05/21/01ISR Number: 3725789-6Report Type:Expedited (15-DaCompany Report #B0107842A  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300MG Per day	2 DAY	Deep Vein Thrombosis		Zyban Kliofem	PS C	Glaxo Wellcome	ORAL ORAL

Date:05/21/01ISR Number: 3726400-0Report Type:Expedited (15-DaCompany Report #A0138535A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG /		Diverticulum Oesophageal	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Initial or Prolonged TWICE PER DAY Disability / ORAL Required Intervention to Prevent Permanent Impairment/Damage 75 MG / TWICE  PER DAY /  ORAL	Dry Mouth  Dyskinesia  Gastric Polyps Gingival Pain Hypokalaemia Hyponatraemia  Pituitary Tumour Benign  Salivary Hypersecretion  Speech Disorder Tardive Dyskinesia	Professional  Company  Representative	Venlafaxine Hydrochloride (Formulation Unknown)	SS	ORAL
			Nicotine Gum Trazodone Clonazepam Dexamphetamine Sulphate Thyroxine Sodium	SS C C C C	

Date:05/21/01ISR Number: 3726417-6Report Type:Expedited (15-DaCompany Report #2001SUS0407  
Age:0 DY Gender:Male I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
TRANSPLACENTAL	TR	Complications Of Maternal Exposure To Therapeutic	Health Professional	Sustiva	PS	Dupont Pharmaceuticals Co	
TRANSPLACENTAL	TR	Drugs Cystic Fibrosis Meconium Ileus		Combivir (Lamivudine/Zidovudi ne)	SS		
TRANSPLACENTAL	TR			Norvir (Ritonavir)	SS		
TRANSPLACENTAL	TR			Crixivan (Indinavir)	SS		
TRANSPLACENTAL	TR			Bupropion			

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TR		(Amfebutamone)	SS
TRANSPLACENTAL	TR	Viramune (Nevirapine)	SS
TRANSPLACENTAL	TR	Epivir (Lamivudine)	SS
TRANSPLACENTAL	TR	Zerit (Stavudine)	SS
TRANSPLACENTAL	TR	Viracept (Nelfinavir)	SS

Date:05/21/01ISR Number: 3726825-3Report Type:Expedited (15-DaCompany Report #B0104518A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG/ PER	Asthenia	Foreign	Zyban	PS	Glaxo Wellcome Inc	
Hospitalization - DAY/		Cough	Health				
Initial or Prolonged Disability		Dizziness	Professional	Atenolol	C		
		Infection		Aspirin	C		
		Nausea		Dipyridamole	C		
		Neck Pain		Atorvastatin Calcium	C		
		Pain		Conjugated Estrogens	C		

Date:05/21/01ISR Number: 3726827-7Report Type:Expedited (15-DaCompany Report #B0099812A  
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG/ TWICE	Asthenia	Foreign	Zyban	PS	Glaxo Wellcome Inc	
PER DAY/		Blood Pressure Increased	Health				
		Dehydration	Professional	Quinapril	C		
		Diplopia		Trisequens	C		
		Dizziness		Ketoconazole	C		
		Dry Mouth		Ethanol	C		
		Dry Skin					
		Feeling Abnormal					
		Food Allergy					

Headache  
 Hypersensitivity  
 Malaise  
 Nasopharyngitis  
 Nausea  
 Pallor  
 Palpitations  
 Paraesthesia  
 Pharyngeal Oedema  
 Sneezing  
 Tongue Disorder  
 Tongue Oedema  
 Vertigo  
 Vision Blurred

Date:05/21/01ISR Number: 3726978-7Report Type:Expedited (15-DaCompany Report #B0092571A  
 Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 1 TABLET / AS Initial or Prolonged DIRECTED / Required ORAL Intervention to Prevent Permanent Impairment/Damage		Overdose Stress Suicide Attempt	Foreign Health Professional	Bupropion Hydrochloride      Paracetamol (Formulation Unknown)	PS	Glaxo Wellcome Inc	ORAL

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(Acetaminophen) SS  
 Ethanol (Formulation  
 Unknown) (Alcohol) SS

Date:05/21/01ISR Number: 3726979-9Report Type:Expedited (15-DaCompany Report #B0101422A  
 Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Therapeutic Response Unexpected	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER DAY / ORAL		Vitreous Haemorrhage		Amitriptyline	C		
				Isosorbide	C		
				Co-Codamol	C		
				Cerivastatin	C		
				Clopidogrel	C		
				Frusemide	C		
				Lacidipine	C		
				Fluoxetine	C		
				Aspirin	C		
				Ranitidine			
				Hydrochloride	C		
				Sennosides	C		
				Salbutamol Sulphate	C		
				Naproxen	C		
				Trimethoprim	C		
				Nicorandil	C		

Date:05/21/01ISR Number: 3726980-5Report Type:Expedited (15-DaCompany Report #B0099532A  
 Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Drug Interaction Intermittent Claudication	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER DAY / ORAL		Therapeutic Response Unexpected Vitreous Haemorrhage	Professional	Isosorbide Mononitrate	C		
				Co-Codamol	C		

Cerivastatin	C
Clopidogrel	C
Fruzemide	C
Lacidipine	C
Fluoxetine	C
Aspirin	C
Nitroglycerin	C
Beclomethasone	
Dipropion.	C
Ranitidine	
Hydrochloride	C
Amitriptyline	C
Sennosides	C
Salbutamol Sulphate	C
Naproxen	C
Trimethoprim	C
Nicorandil	C
Cephradine	C
Metronidazole	C

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Freedom Of Information (FOI) Report

Date:05/21/01ISR Number: 3726988-XReport Type:Expedited (15-DaCompany Report #B0107423A

Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Abdominal Distension	Foreign	Bupropion			
Intervention to		Depression		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
Prevent Permanent		Drug Withdrawal Syndrome					
TWICE PER DAY							
Impairment/Damage							
/ ORAL							

Date:05/22/01ISR Number: 3726477-2Report Type:Expedited (15-DaCompany Report #A0148065A

Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression		Wellbutrin	PS	Glaxo Wellcome	ORAL
Other		Amnesia					
150MG Twice		Asthenia		Birth Control Pills	C		
per day	9 DAY	Bite					
		Convulsion					
		Decreased Activity					
		Difficulty In Walking					
		Eyelid Disorder					
		Grand Mal Convulsion					
		Haematuria					
		Hypoaesthesia					
		Insomnia					
		Loss Of Consciousness					
		Mouth Haemorrhage					
		Muscle Twitching					
		Tongue Disorder					
		Tremor					
		Vision Blurred					

Date:05/22/01ISR Number: 3726481-4Report Type:Expedited (15-DaCompany Report #B0100055A

Age:27 YR Gender:Male I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Alanine Aminotransferase Increased		Zyban	PS	Glaxo Wellcome	ORAL

Arthralgia  
 Blood Pressure Decreased  
 Dermatitis  
 Heart Rate Decreased  
 Hypersensitivity  
 Infection  
 Joint Swelling  
 Malaise  
 Palpitations  
 Vomiting

Date:05/22/01ISR Number: 3726482-6Report Type:Expedited (15-DaCompany Report #B0100789A  
 Age:29 YR Gender:Female I/FU:F

Outcome	PT
	Anxiety Arthralgia Cyanosis Hyperhidrosis Musculoskeletal Stiffness Nausea

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Dose	Duration	Peripheral Coldness Tremor	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day				Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL

Date:05/22/01ISR Number: 3726485-1Report Type:Expedited (15-DaCompany Report #B0107427A  
Age:55 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		55 DAY	Pneumothorax		Zyban	PS	Glaxo Wellcome	ORAL
2PUFF Twice per day					Salmeterol	C	Glaxo Wellcome	
2PUFF Twice per day					Ipratropium	C	Glaxo Wellcome	
2PUFF Twice per day					Terbutaline	C		

Date:05/22/01ISR Number: 3726488-7Report Type:Expedited (15-DaCompany Report #B0108509A  
Age:57 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Per day		62 DAY	Diabetes Mellitus		Zyban	PS	Glaxo Wellcome	ORAL
UNKNOWN	150MG		Non-Insulin-Dependent Unknown Malaise		Ranitidine	C	Glaxo Wellcome	

Date:05/22/01ISR Number: 3726489-9Report Type:Expedited (15-DaCompany Report #B0108517A  
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death		Zyban	PS	Glaxo Wellcome	

Date:05/22/01ISR Number: 3726865-4Report Type:Expedited (15-DaCompany Report #B0107381A  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Ecchymosis Excoriation Fall Overdose Periorbital Haematoma Subdural Haematoma	Foreign Health Professional	Bupropion Hydrochloride Fluoxetine Ethanol	PS C C	Glaxo Wellcome Inc	

Date:05/22/01ISR Number: 3726866-6Report Type:Expedited (15-DaCompany Report #B0107247A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG		Hemiplegia Paraesthesia	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

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Freedom Of Information (FOI) Report

Date:05/22/01ISR Number: 3727624-9Report Type:Direct  
Age:32 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 6 CAPS PER Initial or Prolonged DAY	3	WK	Cardiac Arrest	Metabolife	PS		
BID	3	WK	Grand Mal Convulsion Metabolic Acidosis Respiratory Arrest Ventricular Fibrillation Vomiting	Wellbutrin	SS		

Date:05/23/01ISR Number: 3727296-3Report Type:Expedited (15-DaCompany Report #A0147746A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 75MG Per day	1	WK	Condition Aggravated Feeling Abnormal Petit Mal Epilepsy	Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:05/23/01ISR Number: 3727318-XReport Type:Expedited (15-DaCompany Report #B0108029A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Atrial Fibrillation Dyspnoea	Zyban	PS	Glaxo Wellcome	

Date:05/23/01ISR Number: 3727319-1Report Type:Expedited (15-DaCompany Report #B0108034A  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG Per day			Postmenopausal	Zyban	PS	Glaxo Wellcome	ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG		Haemorrhage		Ranitidine	C	Glaxo Wellcome	ORAL
Variable dose				Trisequens	C		
Date:05/23/01ISR Number: 3727320-8Report Type:Expedited (15-DaCompany Report #B0108210A Age:37 YR Gender:Male I/FU:I							
Disability 150MG Twice per day	18 DAY	Fatigue Headache Insomnia Keratitis		Zyban	PS	Glaxo Wellcome	ORAL
Date:05/23/01ISR Number: 3727321-XReport Type:Expedited (15-DaCompany Report #B0108213A Age:50 YR Gender:Female I/FU:I							
Other 150MG Twice per day	9 DAY	Diarrhoea Haemorrhagic		Zyban  Levonorgestrel	PS  C	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/23/01ISR Number: 3727322-1Report Type:Expedited (15-DaCompany Report #B0108220A

Age:50 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged						
	Pulmonary Oedema		Zyban	PS	Glaxo Wellcome	

Date:05/23/01ISR Number: 3727323-3Report Type:Expedited (15-DaCompany Report #B0108504A

Age:71 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day 5 DAY 10MG Three times per day 40MG Per day 1U Four times per day SUBLINGUAL	Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
			Nifedipine	C		ORAL
			Frusemide	C		ORAL
			Potassium Chloride	C		ORAL
			Glyceryl Trinitrate	C	Glaxo Wellcome	

Date:05/23/01ISR Number: 3727324-5Report Type:Expedited (15-DaCompany Report #B0108505A

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day 53 DAY	Dyspnoea Exertional Left Ventricular Failure		Zyban	PS	Glaxo Wellcome	ORAL
			Lisinopril	C		ORAL
			Diltiazem	C		ORAL
			Aspirin	C		ORAL

UNKNOWN Cerivastatin C  
 UNKNOWN 20MG Twice Isosorbide C  
 per day

Date:05/23/01ISR Number: 3727325-7Report Type:Expedited (15-DaCompany Report #B0108747A  
 Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous		Zyban	PS	Glaxo Wellcome	
UNKNOWN	300MG per day 3	WK					
		Pregnancy					

Date:05/23/01ISR Number: 3727868-6Report Type:Expedited (15-DaCompany Report #B0100682A  
 Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Agitation	Foreign	Zyban	PS	Glaxo Wellcome Inc	
150 MG /							
Initial or Prolonged		Blood Glucose Decreased	Health				
TWICE PER DAY							
/		Circulatory Collapse	Professional				
		Dizziness		Coffee	C		
		Headache		Naproxen	C		
		Hypoaesthesia					
		Photophobia					
		Vision Blurred					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/23/01ISR Number: 3727912-6Report Type:Expedited (15-DaCompany Report #B0107842A  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Deep Vein Thrombosis	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
300 MG (PER DAY) ORAL				Kliefem	C		

Date:05/23/01ISR Number: 3727915-1Report Type:Expedited (15-DaCompany Report #B0107510A  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG (PER DAY) ORAL		Asthma Back Pain	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Aminophylline	C		
				Prednisolone	C		
				Diclofenac	C		

Date:05/23/01ISR Number: 3727918-7Report Type:Expedited (15-DaCompany Report #B0107658A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG (TWICE PER DAY) ORAL Impairment/Damage		Agitation Disturbance In Attention Feeling Abnormal Speech Disorder Tinnitus Visual Disturbance	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL



Date:05/23/01ISR Number: 3727924-2Report Type:Expedited (15-DaCompany Report #B0107823A  
Age:20 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG ORAL 1 WK	Decreased Appetite Depression Flat Affect Overdose	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/23/01ISR Number: 3728001-7Report Type:Expedited (15-DaCompany Report #A0139670A  
Age:74 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG (PER Initial or Prolonged DAY) ORAL 2 MON	Cardiac Failure Congestive Convulsion Loss Of Consciousness	Consumer	Wellbutrin Sr  Ceftriaxone Sodium Famotidine Potassium Chloride Captopril Semisodium Valproate Buspirone Hydrochloride Nicotine Quetiapine Fumarate Frusemide Digoxin Salbutamol Sulphate	PS  C C C C C C C C C C C	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Spironolactone C  
 Aspirin C  
 Meclozine  
 Hydrochloride C  
 Hypremellose C  
 Ipratropium Bromide C

Date:05/24/01ISR Number: 3727940-0Report Type:Expedited (15-DaCompany Report #A0093694A  
 Age:46 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Per day 4 MON	Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged Disability	Decreased Activity Depression Grand Mal Convulsion Pain Spinal Fracture		Concurrent Medications	C		

Date:05/24/01ISR Number: 3727951-5Report Type:Expedited (15-DaCompany Report #B0100520A  
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration	Circulatory Collapse Dizziness Hypoaesthesia Musculoskeletal Stiffness Nausea Photophobia Retching		Zyban	PS	Glaxo Wellcome	

Date:05/24/01ISR Number: 3728480-5Report Type:Expedited (15-DaCompany Report #A0148065A  
 Age:20 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG /	Aggression	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

TWICE PER DAY

Amnesia

Difficulty In Walking

/ ORAL

Grand Mal Convulsion

Oral Contraceptive C

Haematuria

Hypoaesthesia

Insomnia

Loss Of Consciousness

Mouth Haemorrhage

Muscle Twitching

Tremor

Vision Blurred

Date:05/24/01ISR Number: 3728699-3Report Type:Expedited (15-DaCompany Report #B0100055A

Age:27 YR Gender:Male I/FU:F

Outcome

PT

Arthralgia

Aspartate

Aminotransferase

Increased

Blood Pressure Decreased

Dermatitis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY / ORAL		Heart Rate Decreased Hypersensitivity Infection Joint Swelling Malaise Palpitations Vomiting	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/24/01ISR Number: 3728700-7Report Type:Expedited (15-DaCompany Report #B0100789A  
Age:29 YR Gender:Female I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Anxiety Arthralgia Hyperhidrosis Musculoskeletal Stiffness Nausea Peripheral Coldness Skin Discolouration Tremor	Foreign Study Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:05/24/01ISR Number: 3728738-XReport Type:Expedited (15-DaCompany Report #B0108517A  
Age:54 YR Gender:Male I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG		Sudden Death	Foreign	Zyban	PS	Glaxo Wellcome Inc	

Date:05/24/01ISR Number: 3728739-1Report Type:Expedited (15-DaCompany Report #B0108509A  
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG / PER DAY / ORAL	Diabetes Mellitus Non-Insulin-Dependent Malaise	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Ranitidine Hydrochloride	C		

Date:05/24/01ISR Number: 3728740-8Report Type:Expedited (15-DaCompany Report #B0107427A  
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY / ORAL		Pneumothorax	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Salmeterol Xinafoate Ipratropium Bromide Terbutaline	C C C		

Date:05/25/01ISR Number: 3728381-2Report Type:Expedited (15-DaCompany Report #B0107824A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 10 DAY Initial or Prolonged UNKNOWN		Hypothermia Mood Altered Overdose Social Avoidant Behaviour Stress		Zyban Aspirin	PS SS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/25/01ISR Number: 3728819-0Report Type:Expedited (15-DaCompany Report #A0147746A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
75 MG (PER DAY)		Feeling Abnormal					

Date:05/25/01ISR Number: 3729339-XReport Type:Expedited (15-DaCompany Report #B0108220A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Pulmonary Oedema	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
150 MG			Professional				

Date:05/25/01ISR Number: 3729340-6Report Type:Expedited (15-DaCompany Report #B0108213A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required Intervention to Prevent Permanent Impairment/Damage		Diarrhoea Haemorrhagic	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE PER DAY/ ORAL				Levonorgestrel	C		

Date:05/25/01ISR Number: 3729341-8Report Type:Expedited (15-DaCompany Report #B0108210A  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Fatigue Headache	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							

Insomnia

PER DAY/ ORAL

Keratitis

Date:05/25/01ISR Number: 3729390-XReport Type:Expedited (15-DaCompany Report #B0108034A

Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Postmenopausal	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
300 MG PER DAY ORAL		Haemorrhage					
Prevent Permanent Impairment/Damage				Ranitidine Hydrochloride	C		
				Trisequens	C		

Date:05/25/01ISR Number: 3729398-4Report Type:Expedited (15-DaCompany Report #B0108504A

Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Angina Pectoris	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE PER DAY ORAL							
Prevent Permanent Impairment/Damage				Nifedipine	C		
				Frusemide	C		
				Potassium Chloride	C		
				Nitroglycerin	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/25/01ISR Number: 3729399-6Report Type:Expedited (15-DaCompany Report #B0108029A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Atrial Fibrillation	Foreign	Zyban	PS	Glaxo Wellcome Inc	
Other	150 MG	Dyspnoea	Consumer				
UNKNOWN							
UNKNOWN							

Date:05/25/01ISR Number: 3729400-XReport Type:Expedited (15-DaCompany Report #B0108747A

Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abortion Spontaneous	Foreign	Zyban	PS	Glaxo Wellcome Inc	
Other		3 WK					
UNKNOWN		Complications Of Maternal	Health				
		Exposure To Therapeutic	Professional				
		Drugs					
		Pregnancy					

Date:05/25/01ISR Number: 3729401-1Report Type:Expedited (15-DaCompany Report #B0108505A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Left Ventricular Failure	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Required							
150 MG TWICE							
Intervention to							
PER DAY ORAL							
Prevent Permanent				Lisinopril	C		
Impairment/Damage				Diltiazem	C		
				Aspirin	C		
				Cerivastatin	C		
				Isosorbide	C		

Date:05/25/01ISR Number: 3733469-6Report Type:Periodic

Age:52 YR Gender:Female I/FU:I

Company Report #HQ8388514MAR2001



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
40 MG 1 X PER	1 DAY, ORAL	Asthenia	Consumer	Protonix	PS		ORAL
		Dizziness					
		Drug Interaction		Estratest (Estorgens Esterified/Methyltestosterone)	SS		ORAL
1 TABLET 1X	PER 1 DAY, ORAL						
15 MG 1X PER	1 DAY, ORAL			Univasc (Moexipril Hydrochloride)	SS		ORAL
150 MG 2X PER	1 DAY, ORAL			Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL

Date:05/29/01ISR Number: 3729842-2Report Type:Expedited (15-DaCompany Report #S01-USA-00846-01  
Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anxiety Circulatory Collapse Drug Level Above Therapeutic Parkinson'S Disease	Consumer	Celexa	PS	Forest Laboratories Inc	
INTRAVENOUS	150 MG	BID IV		Wellbutrin (Amfebutamone Hydrochloride)	SS		
				Depakote (Valproate)			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Semisodium) SS  
 Paxil (Paroxetine  
 Hydrochloride) SS

Date:05/29/01ISR Number: 3730019-5Report Type:Expedited (15-DaCompany Report #A0093694A  
 Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ PER Initial or Prolonged DAY/ ORAL Disability		Convulsion Depression Grand Mal Convulsion Pain Spinal Compression Fracture	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:05/29/01ISR Number: 3730364-3Report Type:Expedited (15-DaCompany Report #B0100520A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG		Circulatory Collapse Dizziness Hypoaesthesia Musculoskeletal Stiffness Nausea Photophobia Retching	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:05/30/01ISR Number: 3729804-5Report Type:Expedited (15-DaCompany Report #A0148248A  
 Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Twice Hospitalization - per day 11 DAY		Amnesia Cardio-Respiratory Arrest		Wellbutrin	PS	Glaxo Wellcome	ORAL

Initial or Prolonged	Coma	Flomax	SS	ORAL
.4MG Per day	1 DAY			
2 YR	Convulsion	Viagra	SS	
YR	Drug Interaction	Lipitor	C	
YR	Fall	Lorazepam	C	
YR	Head Injury	Baby Aspirin	C	
8 DAY		Msm	C	
		Co Q10	C	
		Multivitamins	C	
		Fish Oil	C	
		Pentatonic Acid	C	

Date:05/30/01ISR Number: 3729814-8Report Type:Expedited (15-DaCompany Report #B0103784A  
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150MG Twice per day	21 DAY	Alcohol Intolerance		Zyban	PS	Glaxo Wellcome	ORAL
		Haematuria					
		Rhabdomyolysis		Alcohol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/30/01ISR Number: 3729819-7Report Type:Expedited (15-DaCompany Report #B0106830A  
Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 8 DAY Initial or Prolonged		Abdominal Discomfort Diarrhoea Lung Consolidation Nephritis Red Blood Cell Sedimentation Rate Increased Vomiting White Blood Cell Count Increased		Zyban	PS	Glaxo Wellcome	

Date:05/30/01ISR Number: 3729822-7Report Type:Expedited (15-DaCompany Report #B0108202A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Zyban	PS	Glaxo Wellcome	ORAL

Date:05/30/01ISR Number: 3729825-2Report Type:Expedited (15-DaCompany Report #B0108364A  
Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Pain Upper Arthralgia Blister Chest Discomfort Constipation Cough Dizziness Dyspnoea Face Oedema Feeling Cold Haemoptysis Hypoventilation Oliguria Pharyngeal Oedema		Zyban	PS	Glaxo Wellcome	

Urticaria

Date:05/30/01ISR Number: 3729826-4Report Type:Expedited (15-DaCompany Report #B0108503A  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Haemorrhage		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
		Proctitis Ulcerative					
per day	55	DAY					
		Purulence		Mefanamic Acid	C		ORAL
15	WK						

Date:05/30/01ISR Number: 3730651-9Report Type:Expedited (15-DaCompany Report #B0107824A  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Hypothermia	Foreign	Bupropion			
Initial or Prolonged		Mood Altered	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL	10	DAY					
		Overdose	Professional	Aspirin (Aspirin)	SS		
		Stress					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/30/01  
 Age:44 YR  
 Gender:Male  
 I/FU:I

Report Type:Direct  
 Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1 MON	Abdominal Pain		Wellbutrin	PS		
		Back Pain					
		Cardio-Respiratory Arrest					
		Electrocardiogram Qt Prolonged					
		Sinus Tachycardia					
		Vomiting					

Date:05/30/01  
 Age:39 YR  
 Gender:Female  
 I/FU:I

Report Type:Expedited (15-Da  
 Company Report #A0148393A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - SEE TEXT / Initial or Prolonged	1 MON	Asthenia	Literature	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
ORAL		Heart Rate Increased	Health				
		Pyrexia	Professional	Valproic Acid	C		
		Supraventricular Tachycardia		Mirtazapine	C		

Date:05/31/01  
 Age:28 YR  
 Gender:Female  
 I/FU:F

Report Type:Expedited (15-Da  
 Company Report #A0097619A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	23 DAY	Angioneurotic Oedema		Zyban	PS	Glaxo Wellcome	ORAL
200MG As required	3 DAY	Arthralgia					
		Heart Rate Increased		Ibuprofen	C		ORAL
		Malaise					
		Oedema Peripheral					
		Pruritus					
		Serum Sickness					

Skin Discolouration  
Urticaria

Date:05/31/01ISR Number: 3731140-8Report Type:Expedited (15-DaCompany Report #A0103746A  
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	16 DAY	Analgesic Drug Level Angioneurotic Oedema		Zyban	PS	Glaxo Wellcome	ORAL
		Arthralgia Asthma Chills Dyspepsia Face Oedema Lymphadenopathy Myalgia Odynophagia Oedema Oedema Peripheral Periorbital Oedema Pruritus Pyrexia Rash Erythematous Serum Sickness Synovitis Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/01/01ISR Number: 3731735-1Report Type:Expedited (15-DaCompany Report #B0108772A

Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Interaction		Bupropion Hcl	PS	Glaxo Wellcome	ORAL
150MG Per day				Aspirin	C		ORAL
1UNIT Per day				Atenolol	C		ORAL
RESPIRATORY				Beclomethasone	C	Glaxo Wellcome	
(INHALATION)	200MCG Twice						
per day				Co-Amilofruse	C		ORAL
1UNIT Per day				Imdur	C		ORAL
60MG Per day				Lactulose	C		ORAL
15ML Twice				Lisinopril	C		ORAL
per day				Zydol	C		ORAL
20MG Twice				Fluoxetine	C		ORAL
per day				Cetirizine	C	Glaxo Wellcome	ORAL
20MG Per day				Betnovate	C	Glaxo Wellcome	
10MG Per day							
TOPICAL	.1PCT Twice						
per day							

Date:06/01/01ISR Number: 3731736-3Report Type:Expedited (15-DaCompany Report #B0108775A

Age:38 YR Gender:Male I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Bupropion Hcl	PS	Glaxo Wellcome	ORAL
Hospitalization - 5MG Three Initial or Prolonged times per day		Hallucination  Overdose  Tremor		Nifedipine	C		ORAL

Date:06/01/01ISR Number: 3731738-7Report Type:Expedited (15-DaCompany Report #B0109122A  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Per day 3 DAY		Alanine Aminotransferase Increased Alpha 1 Foetoprotein Increased Aspartate Aminotransferase Increased Asthenia Blood Bilirubin Increased Gamma-Glutamyltransferase Increased Laboratory Test Abnormal Malaise		Zyntabac	PS	Glaxo Wellcome	ORAL

Date:06/01/01ISR Number: 3731739-9Report Type:Expedited (15-DaCompany Report #B0109126A  
Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Zyban Indoramin Beclomethasone Dipropionate Diazepam	PS C C C	Glaxo Wellcome Glaxo Wellcome Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Salbutamol C Glaxo Wellcome

Date:06/01/01ISR Number: 3731741-7Report Type:Expedited (15-DaCompany Report #B0109574A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	11 DAY	Agitation		Zyban	PS	Glaxo Wellcome	ORAL
1.25MG Per		Asthenia		Prempak-C	C		ORAL
day		Dizziness					
300MCG Weekly		Flight Of Ideas		Thyroxine	C	Glaxo Wellcome	ORAL
		Insomnia					
		Memory Impairment					
		Palpitations					
		Speech Disorder					
		Tremor					
		Vertigo					

Date:06/01/01ISR Number: 3731742-9Report Type:Expedited (15-DaCompany Report #B0109579A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Twice	Colitis Ulcerative		Zyban	PS	Glaxo Wellcome	ORAL
per day	62 DAY	Diarrhoea Haemorrhagic					

Date:06/01/01ISR Number: 3731743-0Report Type:Expedited (15-DaCompany Report #B0109588A  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Twice	Supraventricular		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -	per day	Tachycardia					

Initial or Prolonged  
Other

Aspirin  
Ketoprofen

C  
C

Date:06/01/01ISR Number: 3731744-2Report Type:Expedited (15-DaCompany Report #B0109706A  
Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Haemorrhagic Stroke Status Epilepticus		Zyban	PS	Glaxo Wellcome	

Date:06/01/01ISR Number: 3732214-8Report Type:Expedited (15-DaCompany Report #B0103784A  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150 MG/TWICE PER DAY/ ORAL		Alcohol Intolerance Haematuria Rhabdomyolysis	Foreign Health Professional	Zyban  Ethanol	PS  C	Glaxo Wellcome Inc	ORAL

Date:06/01/01ISR Number: 3732215-XReport Type:Expedited (15-DaCompany Report #B0108503A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150 MG/ TWICE		Proctitis Haemorrhagic Proctitis Ulcerative	Foreign	Zyban Tablet- Zyban (Bupropion Hydrochloride)	PS	Glaxo Wellcome Inc	ORAL

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Freedom Of Information (FOI) Report

PER DAY/ ORAL

Mefenamic Acid C

Date:06/01/01ISR Number: 3732222-7Report Type:Expedited (15-DaCompany Report #B0108364A  
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Pain Upper Arthralgia Blister Chest Discomfort Constipation Cough Dizziness Dyspnoea Face Oedema Feeling Cold Haemoptysis Hypoventilation Oliguria Pharyngeal Oedema Urticaria	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	

Date:06/01/01ISR Number: 3732228-8Report Type:Expedited (15-DaCompany Report #B0106830A  
Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Discomfort Blood Test Abnormal Chromaturia Diarrhoea Ill-Defined Disorder Inflammation Lung Consolidation Malaise Nephritis Vomiting White Blood Cell Count Increased	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:06/01/01ISR Number: 3732276-8Report Type:Expedited (15-DaCompany Report #B0108202A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
ORAL			Health Professional				

Date:06/01/01ISR Number: 3732277-XReport Type:Expedited (15-DaCompany Report #A0148248A  
Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Coma	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
150 MG / Hospitalization - TWICE PER DAY		Convulsion					
Initial or Prolonged / ORAL		Drug Interaction					
		Fall		Tamsulosin Hcl			
.4 MG / PER DAY / ORAL		Head Injury		Capsule	SS		ORAL
		Tachycardia					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sildenafil Citrate  
 (Formulation  
 Unknown) SS  
 Atorvastatin Calcium C  
 Lorazepam C  
 Aspirin C  
 Dimethyl Sulfone C  
 Ubidecarenone C  
 Multivitamin C  
 Fish Oil C  
 Pentatonic Acid C

Date:06/01/01ISR Number: 3738398-XReport Type:Expedited (15-DaCompany Report #A0149544A  
 Age:59 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - PER DAY/ORAL Initial or Prolonged	Amnesia Brain Damage Concussion Convulsion Fall Head Injury Loss Of Consciousness Spinal Fracture	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL

Date:06/04/01ISR Number: 3732293-8Report Type:Expedited (15-DaCompany Report #B0100818A  
 Age:71 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death 150MG Per day 10ML Three times per day	Nausea Vomiting		Zyban Magnesium Trisilicate	PS C	Glaxo Wellcome	ORAL ORAL

Date:06/04/01ISR Number: 3732296-3Report Type:Expedited (15-DaCompany Report #B0103185A  
Age:48 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Abdominal Distension Abdominal Tenderness Chest Pain Dry Mouth Fatigue Lethargy Malaise Red Blood Cell Count Decreased Tinnitus White Blood Cell Count Decreased		Zyban Anti-Inflammatory Agents	PS C	Glaxo Wellcome	ORAL

Date:06/04/01ISR Number: 3732297-5Report Type:Expedited (15-DaCompany Report #B0103385A  
Age:69 YR Gender:Female I/FU:F

Outcome	PT
	Anorexia Cough

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	12 DAY	Dizziness Lower Respiratory Tract Infection Nausea Vomiting		Zyban	PS	Glaxo Wellcome	ORAL
75MG Unknown				Aspirin	C		ORAL
20MG Unknown				Simvastatin	C		ORAL

Date:06/04/01ISR Number: 3732298-7Report Type:Expedited (15-DaCompany Report #B0108028A  
Age:62 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged per day	150MG Twice	13 DAY	Dizziness Headache Palpitations Tremor		Zyban	PS	Glaxo Wellcome	ORAL
					Monoplus	C		

Date:06/04/01ISR Number: 3732302-6Report Type:Expedited (15-DaCompany Report #B0109127A  
Age:37 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other per day	150MG Twice	15 DAY	Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs Pregnancy		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/04/01ISR Number: 3732303-8Report Type:Expedited (15-DaCompany Report #B0109589A  
Age:39 YR Gender:Female I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day	Amnesia		Zyban	PS	Glaxo Wellcome	
		Clonic Convulsion		Lorazepam	C		
		Convulsion		Irbesartan	C		
		Dizziness					
		Feeling Abnormal					
		Loss Of Consciousness					
		Malaise					
		Tremor					

Date:06/04/01ISR Number: 3732304-XReport Type:Expedited (15-DaCompany Report #B0109704A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day 36 DAY	Anaemia		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged		Cerebral Ischaemia					
		Epilepsy					
		Lipase Increased					
		Liver Function Test Abnormal					
		Red Blood Cell Sedimentation Rate Increased					
		Thrombocytopenia					
		White Blood Cell Count Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/04/01ISR Number: 3732306-3Report Type:Expedited (15-DaCompany Report #D0017224A

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Intracranial Haemangioma		Zyban	PS	Glaxo Wellcome	ORAL
UNKNOWN		Movement Disorder		Ass	C		
UNKNOWN		Nervous System Disorder		Amaryl	C		

Date:06/04/01ISR Number: 3733442-8Report Type:Periodic Company Report #A108581

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150.00 MG		Anxiety Drug Interaction	Health Professional	Geodon	PS	Pfizer Central Research	ORAL
TOTAL:BID:ORA							
L				Wellbutrin	SS		ORAL
250.00 MG							
TOTAL:BID:ORA							
L							

Date:06/05/01ISR Number: 3733204-1Report Type:Expedited (15-DaCompany Report #B0107381A

Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death UNKNOWN		Alcohol Poisoning		Zyban	PS	Glaxo Wellcome	
		Balance Disorder		Fluoxetine	C		
		Drug Level Above Therapeutic		Alcohol	C		

Ecchymosis  
Excoriation  
Fall  
Head Injury  
Hypoaesthesia  
Overdose  
Periorbital Haematoma  
Subdural Haematoma

Date:06/05/01ISR Number: 3733213-2Report Type:Expedited (15-DaCompany Report #B0109966A

Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Zyban	PS	Glaxo Wellcome	ORAL
3 DAY		Hyperventilation					

Date:06/05/01ISR Number: 3733214-4Report Type:Expedited (15-DaCompany Report #B0109968A

Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cough		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Heart Rate Increased Paraesthesia					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/05/01ISR Number: 3733953-5Report Type:Expedited (15-DaCompany Report #B0109579A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required	150 MG TWICE	Colitis Ulcerative	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Intervention to Prevent Permanent Impairment/Damage							

Date:06/05/01ISR Number: 3733956-0Report Type:Expedited (15-DaCompany Report #B0108775A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG ORAL	Cardiac Arrest	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - Initial or Prolonged		Hallucination Tremor		Nifedipine	C		

Date:06/05/01ISR Number: 3733960-2Report Type:Expedited (15-DaCompany Report #B0109126A  
Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG ORAL	Death	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Indoramin	C		
				Beclomethasone			
				Dipropion	C		
				Diazepam	C		
				Salbutamol Sulphate	C		

Date:06/05/01ISR Number: 3733964-XReport Type:Expedited (15-DaCompany Report #B0109574A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Disability	Agitation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150MG ORAL	Asthenia		Prempak-C	C		
	Insomnia		Thyroxine Sodium	C		
	Memory Impairment					
	Palpitations					
	Speech Disorder					
	Syncope					
	Thinking Abnormal					
	Tremor					
	Vertigo					
	Visual Disturbance					

Date:06/05/01ISR Number: 3733966-3Report Type:Expedited (15-DaCompany Report #B0109122A  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alanine Aminotransferase	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Increased	Health				
DAY ORAL		Aspartate	Professional				
		Aminotransferase					
		Increased					
		Asthenia					
		Blood Bilirubin Increased					
		Gamma-Glutamyltransferase					
		Increased					
		Malaise					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/05/01ISR Number: 3733967-5Report Type:Expedited (15-DaCompany Report #B0109588A  
Age:60 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150 MG TWICE	Arrhythmia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - PER DAY ORAL	Supraventricular	Health				
Initial or Prolonged Required	Tachycardia	Professional	Aspirin Ketoprofen	C C		
Intervention to Prevent Permanent Impairment/Damage						

Date:06/05/01ISR Number: 3734139-0Report Type:Expedited (15-DaCompany Report #2012743  
Age:47 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Grand Mal Convulsion Impaired Work Ability	Foreign Health	Dhc Plus	PS	Purdue Frederick Cp.	
150 MG		Professional Company	Zyban (Bupropion Hydrochloride)	SS		ORAL
1-2/DAY PO		Representative				
			Co-Amoxiclav (Amoxicillin/Clavula nic Acid	C		
			Salbutamol Sulphate, Budesonide	C		

Date:06/05/01ISR Number: 3734565-XReport Type:Expedited (15-DaCompany Report #A0097619A  
Age:28 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose 150 MG (TWICE	Difficulty In Walking Heart Rate Increased	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY) ORAL	Malaise	Professional				

Oedema Peripheral  
Pain In Extremity  
Serum Sickness  
Skin Discolouration

Ibuprofen

C

Date:06/05/01ISR Number: 3734567-3Report Type:Expedited (15-DaCompany Report #A0103746A  
Age:24 YR Gender:Male I/FU:F

Outcome PT  
Hospitalization - Angioneurotic Oedema  
Initial or Prolonged Arthralgia  
Asthma  
Chills  
Dyspepsia  
Erythema  
Face Oedema  
Lymphadenopathy  
Movement Disorder  
Myalgia  
Odynophagia  
Oedema Peripheral  
Periorbital Oedema  
Pruritus  
Pyrexia  
Serum Sickness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Synovitis Urticaria				
Dose	Duration		Report Source	Product	Role	Manufacturer
			Foreign Literature	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc
150 MG (TWICE PER DAY) ORAL	16 DAY		Health Professional			ORAL

Date:06/05/01ISR Number: 3734657-5Report Type:Expedited (15-DaCompany Report #B0109706A  
Age:31 YR Gender:Female I/FU:I

		PT				
Outcome	Duration		Report Source	Product	Role	Manufacturer
Dose		Haemorrhagic Stroke	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc
Hospitalization - Initial or Prolonged 150 MG		Status Epilepticus	Professional Company Representative			

Date:06/05/01ISR Number: 3734829-XReport Type:Expedited (15-DaCompany Report #10819928  
Age: Gender:Female I/FU:F

		PT				
Outcome	Duration		Report Source	Product	Role	Manufacturer
Dose		Hypersensitivity	Health Professional	Paraplatin	PS	Bristol Myers Squibb Co Pharmaceutical Research Institute
Hospitalization - Initial or Prolonged Other INTRAVENOUS	640	Loss Of Consciousness Petit Mal Epilepsy				

MILLIGRAM,

1/1 CYCLE IV

Wellbutrin (Bupropion Hcl)	SS
Decadron	C
Anzemet	C
Gemzar	C



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Interaction	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL				Aspirin	C		
				Atenolol	C		
				Beclomethasone			
				Dipropion.	C		
				Co-Amilofruse	C		
				Isosorbide			
				Mononitrate	C		
				Lactulose	C		
				Lisinopril	C		
				Tramadol			
				Hydrochloride	C		
				Fluoxetine	C		
				Cetirizine			
				Hydrochloride	C		
				Betamethasone			
				Valerate	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/05/01ISR Number: 3735639-XReport Type:Expedited (15-DaCompany Report #2012618

Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Atherosclerosis	Health	Oxycontin	PS	Purdue Pharma Lp	ORAL
MG PO							
Hospitalization -		Blood Creatine	Professional	Bupropion	SS		
Initial or Prolonged		Phosphokinase Increased	Other	Carisoprodol	SS		
		Blood Glucose Increased		Amitriptyline	SS		
		Bronchopneumonia		Benzodiazepine	SS		
		Cardiomegaly		Ibuprofen	SS		
		Drug Abuser		Thorazine			
		Haemodialysis		(Chlorpromazine Hcl)	C		
		Hepatitis C		Plendil	C		
		Hypotension		Risperdal	C		
		Hypothermia					
		Overdose					
		Peritonitis					
		Renal Failure					

Date:06/06/01ISR Number: 3734042-6Report Type:Expedited (15-DaCompany Report #A0149363A

Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Blood Pressure Increased		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice							
Initial or Prolonged		Disturbance In Attention					
per day							
		Drug Ineffective		Wellbutrin	SS	Glaxo Wellcome	
		Electroencephalogram		Tylenol	SS		
		Abnormal		Imodium	SS		
		Grand Mal Convulsion		Paxil	C	Glaxo Wellcome	ORAL
		Hallucination		Tetracycline	C		
		Intentional Misuse		Retin-A	C		
		Irritability					
		Memory Impairment					
		Sedation					
		Tachycardia					

TOPICAL

Date:06/06/01ISR Number: 3734045-1Report Type:Expedited (15-DaCompany Report #A0149803A  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Zyban	PS	Glaxo Wellcome	ORAL
7.5G See text		Catatonia		Amphetamine	C		
		Convulsion		Cannabis	C		
		Intentional Misuse					
		Sedation					
		Tachycardia					

Date:06/06/01ISR Number: 3734046-3Report Type:Expedited (15-DaCompany Report #A0149863A  
Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dry Mouth		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Dry Throat					
per day		Emphysema					
		Laryngitis					
		Speech Disorder					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/06/01ISR Number: 3734050-5Report Type:Expedited (15-DaCompany Report #A0150080A  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day		Decreased Activity		Zyban	PS	Glaxo Wellcome	ORAL
		Feeling Cold Hypoaesthesia					

Date:06/06/01ISR Number: 3734059-1Report Type:Expedited (15-DaCompany Report #B0109581A  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day		Anxiety		Zyban	PS	Glaxo Wellcome	
		Convulsion					
150MG Twice per day		Drug Effect Decreased		Lithium Carbonate	C	Glaxo Wellcome	
		Dysgeusia					
50MCG Twice per day		Joint Dislocation		Thyroxine	C	Glaxo Wellcome	
				Nortriptyline Hydrochloride	C		
10MG Twice per day							

Date:06/06/01ISR Number: 3734062-1Report Type:Expedited (15-DaCompany Report #B0110167A  
Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Twice per day	21 DAY	Decreased Activity		Zyban	PS	Glaxo Wellcome	ORAL
		Staphylococcal Scalded Skin Syndrome					

Swelling

Date:06/06/01ISR Number: 3734063-3Report Type:Expedited (15-DaCompany Report #B0110168A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day	Psoriasis		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/06/01ISR Number: 3734064-5Report Type:Expedited (15-DaCompany Report #B0110170A  
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	48 DAY	Retroperitoneal Haemorrhage		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/06/01ISR Number: 3734065-7Report Type:Expedited (15-DaCompany Report #B0110171A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	8 DAY	Abdominal Distension Oedema		Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/06/01ISR Number: 3735445-6Report Type:Expedited (15-DaCompany Report #B0109704A

Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anaemia	Foreign	Bupropion			
Hospitalization - 150 MG/PER Initial or Prolonged DAY/ORAL		Aspartate		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Aminotransferase Increased Blood Alkaline Phosphatase Increased Blood Bilirubin Increased Blood Creatine Phosphokinase Increased Cerebral Ischaemia Epilepsy Lipase Increased Liver Function Test Abnormal Red Blood Cell Sedimentation Rate Increased Thrombocytopenia White Blood Cell Count Increased					

Date:06/06/01ISR Number: 3735446-8Report Type:Expedited (15-DaCompany Report #B0109127A

Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG/TWICE Prevent Permanent PER DAY/ORAL Impairment/Damage		Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs Pregnancy	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:06/06/01ISR Number: 3735447-XReport Type:Expedited (15-DaCompany Report #B0103185A

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Abdominal Distension Chest Pain	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ORAL		Dry Mouth Fatigue Lethargy Malaise Red Blood Cell Count Decreased Tinnitus White Blood Cell Count Decreased	Professional Company Representative	Anti-Inflammatory	C		

Date:06/06/01ISR Number: 3735448-1Report Type:Expedited (15-DaCompany Report #B0108028A  
Age:62 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Balance Disorder Dizziness Headache Palpitations

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Foreign Health Professional	Bupropion Hydrochloride Monoplus	PS C	Glaxo Wellcome Inc	ORAL

Date:06/06/01ISR Number: 3735449-3Report Type:Expedited (15-DaCompany Report #B0100818A  
Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Lung Neoplasm Malignant Metastasis	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER DAY/ORAL		Nausea Vomiting		Magnesium Trisilicate	C		

Date:06/06/01ISR Number: 3735450-XReport Type:Expedited (15-DaCompany Report #B0109589A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Amnesia Clonic Convulsion	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
150 MG/PER DAY		Convulsion Dizziness Loss Of Consciousness Malaise Tremor		Lorazepam Irbesartan	C C		

Date:06/06/01ISR Number: 3735451-1Report Type:Expedited (15-DaCompany Report #B0109585A  
Age:19 YR Gender:Female I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 3.75 G/SINGLE		Agitation	Foreign	Zyban	PS	Glaxo Wellcome Inc	
Initial or Prolonged DOSE		Convulsion	Literature				
		Hallucinations, Mixed Hyperreflexia Hypertonia Mental Disorder Overdose Tachycardia	Health Professional				

Date:06/06/01ISR Number: 3735452-3Report Type:Expedited (15-DaCompany Report #D0017224A  
Age:69 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ TWICE		Movement Disorder	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged PER DAY /			Health Professional				
ORAL				Aspirin Glimepiride	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/06/01ISR Number: 3735456-0Report Type:Expedited (15-DaCompany Report #B0103385A  
 Age:69 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia Cough	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Dizziness	Professional				
TWICE PER DAY		Lower Respiratory Tract					
/ ORAL		Infection		Aspirin	C		
		Nausea		Simvastatin	C		
		Vomiting					

Date:06/07/01ISR Number: 3734968-3Report Type:Expedited (15-DaCompany Report #B0109628A  
 Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cardiac Arrest		Zyban	PS	Glaxo Wellcome	ORAL
Death		Confusional State					
1 DAY		Dysarthria					
		Hallucination					
		Overdose					
		Tremor					
		Vomiting					

Date:06/07/01ISR Number: 3735561-9Report Type:Direct  
 Age:17 YR Gender:Male I/FU:I Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin Sr 100	PS		ORAL
Required							
2/2/ORAL							
Intervention to							
Prevent Permanent							
Impairment/Damage							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged INTRAVENOUS Other IV	125 MG, QID,	Headache Pruritus Urticaria	Health Professional	Solu-Medrol	PS	Pharmacia And Upjohn Co	
ORAL				Hydroxyzine (Hydroxyzine)	SS		ORAL
ORAL				Prednisone (Prednisone)	SS		ORAL
150 MG, BID, ORAL				Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL
				Ortho Tri-Cyclen	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 200MG per day	98 DAY	Abortion Spontaneous Convulsion		Bupropion	PS	Glaxo Wellcome	ORAL
				Prozac	SS		ORAL
				Klonopin	SS		ORAL
				Multivitamin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/08/01ISR Number: 3735598-XReport Type:Expedited (15-DaCompany Report #A0138183B

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	200MG per day 98 DAY	Abortion Spontaneous		Bupropion	PS	Glaxo Wellcome	ORAL
Congenital Anomaly		Complications Of Maternal Exposure To Therapeutic Drugs		Prozac	SS		ORAL
		Convulsion		Klonopin	SS		ORAL
		Skull Malformation		Multivitamins	C		

Date:06/08/01ISR Number: 3735602-9Report Type:Expedited (15-DaCompany Report #A0149544A

Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Amnesia		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Brain Damage					
		Concussion					
		Convulsion					
		Fall					
		Head Injury					
		Loss Of Consciousness					
		Spinal Fracture					

Date:06/08/01ISR Number: 3735607-8Report Type:Expedited (15-DaCompany Report #A0150205A

Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Complications Of Maternal Exposure To Therapeutic Drugs		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day		Intra-Uterine Death		Ferrous Sulfate	C		ORAL
500MG Per day		Pregnancy		Folic Acid	C		ORAL

Date:06/08/01ISR Number: 3735613-3Report Type:Expedited (15-DaCompany Report #B0104336A  
Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Zyban Salbutamol	PS C	Glaxo Wellcome Glaxo Wellcome	

Date:06/08/01ISR Number: 3735618-2Report Type:Expedited (15-DaCompany Report #B0109566A  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Twice per day	29 DAY	Headache Vision Blurred		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/08/01ISR Number: 3736781-XReport Type:Expedited (15-DaCompany Report #B0107381A  
Age:40 YR Gender:Male I/FU:F

Outcome	PT
Death	Ecchymosis Excoriation Fall Head Injury Overdose

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Subdural Haematoma

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Foreign Health Professional	Zyban Fluoxetine Ethanol	PS C C	Glaxo Wellcome Inc	

Date:06/08/01ISR Number: 3736839-5Report Type:Expedited (15-DaCompany Report #B0109966A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 150 MG / ORAL Intervention to Prevent Permanent Impairment/Damage		Anxiety Hyperventilation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/08/01ISR Number: 3736904-2Report Type:Expedited (15-DaCompany Report #B0109968A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / ORAL Initial or Prolonged		Cough Palpitations Paraesthesia	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/11/01ISR Number: 3736396-3Report Type:Expedited (15-DaCompany Report #A0150427A  
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1TAB Per day 6 WK		Diabetes Mellitus		Wellbutrin No Concurrent Medication	PS C	Glaxo Wellcome	ORAL

Date:06/11/01ISR Number: 3736398-7Report Type:Expedited (15-DaCompany Report #B0098184A  
Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG As Initial or Prolonged directed	4 DAY	Chest Pain Hyperhidrosis Hypoaesthesia Monoparesis Paraesthesia Skin Discolouration		Zyban	PS	Glaxo Wellcome	

Date:06/11/01ISR Number: 3736404-XReport Type:Expedited (15-DaCompany Report #B0110444A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - Initial or Prolonged		Arrhythmia Syncope		Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/11/01ISR Number: 3736405-1Report Type:Expedited (15-DaCompany Report #B0110446A

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Palpitations		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/11/01ISR Number: 3736406-3Report Type:Expedited (15-DaCompany Report #B0110451A

Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day	29 DAY	Condition Aggravated Syncope		Zyban	PS	Glaxo Wellcome	ORAL
UNKNOWN		Ventricular Tachycardia		Omeprazole	C		
UNKNOWN	1UNIT At			Atorvastatin	C		
UNKNOWN				Amiodarone	C		
UNKNOWN				Isosorbide	C		

Date:06/11/01ISR Number: 3736407-5Report Type:Expedited (15-DaCompany Report #B0110587A

Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG As directed		Photopsia		Zyntabac	PS	Glaxo Wellcome	ORAL
25MG per day		Retinal Detachment		Atenolol	C		



Date:06/11/01ISR Number: 3736408-7Report Type:Expedited (15-DaCompany Report #B0110591A  
Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG As		Myocardial Infarction		Zyntabac	PS	Glaxo Wellcome	ORAL
directed	10	DAY		Salmeterol + Fluticasone	SS	Glaxo Wellcome	
RESPIRATORY (INHALATION)	2U Twice per						
day	27	DAY					

Date:06/11/01ISR Number: 3736800-0Report Type:Direct Company Report #  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG, ONCE		Confusional State		Wellbutrin	PS		
Initial or Prolonged DAILY		Convulsion		Ambien	C		
		Diarrhoea Insomnia					

Date:06/11/01ISR Number: 3737369-7Report Type:Expedited (15-DaCompany Report #B0109628A  
Age:38 YR Gender:Male I/FU:I

Outcome	PT
Death	Cardiac Arrest Confusional State Dysarthria Hallucination, Visual Overdose Tremor

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vomiting

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/ORAL		Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/11/01ISR Number: 3737802-0Report Type:Expedited (15-DaCompany Report #B0110168A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Psoriasis	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							
DAY/ ORAL							

Date:06/11/01ISR Number: 3737810-XReport Type:Expedited (15-DaCompany Report #A0149803A  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
7.5 MG/ SEE							
TEXT/ ORAL		Catatonia					
		Convulsion		Amphetamine	C		
		Sedation		Cannabis	C		
		Tachycardia					

Date:06/11/01ISR Number: 3737820-2Report Type:Expedited (15-DaCompany Report #B0109581A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Foreign	Zyban	PS	Glaxo Wellcome Inc	
UNKNOWN	150 MG/TWICE						
PER DAY/		Dysgeusia	Health				
UNKNOWN		Electroencephalogram	Professional				

Abnormal  
Hyperventilation  
Joint Dislocation  
Salivary Hypersecretion

Lithium Carbonate C  
Thyroxine Sodium C  
Nortriptyline Hcl C

Date:06/11/01ISR Number: 3737821-4Report Type:Expedited (15-DaCompany Report #A0150080A  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Feeling Cold	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Hypoaesthesia	Consumer				
DAY/ ORAL							

Date:06/11/01ISR Number: 3737823-8Report Type:Expedited (15-DaCompany Report #A0149863A  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dry Mouth	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE		Dry Throat	Health				
PER DAY/ORAL		Emphysema	Professional				
		Laryngitis					
		Speech Disorder					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/11/01ISR Number: 3737824-XReport Type:Expedited (15-DaCompany Report #B0110170A  
 Age:58 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ORAL	Retroperitoneal Haemorrhage	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/11/01ISR Number: 3737826-3Report Type:Expedited (15-DaCompany Report #B0110171A  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability 150 MG/UNK/ORAL	Abdominal Distension Oedema Peripheral	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/11/01ISR Number: 3737827-5Report Type:Expedited (15-DaCompany Report #B0110167A  
 Age:69 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability 150 MG/TWICE PER DAY/ORAL	Staphylococcal Scalded Skin Syndrome	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/11/01ISR Number: 3737869-XReport Type:Expedited (15-DaCompany Report #A0149363A  
 Age:14 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG /TWICE Initial or Prolonged PER DAY / ORAL	Blood Pressure Increased Disturbance In Attention Drug Ineffective	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Grand Mal Convulsion  
Hallucination, Visual  
Intentional Misuse  
Irritability  
Memory Impairment  
Sedation  
Tachycardia

Wellbutrin Tablet  
(Bupropion  
Hydrochloride) SS  
Paracetamol  
(Formulation  
Unknown)  
(Acetaminophen) SS  
Loperamide  
Hydrochloride Tablet  
(Loperamide  
Hydrochloride) SS  
Paroxetine  
Hydrochloride C  
Tetracycline C  
Tretinoin C

Date:06/11/01ISR Number: 3738380-2Report Type:Periodic Company Report #252815  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Induced	Consumer	Accutane	PS	Hlr Technology	ORAL
40 MG DAILY							
ORAL							
				Wellbutrin	SS		
				Effexor	C		
				Spermicide	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/12/01ISR Number: 3737249-7Report Type:Expedited (15-DaCompany Report #B0100125A  
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Chest Pain		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged per day	8 DAY	Ecchymosis					
		Headache		Inhalers	C		

Date:06/12/01ISR Number: 3737253-9Report Type:Expedited (15-DaCompany Report #B0108029A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 50MG Twice		Atrial Flutter Cardiac Failure		Zyban Metoprolol Tartrate	PS C	Glaxo Wellcome	ORAL
per day		Dyspnoea		Diltiazem Hydrochloride	C		ORAL

Date:06/12/01ISR Number: 3737263-1Report Type:Expedited (15-DaCompany Report #B0109965A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 15 DAY		Formication		Zyban	PS	Glaxo Wellcome	ORAL
		Headache Nausea Pruritus Sleep Disorder					

Date:06/12/01ISR Number: 3737265-5Report Type:Expedited (15-DaCompany Report #B0110766A  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Chest Discomfort	Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	4 DAY				
	Chest Pain	Enalapril	C		ORAL
20MG Per day		Beclomethasone	C	Glaxo Wellcome	
RESPIRATORY					
(INHALATION)	50MCG Twice				
per day		Nifedipine	C		ORAL
60MG Per day		Bendrofluazide	C	Glaxo Wellcome	ORAL
2.5MG Per day		Salbutamol	C	Glaxo Wellcome	
RESPIRATORY					
(INHALATION)	100MCG Twice				
per day		Insulin Lispro	C		
SUBCUTANEOUS		Aspirin	C		ORAL
75MG Per day					

Date:06/12/01ISR Number: 3737266-7Report Type:Expedited (15-DaCompany Report #B0110768A  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxo Wellcome	ORAL
Other		International Normalised Ratio Increased		Warfarin	C	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/12/01ISR Number: 3737267-9Report Type:Expedited (15-DaCompany Report #B0110774A  
 Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Asthenia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Hospitalization -		Confusional State					
per day	17 DAY						
Initial or Prolonged		Depression		Captopril	C	Glaxo Wellcome	ORAL
100MG per day				Zafirlukast	C		ORAL
20MG Twice		Disturbance In Attention					
per day		Haemophilus Infection					
600MG per day		Hypercapnia		Theophylline	C		ORAL
RESPIRATORY		Lung Disorder		Salmeterol	C	Glaxo Wellcome	
(INHALATION)		Respiratory Distress					
per day	50UG Twice	Thinking Abnormal		Fluticasone	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)							
per day	500UG Twice						
				Duovent	C		

Date:06/12/01ISR Number: 3738140-2Report Type:Direct Company Report #  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Pancytopenia		Wellbutrin / 150mg /			
Required				Glaxo Wellcome	PS	Glaxo Wellcome	ORAL
150 MG TID							
Intervention to							
ORAL							
Prevent Permanent				Celexa	SS	Forest	ORAL
20 MG QD ORAL							
Impairment/Damage				Lithium	C		



Date:06/12/01ISR Number: 3738768-XReport Type:Expedited (15-DaCompany Report #B0091930A  
Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Overdose	Foreign Literature Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:06/12/01ISR Number: 3738770-8Report Type:Expedited (15-DaCompany Report #B0088230A  
Age:18 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 50 Initial or Prolonged TABLET/SINGLE	Acidosis Agitation Coma Grand Mal Convulsion	Foreign Literature Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
DOSE/ORAL  24 TABLET/SINGLE	Hypotension Intentional Misuse Pyrexia		Anadin Extra Tablet (Anadin Extra)	SS		
DOSE/24	Sinus Tachycardia					
	Suicide Attempt Vomiting		Ethanol (Formulation Unknown) (Alcohol)	SS		

Date:06/12/01ISR Number: 3738773-3Report Type:Expedited (15-DaCompany Report #B0104336A  
Age:34 YR Gender:Male I/FU:F

Outcome	PT	Report Source
Death	Completed Suicide	Foreign Health

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Professional

Dose	Duration	Product	Role	Manufacturer	Route
150 MG		Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
		Salbutamol Sulphate	C		

Date:06/12/01ISR Number: 3738782-4Report Type:Expedited (15-DaCompany Report #A0150205A  
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG / TWICE PER DAY / ORAL		Complications Of Maternal Exposure To Therapeutic Drugs Intra-Uterine Death Pregnancy Uterine Dilation And Curettage	Foreign Study Health Professional	Zyban   Ferrous Sulfate Folic Acid	PS   C C	Glaxo Wellcome Inc	ORAL

Date:06/12/01ISR Number: 3738919-7Report Type:Expedited (15-DaCompany Report #MPI-2001-05634(0)  
 Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 9 TABLETS, Required TAKEN OVER 72 Intervention to HOURS, PO Prevent Permanent TAKEN OVER 72 Impairment/Damage HOURS		Angina Pectoris Chest Pain Myocardial Infarction	Foreign Literature Health Professional	Semprex-D   Bupropion  Erythromycin (Erythromycin)	PS   SS  C	Celltech Pharmaceuticals Inc	ORAL

Date:06/12/01ISR Number: 3738946-XReport Type:Expedited (15-DaCompany Report #B0109566A  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Headache	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Vision Blurred					
TWICE PER DAY							
/ ORAL							

Date:06/12/01ISR Number: 3738981-1Report Type:Expedited (15-DaCompany Report #A0138183B  
Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Abortion Spontaneous	Study	Wellbutrin	PS	Glaxo Wellcome Inc	
TRANSPLACENTAL	SEE TEXT/						
Congenital Anomaly		Complications Of Maternal	Health				
ORAL		Exposure To Therapeutic	Professional	Fluoxetine			
		Drugs		Hydrochloride			
		Convulsion		Capsule (Fluoxetine			
		Skull Malformation		Hydrochloride)	SS		
TRANSPLACENTAL	ORAL						
				Clonazepam Tablet			
				(Clonazepam)	SS		
TRANSPLACENTAL	ORAL						
				Multivitamin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/12/01ISR Number: 3738982-3Report Type:Expedited (15-DaCompany Report #A0138183A

Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Study	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
SEE TEXT/ORAL							
		Complications Of Maternal Exposure To Therapeutic Drugs Convulsion	Health Professional	Fluoxetine Hydrochloride Capsule (Fluoxetine Hydrochloride)	SS		ORAL
ORAL							
				Clonazepam Tablet (Clonazepam)	SS		ORAL
ORAL							
				Multivitamin	C		

Date:06/13/01ISR Number: 3738076-7Report Type:Expedited (15-DaCompany Report #A0107343A

Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
100MG Twice per day							
		Drug Interaction					
		Hypertension Insomnia Sinus Tachycardia		Paxil	SS	Glaxo Wellcome	ORAL

Date:06/13/01ISR Number: 3738077-9Report Type:Expedited (15-DaCompany Report #A0108484A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Aggression Anxiety Dysgeusia Dysphagia Erythema Face Oedema Fatigue Halitosis		Zyban	PS	Glaxo Wellcome	ORAL

Insomnia  
 Nail Disorder  
 Oedema Peripheral  
 Oral Pain  
 Oral Soft Tissue Disorder  
 Pain In Extremity  
 Pruritus  
 Salivary Hypersecretion  
 Stomatitis

Date:06/13/01ISR Number: 3738078-0Report Type:Expedited (15-DaCompany Report #A0120078A  
 Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	1 MON	Bipolar I Disorder Condition Aggravated		Zyban	PS	Glaxo Wellcome	ORAL
.5MG Twice per day		Convulsion Mood Swings		Clonazepam	C		ORAL
20MG Per day		Psychotic Disorder		Paxil	C	Glaxo Wellcome	ORAL
2MG Twice per day		Visual Disturbance		Perphenazine	C	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/13/01ISR Number: 3738079-2Report Type:Expedited (15-DaCompany Report #A0124641A  
Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice			Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
	per day	6 MON	Dehydration				
			Heat Stroke				
			Sudden Death	Alcohol	SS		

Date:06/13/01ISR Number: 3738080-9Report Type:Expedited (15-DaCompany Report #A0130642A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Twice			Zyban	PS	Glaxo Wellcome	ORAL
	per day	15 DAY	Blood Thyroid Stimulating				
			Hormone Increased				
			Dermatitis				
			Malaise				
			Pyrexia				
			Red Blood Cell				
			Sedimentation Rate				
			Increased				

Date:06/13/01ISR Number: 3738084-6Report Type:Expedited (15-DaCompany Report #A0147746A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	75MG Per day			Wellbutrin	PS	Glaxo Wellcome	ORAL
	1 WK		Condition Aggravated				
			Crying	Klonopin	C		
			Feeling Abnormal				
			Nervousness				
			Petit Mal Epilepsy				

Date:06/13/01ISR Number: 3738087-1Report Type:Expedited (15-DaCompany Report #A0150616A  
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypersensitivity		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	19	DAY					
				Ginseng Vitamins	C C		ORAL

Date:06/13/01ISR Number: 3738089-5Report Type:Expedited (15-DaCompany Report #A0150720A  
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Haematemesis		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day		Pancreatitis					
				Aspirin	C		ORAL
325MG Per day							
				Colace	C		
100MG Twice							
per day							
14MG Per day				Habitrol	C		
15MG Per day				Oxazepam	C		
				Parafon Forte	C		
				Zocor	C		
10MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/13/01ISR Number: 3738090-1Report Type:Expedited (15-DaCompany Report #A0150723A  
 Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Petechiae		Zyban	PS	Glaxo Wellcome	ORAL
150TAB Twice		Rash Pruritic					
per day							

Date:06/13/01ISR Number: 3738091-3Report Type:Expedited (15-DaCompany Report #A0150739A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Physical Assault		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:06/13/01ISR Number: 3738092-5Report Type:Expedited (15-DaCompany Report #A0150823A  
 Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alanine Aminotransferase		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Increased					
per day	17 DAY	Angioneurotic Oedema					
		Arthralgia					
		Aspartate					
		Aminotransferase					
		Increased					
		Asthenia					
		Dizziness					
		Dysphagia					
		Dyspnoea					
		Hypoaesthesia					
		Osteoarthritis					
		Paraesthesia					
		Pruritus					
		Urticaria					



Date:06/13/01ISR Number: 3738093-7Report Type:Expedited (15-DaCompany Report #A0150828A  
Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day				Airomir	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)							

Date:06/13/01ISR Number: 3738094-9Report Type:Expedited (15-DaCompany Report #A0150884A  
Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Retroperitoneal Fibrosis		Bupropion	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	3	MON					

Date:06/13/01ISR Number: 3738095-0Report Type:Expedited (15-DaCompany Report #A0150891A  
Age:42 YR Gender:Female I/FU:F

Outcome	PT
Other	Agitation Confusional State Epileptic Aura

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Flushing Paraesthesia Psychotic Disorder	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day				Bupropion	PS	Glaxo Wellcome	ORAL

Date:06/13/01ISR Number: 3738096-2Report Type:Expedited (15-DaCompany Report #A0150908A  
Age:28 YR Gender:Female I/FU:F

Outcome Dose Other	Duration	PT Anxiety Crying Dizziness Fear Feeling Jittery Hyperventilation Panic Reaction	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	5 DAY			Bupropion	PS	Glaxo Wellcome	ORAL
				Triphasil	C		

Date:06/13/01ISR Number: 3738098-6Report Type:Expedited (15-DaCompany Report #A0150917A  
Age:38 YR Gender:Female I/FU:F

Outcome Dose Other	Duration	PT Anxiety Hallucination, Auditory Panic Reaction	Report Source	Product	Role	Manufacturer	Route
75MG Twice per day				Bupropion	PS	Glaxo Wellcome	ORAL
.5MG Twice per day				Clonazepam	C		ORAL
500MG Twice per day				Epival	C		ORAL

Date:06/13/01ISR Number: 3738101-3Report Type:Expedited (15-DaCompany Report #B0098815A  
Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day 3 DAY	Anaphylactic Reaction		Zyban	PS	Glaxo Wellcome	ORAL
	5MG Per day	Arrhythmia		Bisoprolol	C		
	75MG Per day	Chest Discomfort		Aspirin	C		ORAL
	1G Twice per day	Decreased Appetite		Metformin	C		ORAL
	160MG Per day	Dehydration					
		Dry Skin		Gliclazide	C		ORAL
		Feeling Abnormal					
		Insomnia					
		Lethargy					
		Malaise					
		Myocardial Infarction					
		Oedema					
		Pain					

Date:06/13/01ISR Number: 3738110-4Report Type:Expedited (15-DaCompany Report #B0110962A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	10ML Three times per day	Small Cell Lung Cancer Stage Unspecified		Zyban	PS	Glaxo Wellcome	ORAL
		Tonsillitis		Orciprenaline	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/13/01ISR Number: 3738111-6Report Type:Expedited (15-DaCompany Report #B0110963A  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anger		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Depression					
per day	32 DAY	Insomnia		Alcohol	C		
		Self Mutilation					

Date:06/13/01ISR Number: 3739226-9Report Type:Expedited (15-DaCompany Report #HQ5327428DEC2000  
Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Craniopharyngioma	Health	Effexor Xr	PS	Wyeth Ayerst Laboratories	ORAL
Initial or Prolonged		Diverticulum Oesophageal	Professional				
112.5 MG		Gastric Polyps					
Other		Hypokalaemia		Nicotine (Nicotine)	SS		
DAILY, ORAL		Hyponatraemia		Wellbutrin			
		Pituitary Tumour Benign		(Amfebutamone			
		Salivary Hypersecretion		Hydrochloride)	SS		ORAL
150 MG 2X PER		Speech Disorder					
1 DAY, ORAL		Tardive Dyskinesia		Trazodone			
		Visual Field Defect		(Trazodone)	C		
				Clonazepam(Clonazepa			
				m0	C		
				Dexamfetamine			
				(Dexamfetamine)	C		
				Thyroxine Sodium			
				(Levothyroxine			
				Sodium)	C		

Date:06/13/01ISR Number: 3739464-5Report Type:Expedited (15-DaCompany Report #B0110587A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 150 MG / AS		Retinal Detachment	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Intervention to DIRECTED / Prevent Permanent ORAL Impairment/Damage			Consumer				
				Atenolol	C		

Date:06/13/01ISR Number: 3739468-2Report Type:Expedited (15-DaCompany Report #B0110444A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 150 MG ORAL		Arrhythmia Syncope	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged			Professional				

Date:06/13/01ISR Number: 3739469-4Report Type:Expedited (15-DaCompany Report #B0110446A  
Age: Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG ORAL		Heart Rate Irregular	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/13/01ISR Number: 3739470-0Report Type:Expedited (15-DaCompany Report #B0110591A  
 Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (AS DIRECTED)							
ORAL							
RESPIRATORY (INHALATION)	2 UNKNOWN			Salmeterol + Fluticasone (Salmeterol + Fluticasone)	SS		
(TWICE PER DAY) INHALED							

Date:06/13/01ISR Number: 3739473-6Report Type:Expedited (15-DaCompany Report #B0110451A  
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Condition Aggravated Syncope	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (TWICE PER DAY) ORAL		Ventricular Tachycardia					
				Omeprazole	C		
				Atorvastatin Calcium	C		
				Amiodarone	C		
				Isosorbide	C		

Date:06/13/01ISR Number: 3739499-2Report Type:Expedited (15-DaCompany Report #B0098184A  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG (AS DIRECTED)		Chest Pain Hyperhidrosis Hypoaesthesia Monoparesis Paraesthesia Skin Discolouration	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:06/13/01ISR Number: 3739625-5Report Type:Expedited (15-DaCompany Report #A0150427A  
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1 TABLET / PER DAY / ORAL		Blood Chloride Decreased Blood Sodium Decreased Blood Urea Increased Diabetes Mellitus	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Date:06/15/01ISR Number: 3740147-6Report Type:Expedited (15-DaCompany Report #A0106360A  
Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day .625MG Per day		Facial Palsy Hypoaesthesia Urticaria		Zyban  Premarin	PS  C	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/15/01ISR Number: 3740148-8Report Type:Expedited (15-DaCompany Report #A0128949A  
Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	1 WK	Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/15/01ISR Number: 3740149-XReport Type:Expedited (15-DaCompany Report #A0141486A  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged		Atrial Fibrillation Atrial Flutter Cardiac Failure Congestive		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/15/01ISR Number: 3740150-6Report Type:Expedited (15-DaCompany Report #A0144603A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day		Arthralgia Bronchospasm Burning Sensation Chest Pain Dermatitis Dyspnoea Face Oedema Insomnia Joint Stiffness Myocardial Infarction Oedema Peripheral Pruritus Urticaria		Zyban	PS	Glaxo Wellcome	ORAL



Date:06/15/01ISR Number: 3740153-1Report Type:Expedited (15-DaCompany Report #A0149803A  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Zyban	PS	Glaxo Wellcome	ORAL
7.5G	See text	Catatonia		Amphetamine	C		
		Convulsion		Cannabis	C		
		Intentional Misuse					
		Sedation					
		Tachycardia					

Date:06/15/01ISR Number: 3740156-7Report Type:Expedited (15-DaCompany Report #A0150913A  
Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Unstable		Zyban	PS	Glaxo Wellcome	ORAL
1 WK		Asthenia		Alprazolam	C		
.5MG As		Chest Discomfort					
required		Dyspnoea		Aspirin	C		ORAL
80MG Per day		Heart Rate Increased		Atenolol	C		ORAL
50MG Per day		Hypertension		Baycol	C		ORAL
.2MG Per day		Hypoaesthesia		Dalteparin	C		
SUBCUTANEOUS	10000IU Twice						

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

per day 4 DAY

Prinzide C ORAL

Date:06/15/01ISR Number: 3740157-9Report Type:Expedited (15-DaCompany Report #A0150949A  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Twice			Abdominal Pain	Zyban	PS	Glaxo Wellcome	ORAL
per day			Arthralgia				
			Chest Pain	Nefazodone	SS		
			Diarrhoea	Aspirin	C		
			Drug Interaction	Atenolol	C		
			Intestinal Obstruction	Lipitor	C		ORAL
			Malaise				
			Neck Pain				

Date:06/15/01ISR Number: 3740158-0Report Type:Expedited (15-DaCompany Report #B0104342A  
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Twice			Circulatory Collapse	Zyban	PS	Glaxo Wellcome	ORAL
per day	13	DAY	Convulsion				
				Alcohol	SS		
				Triazolam	SS		
.25MG Per day				Mercilon	SS		ORAL

Date:06/15/01ISR Number: 3740159-2Report Type:Expedited (15-DaCompany Report #B0109704A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death			Bone Marrow Depression	Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	36	DAY					
Hospitalization -			Cerebral Ischaemia	Sulpiride	C		

Initial or Prolonged

Coma  
Convulsion  
Haemolytic Anaemia  
Hepatitis  
Hypertonia  
Lipase Increased  
Metabolic Acidosis  
Pancytopenia  
Paralysis Flaccid  
Peripheral Ischaemia  
Petechiae  
Respiratory Failure  
Shock  
Tachycardia  
Thrombotic  
Thrombocytopenic Purpura  
Trismus  
Urinary Incontinence

Migril

C

Glaxo Wellcome

Date:06/15/01ISR Number: 3740163-4Report Type:Expedited (15-DaCompany Report #B0110588A  
Age:42 YR Gender:Male I/FU:I

Outcome PT  
Hospitalization - Convulsion  
Initial or Prolonged Fall  
Headache

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
13 DAY		Insomnia Muscle Spasms Salivary Hypersecretion Sensation Of Pressure		Zyban	PS	Glaxo Wellcome	

Date:06/15/01ISR Number: 3740198-1Report Type:Expedited (15-DaCompany Report #A0075655A  
Age:49 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	500MG Per day 1.25MG Per day	5 DAY	Amnesia Dyspnoea Emotional Disorder Grand Mal Convulsion Status Epilepticus Suicidal Ideation		Zyban Calcium Ces St. Johns Wort Vitamin C	PS C C C C	Glaxo Wellcome	ORAL      ORAL

Date:06/15/01ISR Number: 3740199-3Report Type:Expedited (15-DaCompany Report #A0082610A  
Age:55 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day Initial or Prolonged 4 YR	6 DAY		Anaphylactoid Reaction Angioneurotic Oedema		Zyban Synthroid Hormone Replacement Therapy	PS C C	Glaxo Wellcome	ORAL    ORAL

Date:06/15/01ISR Number: 3740200-7Report Type:Expedited (15-DaCompany Report #A0086553A  
Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Other		Myocardial Infarction					
per day	11	DAY					
				Glucophage	C		
				Premarin	C		
				Lipitor	C		
				Diabeta	C		

Date:06/15/01ISR Number: 3740201-9Report Type:Expedited (15-DaCompany Report #A0090004A  
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardiac Arrest		Zyban	PS	Glaxo Wellcome	ORAL
150MG Single							
dose	1	DAY					
		Chest Discomfort					
.5MG Twice		Dyspnoea		Ativan	C		ORAL
per day		Pulmonary Embolism					
				Ceftin	C	Glaxo Wellcome	ORAL
500MG Twice							
per day							
RESPIRATORY				Combivent	C		
(INHALATION)	2	UNIT Twice					
per day							
RESPIRATORY				Pulmicort	C		
(INHALATION)	.5	MG Four					
times per day							
250MG Four				Tetracycline	C		ORAL
times per day							
400MG Per day				Uniphyl	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/15/01ISR Number: 3740202-0Report Type:Expedited (15-DaCompany Report #A0090299A  
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other	100MG Per day 7 DAY	Abdominal Pain		Zyban	PS	Glaxo Wellcome	ORAL
		Chest Pain					
		Pericarditis					
		Vomiting					

Date:06/15/01ISR Number: 3740203-2Report Type:Expedited (15-DaCompany Report #A0096983A  
Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -	100MG per day 33 DAY	Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	600MG per day	Generalised		Ibuprofen	C		
Other	1000MG per day	Non-Convulsive Epilepsy		Epival	C		ORAL
		Headache					

Date:06/15/01ISR Number: 3740204-4Report Type:Expedited (15-DaCompany Report #A0104189A  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other	150MG Per day MON	Abdominal Distension		Zyban	PS	Glaxo Wellcome	ORAL
		Arthralgia		Triphasil	C		ORAL
		Back Pain					
		Chest Pain					
		Dyspepsia					
		Eye Irritation					
		Hypersensitivity					
		Urticaria					

Date:06/15/01ISR Number: 3740205-6Report Type:Expedited (15-DaCompany Report #A0105423A  
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day 9 DAY Disability		Multiple Sclerosis		Zyban	PS	Glaxo Wellcome	ORAL
				No Concurrent Medications	C		

Date:06/15/01ISR Number: 3740206-8Report Type:Expedited (15-DaCompany Report #A0105871A  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation Overdose		Wellbutrin Nortriptyline Cocaine	PS SS SS	Glaxo Wellcome	

Date:06/15/01ISR Number: 3740207-XReport Type:Expedited (15-DaCompany Report #A0105877A  
Age:26 YR Gender:Female I/FU:F

Outcome	PT
Other	Confusional State Cyanosis Dyspnoea Eye Rolling Movement Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sedation

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day			Zyban	PS	Glaxo Wellcome	ORAL
100MG As required			Imitrex	C	Glaxo Wellcome	
40MG Twice per day			Propranolol	C		
2U Twice per day			Salbutamol	C	Glaxo Wellcome	
21 DAY			Triquilar	C		

Date:06/15/01  
 Age:34 YR Gender:Male I/FU:F  
 ISR Number: 3740208-1  
 Report Type:Expedited (15-DaCompany Report #A0106433A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dermatitis		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice per day	20 DAY	Dyspepsia					
600MG Twice per day	11 DAY	Ecchymosis		Penicillin	SS	Glaxo Wellcome	
		Face Oedema					
		Fatigue					
		Lip Dry					
		Myalgia					
		Pruritus					
		Throat Tightness					
		Urticaria					



Date:06/15/01ISR Number: 3740209-3Report Type:Expedited (15-DaCompany Report #A0129592A  
Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day	Agitation		Zyban	PS	Glaxo Wellcome	ORAL
	20 DAY	Dyspnoea		No Concurrent Medications	C		
		Erythema Multiforme					
		Face Oedema					
		Leukocytoclastic Vasculitis					
		Serum Sickness					

Date:06/15/01ISR Number: 3740214-7Report Type:Expedited (15-DaCompany Report #A0150827A  
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	100MG Per day	Bundle Branch Block Left		Wellbutrin	PS	Glaxo Wellcome	ORAL
	25MG Per day	Chest Pain		Vioxx	C		
	100MG Per day	Dizziness		Thiamine	C		
	20MG Per day	Headache		Ritalin Sr	C		ORAL
	150MG Per day	Hypertension		Effexor Xr	C		ORAL
	.625MG Per day	Vomiting		Conjugated Estrogens	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/15/01ISR Number: 3740215-9Report Type:Expedited (15-DaCompany Report #A0150905A  
 Age:21 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	19 DAY	Aggression Anxiety Coordination Abnormal Depression Dizziness Fatigue Headache Intentional Self-Injury Lethargy Panic Disorder Paranoia Phobia		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/15/01ISR Number: 3740217-2Report Type:Expedited (15-DaCompany Report #A0151169A  
 Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	8 DAY	Facial Palsy Hypoaesthesia Sensory Disturbance		Zyban	PS	Glaxo Wellcome	ORAL
TRANSDERMAL times per week	50MCG Two			Estraderm	C		
				Calcium + Vitamin D Avalide	C C		ORAL ORAL

Date:06/15/01ISR Number: 3740218-4Report Type:Expedited (15-DaCompany Report #A0151174A  
 Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other		Anaemia		Wellbutrin	PS	Glaxo Wellcome	ORAL
250MG Per day							
		Bronchiolitis		Anafranil	SS		
125MG Per day	YR						
		Cough		Lithium	C		
1050MG Per							
day		Drug Interaction					
.25MG Per day		Hyperhidrosis		Rivotril	C		
		Hypoxia					
		Pyrexia					

Date:06/15/01ISR Number: 3740224-XReport Type:Expedited (15-DaCompany Report #B0110444A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia		Zyban	PS	Glaxo Wellcome	ORAL
7 DAY							
Hospitalization - RESPIRATORY		Coma		Fluticasone	C	Glaxo Wellcome	
Initial or Prolonged (INHALATION)		Malaise					
RESPIRATORY		Nausea		Salmeterol	C	Glaxo Wellcome	
(INHALATION)		Pulmonary Embolism					
		Syncope					

Date:06/15/01ISR Number: 3740225-1Report Type:Expedited (15-DaCompany Report #B0110521A  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abnormal Behaviour		Zyban	PS	Glaxo Wellcome	
3 DAY							
Initial or Prolonged		Suicide Attempt		Clonazepam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sodium Valproate C  
Anxiolytic C

Date:06/15/01ISR Number: 3740226-3Report Type:Expedited (15-DaCompany Report #B0110727A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day		Coronary Artery Disease	Bupropion	PS	Glaxo Wellcome	ORAL
			Myocardial Infarction				

Date:06/15/01ISR Number: 3740227-5Report Type:Expedited (15-DaCompany Report #B0110883A  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	10 DAY		Cardiomyopathy	Zyban	PS	Glaxo Wellcome	ORAL
			Left Ventricular Failure	Amitriptyline	C		ORAL
			Pulmonary Oedema	Nitrazepam	C		ORAL
			Sudden Death	Loperamide	C		ORAL
				Co-Codamol	C		ORAL

Date:06/15/01ISR Number: 3740229-9Report Type:Expedited (15-DaCompany Report #B0111230A  
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Twice		Abdominal Pain	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	per day		Hepatic Steatosis				
Other	31 DAY		Hepatomegaly				

Date:06/15/01ISR Number: 3740230-5Report Type:Expedited (15-DaCompany Report #D0016065A  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Formication		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Gastrointestinal Disorder					
per day	10	DAY					
		Headache					
		Paraesthesia					
		Sensation Of Pressure					

Date:06/15/01ISR Number: 3740751-5Report Type:Expedited (15-DaCompany Report #B0108029A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Atrial Fibrillation	Foreign	Bupropion			
Initial or Prolonged		Atrial Flutter	Health	Hydrochloride	PS	Glaxo Wellcome Inc	
150 MG		Cardiac Failure	Professional	Metoprolol Tartrate	C		
				Diltiazem			
				Hydrochloride	C		

Date:06/15/01ISR Number: 3741098-3Report Type:Expedited (15-DaCompany Report #B0110774A  
Age:67 YR Gender:Male I/FU:I

Outcome	PT
Death	Asthenia
Hospitalization -	Confusional State
Initial or Prolonged	Depression

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Disturbance In Attention Haemophilus Infection Hypercapnia	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Lung Disorder Respiratory Distress					
/ ORAL				Captopril	C		
				Zafirlukast	C		
				Theophylline	C		
				Salmeterol Xinafoate	C		
				Fluticasone			
				Propionate	C		
				Duovent	C		

Date:06/15/01ISR Number: 3741110-1Report Type:Expedited (15-DaCompany Report #B0110768A  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG / ORAL Prevent Permanent Impairment/Damage		International Normalised Ratio Increased	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Warfarin Sodium	C		

Date:06/15/01ISR Number: 3741111-3Report Type:Expedited (15-DaCompany Report #B0110766A  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG / PER DAY / ORAL Prevent Permanent Impairment/Damage		Angina Pectoris Chest Discomfort	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Chest Pain					
				Enalapril	C		
				Beclomethasone			
				Dipropion.	C		

Nifedipine	C
Bendrofluazide	C
Salbutamol Sulphate	C
Insulin Lispro	C
Aspirin	C

Date:06/15/01ISR Number: 3741112-5Report Type:Expedited (15-DaCompany Report #B0109965A  
 Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Formication Headache	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL		Nausea Pruritus Sleep Disorder					

Date:06/15/01ISR Number: 3741113-7Report Type:Expedited (15-DaCompany Report #B0100125A  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Chest Pain Ecchymosis	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
150 MG / TWICE PER DAY		Headache	Professional				

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Freedom Of Information (FOI) Report

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Inhaler C

Date:06/18/01ISR Number: 3740564-4Report Type:Expedited (15-DaCompany Report #A0075616A  
 Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypertension		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Hypertensive Crisis					
per day	5	DAY		Ramipril	C		ORAL
2.5MG Per day				Lorazepam	C		ORAL
1MG As							
required							

Date:06/18/01ISR Number: 3740566-8Report Type:Expedited (15-DaCompany Report #A0083661A  
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Hypoglycaemia					
per day	16	DAY		Humulin R	SS		
		Medication Error		Humulin N	C		
				Alcohol	C		

Date:06/18/01ISR Number: 3740567-XReport Type:Expedited (15-DaCompany Report #A0087951A  
 Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Glucose Decreased		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Convulsion					
per day							



Humalog	C	ORAL
Humulin N	C	
Regular Insulin	C	

Date:06/18/01ISR Number: 3740568-1Report Type:Expedited (15-DaCompany Report #A0090158A  
 Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Twice	Leukopenia		Zyban	PS	Glaxo Wellcome	ORAL
per day	13 DAY	Neutropenia					
				Acetaminophen	C		
				Chlorpheniramine	C		
				Cytosar	C		
				Etoposide	C		
INTRAVENOUS	240MG Per day	5 DAY					
				Idarubicin	C		
				Novantrone	C		
24MG Per day	5 DAY						
RESPIRATORY				Salbutamol	C	Glaxo Wellcome	
(INHALATION)							

Date:06/18/01ISR Number: 3740569-3Report Type:Expedited (15-DaCompany Report #A0090923A  
 Age:51 YR Gender:Male I/FU:F

Outcome	PT
Life-Threatening	Alcohol Withdrawal
Hospitalization -	Syndrome
Initial or Prolonged	Aspartate
	Aminotransferase
	Increased



Other		Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
		Tendon Disorder					
per day	16	DAY					
		Urticaria		Ortho 1/35	C		

Date:06/18/01ISR Number: 3740573-5Report Type:Expedited (15-DaCompany Report #A0105879A  
 Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthritis		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	15	DAY					
		Diarrhoea					
		Dizziness					
		Erythema Multiforme					
		Leukocytosis					
		Serum Sickness					
		Tachycardia					
		Tongue Disorder					

Date:06/18/01ISR Number: 3740574-7Report Type:Expedited (15-DaCompany Report #A0106203A  
 Age:40 YR Gender:Male I/FU:F

Outcome	PT
Other	Abdominal Pain
	Proctitis
	Rectal Haemorrhage

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Rectal Tenesmus Vomiting	Report Source	Product	Role	Manufacturer	Route
100MG Per day	1 MON			Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
RECTAL				Betnesol	C	Glaxo Wellcome	
250MG Per day				Desyrel	C		ORAL
100MG Per day				Imuran	C	Glaxo Wellcome	ORAL
75MG As				Voltaren Sr	C		ORAL
required							

Date:06/18/01ISR Number: 3740575-9Report Type:Expedited (15-DaCompany Report #A0106475A  
Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	17 DAY	Arthropathy Crepitations Oedema Urticaria		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/18/01ISR Number: 3740576-0Report Type:Expedited (15-DaCompany Report #A0137061A  
Age:17 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 4500MG per day		Convulsion Suicide Attempt		Wellbutrin	PS	Glaxo Wellcome	ORAL
400MG per day				Celexa	SS		ORAL

Date:06/18/01ISR Number: 3740578-4Report Type:Expedited (15-DaCompany Report #A0143251A  
 Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	60 DAY	Fall					
500MG Twice		Loss Of Consciousness		Naproxen	C		
per day		Malaise					
20MG Per day				Losec	C		
300MG Per day				Aspirin	C		
5MG Per day				Norvasc	C		
50MG At night				Trazodone	C		
10MG Every				Provera	C		
two weeks							
3MG Three				Lectopam	C		
times per day							
200MG Per day				Zoloft	C		
				Fiorinal	C		
				Percocet	C		

Date:06/18/01ISR Number: 3740582-6Report Type:Expedited (15-DaCompany Report #A0150726A  
 Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Bradycardia		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Coma		Cardizem Cd	SS		ORAL
		Oliguria		Clonazepam	SS		ORAL
		Suicide Attempt		Demerol	SS		ORAL
				Desipramine	SS		ORAL
				Loxapine	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Ms Contin	SS	Glaxo Wellcome	ORAL
Trazodone	SS		ORAL
Zopiclone	SS		ORAL

Date:06/18/01ISR Number: 3740584-XReport Type:Expedited (15-DaCompany Report #A0150947A  
 Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Twice			Body Temperature	Zyban	PS	Glaxo Wellcome	ORAL
			Increased				
per day	18	DAY	Chest Pain	Lipitor	C		ORAL
40MG Per day			Face Oedema				
			Periorbital Oedema				
			Pharyngolaryngeal Pain				
			Pruritus				
			Urticaria				

Date:06/18/01ISR Number: 3740587-5Report Type:Expedited (15-DaCompany Report #A0151323A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
			Abdominal Pain	Zyban	PS	Glaxo Wellcome	ORAL
			Angina Pectoris	Lipidil Micro	SS		ORAL
			Drug Interaction				

Date:06/18/01ISR Number: 3740588-7Report Type:Expedited (15-DaCompany Report #A0151352A  
 Age:21 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
15G Single			Electrocardiogram Qrs	Zyban	PS	Glaxo Wellcome	ORAL
			Complex Prolonged				
dose				Ativan	C		

Date:06/18/01ISR Number: 3740589-9Report Type:Expedited (15-DaCompany Report #A0151353A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agranulocytosis		Bupropion	PS	Glaxo Wellcome	ORAL
		Infection		Rivotril	SS		ORAL

Date:06/18/01ISR Number: 3740592-9Report Type:Expedited (15-DaCompany Report #B0099946A  
Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Conductive Deafness		Bupropion			
		Influenza Like Illness		Hydrochloride	PS	Glaxo Wellcome	ORAL
150MG Per day							
		Middle Ear Disorder		Paracetamol	C		ORAL
500MG As							
required							

Date:06/18/01ISR Number: 3740597-8Report Type:Expedited (15-DaCompany Report #B0110591A  
Age:69 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Myocardial Infarction		Zyntabac	PS	Glaxo Wellcome	ORAL
300MG Per day	10 DAY			Salmeterol +			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Fluticasone SS Glaxo Wellcome

RESPIRATORY

(INHALATION) 2U Twice per

day

Date:06/18/01ISR Number: 3740601-7Report Type:Expedited (15-DaCompany Report #B0111327A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cardiomyopathy		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/18/01ISR Number: 3740603-0Report Type:Expedited (15-DaCompany Report #B0111329A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction		Zyntabac	PS	Glaxo Wellcome	ORAL

Date:06/18/01ISR Number: 3741394-XReport Type:Direct Company Report #  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aggression		Luvox	PS		
		Anger		Effexor	SS		
		Muscle Spasms		Trazodone	SS		
				Wellbutrin	SS		
				Tegretol	SS		
				Paxil	SS		

Date:06/18/01ISR Number: 3741492-0Report Type:Expedited (15-DaCompany Report #A0150917A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Required 75 MG TWICE Intervention to PER DAY ORAL Prevent Permanent Impairment/Damage	Anxiety  Hallucination, Auditory  Panic Reaction	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
			Clonazepam Semisodium Valproate	C C		

Date:06/18/01ISR Number: 3741493-2Report Type:Expedited (15-DaCompany Report #B0110962A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG ORAL		Small Cell Lung Cancer	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Stage Unspecified Tonsillitis		Orciprenaline	C		

Date:06/18/01ISR Number: 3741494-4Report Type:Expedited (15-DaCompany Report #B0110963A  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 150 MG TWICE Intervention to PER DAY ORAL Prevent Permanent Impairment/Damage		Anger  Depression  Insomnia Self Mutilation	Foreign	Zyban   Ethanol	PS   C	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/18/01ISR Number: 3741514-7Report Type:Expedited (15-DaCompany Report #A0130642A  
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 150 MG / Intervention to TWICE PER DAY Prevent Permanent / ORAL Impairment/Damage	15 DAY	Hormone Increased Dermatitis Malaise Pyrexia Red Blood Cell Sedimentation Rate Increased	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/18/01ISR Number: 3741516-0Report Type:Expedited (15-DaCompany Report #A0150891A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 150 MG / Intervention to TWICE PER DAY Prevent Permanent / ORAL Impairment/Damage		Agitation Confusional State Epileptic Aura Flushing Paraesthesia Psychotic Disorder	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/18/01ISR Number: 3741554-8Report Type:Expedited (15-DaCompany Report #A0147746A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 75 MG / PER DAY / ORAL		Convulsion Crying Feeling Abnormal Nervousness	Health Professional	Wellbutrin Clonazepam	PS C	Glaxo Wellcome Inc	ORAL

Date:06/18/01ISR Number: 3741567-6Report Type:Expedited (15-DaCompany Report #A0150739A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
ORAL		Convulsion Psychotic Disorder Sexual Offence	Professional				

Date:06/18/01ISR Number: 3741887-5Report Type:Expedited (15-DaCompany Report #A0108484A  
Age: Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Aggression Anxiety Breath Odour Dermatitis Exfoliative Dysgeusia Dysphagia Erythema Face Oedema Fatigue Insomnia Lip Dry Nail Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Oedema Oral Intake Reduced Oral Pain	Report Source	Product	Role	Manufacturer	Route
150 MG ORAL		Pain Pruritus	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Salivary Hypersecretion Skin Lesion Stomatitis					

Date:06/18/01ISR Number: 3741888-7Report Type:Expedited (15-DaCompany Report #A0124641A  
Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alcohol Poisoning	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Arrhythmia	Health				
PER DAY ORAL		Dehydration Heat Stroke Sudden Death	Professional	Ethanol (Formulation Unknown) (Alcohol)	SS		

Date:06/18/01ISR Number: 3741889-9Report Type:Expedited (15-DaCompany Report #B0098815A  
Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anaphylactic Reaction	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG PER		Arrhythmia	Literature				
DAY ORAL		Chest Discomfort Chest Pain Decreased Appetite Dehydration Dermatitis Dizziness Dry Mouth Dyspnoea Fatigue Headache	Health Professional	Bisoprolol Aspirin Metformin Hydrochloride Glicazide	C C C C C		

Insomnia  
Lethargy  
Malaise  
Myocardial Infarction  
Pain  
Swelling  
Tongue Oedema

Date:06/18/01ISR Number: 3741892-9Report Type:Expedited (15-DaCompany Report #A0150884A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 150 MG TWICE		Retroperitoneal Fibrosis	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Intervention to PER DAY ORAL 3	MON						
Prevent Permanent Impairment/Damage							

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Freedom Of Information (FOI) Report

Date:06/18/01ISR Number: 3741903-0Report Type:Expedited (15-DaCompany Report #A0120078A  
 Age:58 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG (TWICE PER DAY) ORAL	Bipolar Disorder Bipolar I Disorder Disorientation Disturbance In Attention Grand Mal Convulsion Mood Swings Psychotic Disorder Visual Disturbance	Foreign Health Professional	Bupropion Hydrochloride Clonazepam Paroxetine Hydrochloride Perphenazine	PS  C C C	Glaxo Wellcome Inc	ORAL

Date:06/18/01ISR Number: 3741919-4Report Type:Expedited (15-DaCompany Report #A0150723A  
 Age:37 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to 150 TABLET/ Prevent Permanent TWICE PER Impairment/Damage DAY/ ORAL	Dermatitis Petechiae Pruritus	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:06/18/01ISR Number: 3741920-0Report Type:Expedited (15-DaCompany Report #A0150616A  
 Age:32 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to 150 MG/ TWICE Prevent Permanent PER DAY/ ORAL 19 Impairment/Damage	Drug Hypersensitivity	Foreign	Bupropion Hydrochloride Ginseng Multivitamin	PS  C C	Glaxo Wellcome Inc	ORAL

Date:06/18/01ISR Number: 3741921-2Report Type:Expedited (15-DaCompany Report #A0150823A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Alanine Aminotransferase	Foreign	Bupropion			
Intervention to		Increased		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
Prevent Permanent		Angioneurotic Oedema					
PER DAY/ ORAL 17	DAY						
Impairment/Damage		Aspartate					
		Aminotransferase					
		Increased					
		Asthenia					
		Dizziness					
		Dysphagia					
		Dyspnoea					
		Hypoaesthesia					
		Osteoarthritis					
		Paraesthesia					
		Urticaria					

Date:06/18/01ISR Number: 3741922-4Report Type:Expedited (15-DaCompany Report #A0150828A  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Myocardial Infarction	Foreign	Bupropion			
				Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Salbutamol Sulphate C

Date:06/18/01ISR Number: 3741923-6Report Type:Expedited (15-DaCompany Report #A0150720A  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG / TWICE PER DAY/ ORAL		Haematemesis Pancreatitis	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Aspirin C  
Docusate Sodium C  
Nicotine C  
Oxazepam C  
Parafon Forte C  
Simvastatin C

Date:06/18/01ISR Number: 3741924-8Report Type:Expedited (15-DaCompany Report #A0150908A  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG TWICE Prevent Permanent PER DAY/ ORAL 5 Impairment/Damage	5 DAY	Anxiety Crying Dizziness Fear Feeling Jittery Hyperventilation Panic Disorder	Foreign	Bupropion Hydrochloride  Triphasil	PS  C	Glaxo Wellcome Inc	ORAL

Date:06/18/01ISR Number: 3741925-XReport Type:Expedited (15-DaCompany Report #A0107343A  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Required 100 MG/ TWICE Intervention to PER DAY/ ORAL Prevent Permanent Impairment/Damage	Chest Pain  Drug Interaction  Hypertension Insomnia Sinus Tachycardia	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
SEE TEXT/  ORAL			Paroxetine Hydrochloride (Formulation Unknown) (Paroxetine Hydrochloride)	SS		ORAL

Date:06/18/01ISR Number: 3743005-6Report Type:Expedited (15-DaCompany Report #A0105871A  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death UNKNOWN		Aspiration	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	
UNKNOWN		Overdose		Nortriptyline (Nortriptyline)	SS		
UNKNOWN				Cocaine (Cocaine)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/18/01ISR Number: 3743550-3Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #LBID00201001555

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Coordination Abnormal Paranoia	Health Professional	Lithobid	PS	Solvay Pharmaceuticals	ORAL
900 MG DAILY		Speech Disorder					
PO				Seroquel (Seroquel)	SS		ORAL
600 MG DAILY							
PO				Wellbutrin - Slow Release (Amfebutamone Hydrochloride)	SS		ORAL
150 MG DAILY							
PO				Klonopin (Clonazepam)	SS		ORAL
2 MG DAILY PO							
UNK DAILY PO				Celebrex (Celecoxib)	SS		ORAL

Date:06/19/01ISR Number: 3741347-1Report Type:Expedited (15-DaCompany Report #A0086224A  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Medication Error		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Myocardial Infarction					
per day	2 DAY			No Concurrent Medications	C		

Date:06/19/01ISR Number: 3741348-3Report Type:Expedited (15-DaCompany Report #A0114129A  
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150MG Twice Other per day	8 DAY	Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
		Asthenia					
		Chest Pain		Diuretic	C		
		Coordination Abnormal					
		Cyanosis					
		Depressed Level Of Consciousness					
		Disorientation					
		Dizziness					
		Dyspepsia					
		Fatigue					
		Feeling Abnormal					
		Feeling Hot					
		Feeling Jittery					
		Headache					
		Hearing Impaired					
		Insomnia					
		Neck Pain					
		Oedema Peripheral					
		Pallor					
		Paraesthesia					
		Parkinsonian Gait					
		Pollakiuria					
		Tinnitus					
		Tremor					
		Vertigo					
		Visual Disturbance					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/19/01ISR Number: 3741349-5Report Type:Expedited (15-DaCompany Report #A0130455A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Asthenia		Bupropion	PS	Glaxo Wellcome	ORAL
150MG Unknown		Dysarthria		Fluvoxamine	SS		ORAL
100MG Per day		Gait Disturbance					
		Hypothermia					

Date:06/19/01ISR Number: 3742565-9Report Type:Expedited (15-DaCompany Report #B0110883A  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiomyopathy	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
SEE TEXT /		Left Ventricular Failure					
ORAL		Pulmonary Oedema		Amitriptyline	C		
		Sudden Death		Nitrazepam	C		
				Loperamide			
				Hydrochloride	C		
				Co-Codamol	C		

Date:06/19/01ISR Number: 3742566-0Report Type:Expedited (15-DaCompany Report #A0104189A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Abdominal Distension	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER		Arthralgia					
Intervention to		Back Pain					
DAY/ ORAL		Chest Pain					
Prevent Permanent		Dyspepsia		Triphasil	C		
(DURATION:		Eye Irritation					
Impairment/Damage		Hypersensitivity					
MONTHS)							

Urticaria

Date:06/19/01ISR Number: 3742567-2Report Type:Expedited (15-DaCompany Report #B0111230A  
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ TWICE		Abdominal Pain	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged PER DAY/ ORAL		Hepatic Steatosis					
Required Intervention to Prevent Permanent Impairment/Damage		Hepatomegaly					

Date:06/19/01ISR Number: 3742568-4Report Type:Expedited (15-DaCompany Report #A0150827A  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 100 MG/ PER		Bundle Branch Block Left	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Intervention to DAY/ ORAL		Chest Pain					
Prevent Permanent Impairment/Damage		Dizziness		Rofecoxib	C		
		Headache		Thiamine	C		
		Hypertension		Methylphenidate Hcl	C		
		Vomiting		Venlafaxine			
				Hydrochloride	C		
				Conjugated Estrogens	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/19/01ISR Number: 3742570-2Report Type:Expedited (15-DaCompany Report #A0150905A  
Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Aggression	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Anxiety					
Intervention to	19 DAY	Coordination Abnormal					
PER DAY/ ORAL		Depression					
Prevent Permanent		Dizziness					
Impairment/Damage		Fatigue					
		Headache					
		Lethargy					
		Panic Reaction					
		Paranoia					
		Phobia					
		Suicidal Ideation					

Date:06/19/01ISR Number: 3742571-4Report Type:Expedited (15-DaCompany Report #D0016065A  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abdominal Pain Upper	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Formication	Health				
PER DAY/ ORAL		Headache	Professional				
		Paraesthesia					

Date:06/19/01ISR Number: 3742572-6Report Type:Expedited (15-DaCompany Report #A0106433A  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Dermatitis	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Dyspepsia					
Intervention to	20 DAY	Ecchymosis		Phenoxymethylpenicil			
PER DAY/ ORAL		Face Oedema		lin K (Formulation			
Prevent Permanent							
Impairment/Damage							

Fatigue  
 Lip Dry  
 600 MG/ TWICE  
 Myalgia  
 PER DAY/ 11 DAY  
 Pruritus  
 Throat Tightness  
 Urticaria  
 Unknown) (Penicillin  
 V Potassium) SS

Date:06/19/01ISR Number: 3742577-5Report Type:Expedited (15-DaCompany Report #A0106360A  
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 150 MG/ TWICE		Facial Palsy	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Intervention to PER DAY/ ORAL		Hypoaesthesia					
Prevent Permanent Impairment/Damage		Urticaria		Conjugated Estrogens	C		

Date:06/19/01ISR Number: 3742589-1Report Type:Expedited (15-DaCompany Report #A0129592A  
 Age:31 YR Gender:Female I/FU:I

Outcome	PT
Disability	Agitation Arthralgia Dyspnoea Erythema Multiforme

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Leukocytoclastic Vasculitis Serum Sickness	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL			Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/19/01ISR Number: 3742590-8Report Type:Expedited (15-DaCompany Report #A0151174A  
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anaemia	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Required		Bronchiolitis					
250 MG/ PER Intervention to DAY/ ORAL		Cough		Clomipramine Hcl (Formulation Unknown)			
Prevent Permanent Impairment/Damage		Drug Interaction		(Clomipramine Hcl)	SS		
125 MG/ PER DAY		Hyperhidrosis Hypoxia Pyrexia					

(DURATION:

YEARS)

Lithium Salt	C
Clonazepam	C

Date:06/19/01ISR Number: 3742593-3Report Type:Expedited (15-DaCompany Report #A0151169A  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Facial Palsy	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Required		Hypoaesthesia					
150 MG/ TWICE Intervention to PER DAY/ ORAL 8 DAY		Sensory Disturbance		Oestradiol	C		
Prevent Permanent							



Impairment/Damage

Calcium With Vitamin  
D C  
Avalide C

Date:06/19/01ISR Number: 3742601-XReport Type:Expedited (15-DaCompany Report #B0110588A  
Age:42 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Abnormal Sensation In Eye Convulsion Fall Headache Insomnia Muscle Spasms Salivary Hypersecretion	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:06/19/01ISR Number: 3742606-9Report Type:Expedited (15-DaCompany Report #A0150949A  
Age:52 YR Gender:Male I/FU:I

Outcome	PT
Required Intervention to Prevent Permanent Impairment/Damage	Abdominal Pain Arthralgia Chest Pain Diarrhoea Drug Interaction Intestinal Obstruction Malaise

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Neck Pain

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
			Nefazodone Hydrochloride	SS		
			Aspirin	C		
			Atenolol	C		
			Atorvastatin Calcium	C		

Date:06/19/01ISR Number: 3742607-0Report Type:Expedited (15-DaCompany Report #A0150913A  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent DAY/ORAL Impairment/Damage	1 WK	Angina Unstable Asthenia Dyspnoea Heart Rate Increased Hypoaesthesia	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Alprazolam	C		
				Aspirin	C		
				Atenolol	C		
				Cerivastatin	C		
				Dalteparin Sodium	C		
				Prinzide	C		

Date:06/19/01ISR Number: 3742610-0Report Type:Expedited (15-DaCompany Report #B0110727A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG/ PER DAY/ ORAL		Coronary Artery Disease Myocardial Infarction	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Outcome	PT
Death	Aspartate
Hospitalization -	Aminotransferase
Initial or Prolonged	Increased
	Blood Alkaline
	Phosphatase Increased
	Blood Bilirubin Increased
	Blood Creatine
	Phosphokinase Increased
	Blood Lactate
	Dehydrogenase Increased
	Cerebral Ischaemia
	Coma
	Haemolytic Anaemia
	Hypertonia
	Lipase Increased
	Metabolic Acidosis
	Pancytopenia
	Respiratory Failure
	Shock
	Sinus Tachycardia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Thrombotic Thrombocytopenic Purpura Urinary Incontinence	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL			Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Sulpiride Migril	C C		

Date:06/19/01ISR Number: 3742619-7Report Type:Expedited (15-DaCompany Report #B0104342A  
Age:46 YR Gender:Female I/FU:F

Outcome Dose Other	Duration	PT Convulsion	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL			Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Ethanol (Formulation Unknown) (Alcohol) Triazolam Tablet (Triazolam)	SS SS		
.25 MG/ PER DAY/ ORAL				Mercilon (Formulation Unknown) (Mercilon)	SS		ORAL

Date:06/19/01ISR Number: 3742623-9Report Type:Expedited (15-DaCompany Report #A0128949A  
Age:28 YR Gender:Female I/FU:F

Outcome Dose Other	Duration	PT Abortion Spontaneous Complications Of Maternal	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE			Foreign Study	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

PER DAY/ORAL 1 WK Exposure To Therapeutic Health  
Drugs Professional

Date:06/19/01ISR Number: 3742629-XReport Type:Expedited (15-DaCompany Report #A0141486A  
Age:51 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Atrial Fibrillation	Foreign	Bupropion			
Initial or Prolonged	Atrial Flutter	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL	Cardiac Failure Congestive	Professional				

Date:06/19/01ISR Number: 3742630-6Report Type:Expedited (15-DaCompany Report #A0144603A  
Age: Gender:Male I/FU:F

Outcome	PT
Other	Arthralgia Bronchospasm Burning Sensation Chest Pain Dermatitis Dyspnoea Face Oedema Insomnia Joint Stiffness Myocardial Infarction

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Oedema Pruritus Urticaria	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE			Professional				
PER DAY/ ORAL							

Date:06/19/01ISR Number: 3742632-XReport Type:Expedited (15-DaCompany Report #A0149803A  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Other		Catatonia Convulsion					
7.5 G/ SEE		Intentional Misuse					
TEXT/ ORAL		Sedation Tachycardia		Amphetamine Cannabis	C C		

Date:06/19/01ISR Number: 3742655-0Report Type:Expedited (15-DaCompany Report #B0110521A  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
Hospitalization - Initial or Prolonged		Abnormal Behaviour Suicide Attempt	Consumer				
150 MG	3 DAY			Clonazepam Valproate Sodium Anxiolytic	C C C		

Date:06/19/01ISR Number: 3742814-7Report Type:Expedited (15-DaCompany Report #A0090004A  
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Foreign	Bupropion			
Death		Cardiac Arrest					

Pulmonary Embolism  
 Hydrochloride PS Glaxo Wellcome Inc ORAL  
 150 MG /  
 SINGLE DOSE /  
 ORAL 1 DAY  
 Lorazepam C  
 Cefuroxime Axetil C  
 Combivent C  
 Budesonide C  
 Tetracycline C  
 Theophylline C

Date:06/19/01ISR Number: 3742816-0Report Type:Expedited (15-DaCompany Report #A0086553A  
 Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Angina Pectoris	Foreign	Bupropion			
Other		Myocardial Infarction	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG /			Professional				

TWICE PER DAY

/ ORAL

Metformin  
 Hydrochloride C  
 Conjugated Estrogens C  
 Atorvastatin Calcium C  
 Glibenclamide C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/19/01ISR Number: 3742818-4Report Type:Expedited (15-DaCompany Report #A0090299A  
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Pain Chest Pain	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
100 MG / PER DAY / ORAL	7 DAY	Pericarditis Vomiting					

Date:06/19/01ISR Number: 3742819-6Report Type:Expedited (15-DaCompany Report #A0105877A  
Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Confusional State Cyanosis	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY / ORAL		Dyspnoea Eye Rolling Movement Disorder Sedation		Sumatriptan Succinate Propranolol Hydrochloride Salbutamol Sulphate Triquilar	C C C C		

Date:06/19/01ISR Number: 3742988-8Report Type:Expedited (15-DaCompany Report #A0105423A  
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Disability PER DAY ORAL		Multiple Sclerosis Sensory Loss	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL



Date:06/19/01ISR Number: 3743006-8Report Type:Expedited (15-DaCompany Report #A0075655A  
Age:49 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL	Amnesia  Convulsion  Dyspnoea Grand Mal Convulsion Status Epilepticus Suicidal Ideation	Foreign  Consumer	Zyban   Calcium Salt Conjugated Estrogens Hypericum Ascorbic Acid	PS   C C C C	Glaxo Wellcome Inc	ORAL

Date:06/19/01ISR Number: 3743007-XReport Type:Expedited (15-DaCompany Report #A0082610A  
Age:55 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG PER Initial or Prolonged DAY ORAL	Anaphylactoid Reaction  Angioneurotic Oedema  Dermatitis Pruritus Rash Erythematous Tongue Oedema	Foreign  Health  Professional	Zyban   Thyroxine Sodium Hrt	PS   C C	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/19/01ISR Number: 3743009-3Report Type:Expedited (15-DaCompany Report #A0096983A

Age:23 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 100 MG ORAL	Convulsion	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged Other	Headache		Ibuprofen Semisodium Valproate	C C		

Date:06/19/01ISR Number: 3743234-1Report Type:Expedited (15-DaCompany Report #2001SUS0407

Age:0 DY Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration TRANSPLACENTAL TR	Complications Of Maternal Exposure To Therapeutic Drugs Cystic Fibrosis Meconium Ileus	Health Professional	Sustiva  Combivir (Lamivudine/Zidovudine)	PS  SS	Dupont Pharmaceuticals Co	
TRANSPLACENTAL TR			Norvir (Ritonavir)	SS		
TRANSPLACENTAL TR			Crixivan (Indinavir)	SS		
TRANSPLACENTAL TR			Bupropion (Amfebutamone)	SS		
TRANSPLACENTAL TR			Viramune (Nevirapine)	SS		
TRANSPLACENTAL TR			Epivir (Lamivudine)	SS		
TRANSPLACENTAL TR			Zerit (Stavudine)	SS		
TRANSPLACENTAL TR			Viracept (Nelfinavir)	SS		

Date:06/19/01ISR Number: 3743704-6Report Type:Expedited (15-DaCompany Report #B0110444A

Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia	Foreign	Bupropion			
Hospitalization -		Coma	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL 7	DAY						
Initial or Prolonged		Malaise	Professional	Fluticasone			
		Nausea		Propionate	C		
		Pulmonary Embolism		Salmeterol Xinafoate	C		

Date:06/20/01ISR Number: 3742240-0Report Type:Expedited (15-DaCompany Report #A0150739A  
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Bipolar Disorder		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Convulsion		Zyban	SS	Glaxo Wellcome	ORAL
2	MON						
		Psychotic Disorder					
		Sexual Offence					
		Therapeutic Response					
		Unexpected					

Date:06/20/01ISR Number: 3742247-3Report Type:Expedited (15-DaCompany Report #B0108775A  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT
Death		Cardiac Arrest
Hospitalization -		Confusional State
Initial or Prolonged		Dysarthria
		Hallucination, Visual
		Overdose

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
5MG Three times per day		Tremor Vomiting		Bupropion Hcl Nifedipine	PS C	Glaxo Wellcome	ORAL ORAL

Date:06/20/01ISR Number: 3742248-5Report Type:Expedited (15-DaCompany Report #B0109628A  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 1 DAY		Cardiac Arrest  Confusional State Dysarthria Hallucination, Visual Overdose Tremor Vomiting		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/20/01ISR Number: 3743639-9Report Type:Expedited (15-DaCompany Report #A0083661A  
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG /  TWICE PER DAY  / ORAL		Convulsion  Eye Rolling  Hypoglycaemia	Foreign  Health  Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Human Short-Act. Insulin (Formulation Unknown) (Human Short-Act. Insulin) Human Int/Long Insulin Ethanol	SS   C C		

Date:06/20/01ISR Number: 3743641-7Report Type:Expedited (15-DaCompany Report #B0110591A  
Age:69 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
300 MG / PER							

DAY / ORAL

Fluticasone+Salmeterol (Formulation Unknown) (Fluticasone+Salmeterol) SS

RESPIRATORY

(INHALATION) 2 / TWICE PER

DAY / INHALED

Date:06/20/01ISR Number: 3743642-9Report Type:Expedited (15-DaCompany Report #B0099946A  
Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Conductive Deafness	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							

DAY / ORAL

Conjunctivitis Health  
Ear Disorder Professional

SEE TEXT

Influenza Like Illness Paracetamol C  
Insomnia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/20/01ISR Number: 3743643-0Report Type:Expedited (15-DaCompany Report #A0106475A  
Age:33 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY  / ORAL	Urticaria	Foreign  Health  Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/20/01ISR Number: 3743644-2Report Type:Expedited (15-DaCompany Report #A0090923A  
Age:51 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150 MG / Hospitalization - TWICE PER DAY Initial or Prolonged / ORAL	Alanine Aminotransferase  Increased  Alcohol Withdrawal  Syndrome Aspartate Aminotransferase Increased Blood Alkaline Phosphatase Increased Coma Convulsion Gamma-Glutamyltransferase Increased Hypoxia Pneumonia Aspiration	Foreign  Health  Professional  Company Representative	Zyban      Ethanol (Formulation Unknown) (Alcohol)	PS      SS	Glaxo Wellcome Inc	ORAL

Date:06/20/01ISR Number: 3743645-4Report Type:Expedited (15-DaCompany Report #A0087951A  
Age:24 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required 150 MG /	Blood Glucose Decreased	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Intervention to Convulsion Health  
 TWICE PER DAY  
 Prevent Permanent Fall Professional  
 / ORAL  
 Impairment/Damage

Insulin Lispro C  
 Human Int/Long  
 Insulin C  
 Short-Acting Insulin C

Date:06/20/01ISR Number: 3743646-6Report Type:Expedited (15-DaCompany Report #A0075616A  
 Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypertension	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /			Health				
TWICE PER DAY			Professional				
/ ORAL				Ramipril	C		
				Lorazepam	C		

Date:06/20/01ISR Number: 3743648-XReport Type:Expedited (15-DaCompany Report #A0103586A  
 Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Coma	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
300 MG/ PER							
Initial or Prolonged		Grand Mal Convulsion					
DAY/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/20/01ISR Number: 3743655-7Report Type:Expedited (15-DaCompany Report #A0150726A

Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Bradycardia	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							
Intervention to Prevent Permanent Impairment/Damage		Coma Oliguria Suicide Attempt		Diltiazem Hydrochloride Capsule (Diltiazem Hydrochloride)	SS		ORAL
300 MG/ ORAL							
.5 MG / ORAL				Clonazepam Tablet (Cloazepam)	SS		ORAL
50 MG / ORAL				Pethidine Hydrochloride (Meperidine Hydrochloride)	SS		ORAL
750 MG / ORAL				Desipramine Tablet (Desipramine)	SS		ORAL
10 MG / ORAL				Loxapine (Formulation Unknown) (Loxapine)	SS		ORAL
30 MG / ORAL				Morphine Sulphate (Formulation Unknown) (Morphine Sulfate)	SS		ORAL
50 MG / ORAL				Trazodone Tablet (Trazodone)	SS		ORAL
7.5 MG / ORAL				Zopiclone (Formulation Unknown) (Zopiclone)	SS		ORAL

Date:06/20/01ISR Number: 3743656-9Report Type:Expedited (15-DaCompany Report #A0104181A

Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Required Intervention to 150 MG / Prevent Permanent TWICE PER DAY Impairment/Damage / ORAL	16 DAY	Arthralgia Tendon Disorder  Urticaria	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Ortho 1/35	C		

Date:06/20/01ISR Number: 3743657-0Report Type:Expedited (15-DaCompany Report #A0151352A  
Age:21 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 15 G/ SINGLE Prevent Permanent DOSE / ORAL Impairment/Damage		Electrocardiogram Qrs Complex Prolonged  Overdose	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Lorazepam	C		

Date:06/20/01ISR Number: 3743658-2Report Type:Expedited (15-DaCompany Report #A0151323A  
Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to ORAL Prevent Permanent Impairment/Damage 200 MG / ORAL		Abdominal Pain Angina Pectoris  Drug Interaction	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Fenofibrate Capsule (Fenofibrate)	SS		ORAL



Other	Convulsion	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
SEE TEXT/ORAL						
	Sinus Tachycardia		Citalopram			
	Suicide Attempt		Hydrobromide Tablet	SS		ORAL
SEE TEXT/ORAL						
	Tremor					

Date:06/20/01ISR Number: 3743758-7Report Type:Expedited (15-DaCompany Report #B0111327A  
Age:45 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG/ORAL	Cardiomyopathy	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:06/20/01ISR Number: 3743760-5Report Type:Expedited (15-DaCompany Report #A0106203A  
Age:40 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required 100 MG/PER Intervention to DAY/ORAL 1 MON Prevent Permanent Impairment/Damage	Abdominal Pain  Proctitis Rectal Haemorrhage Rectal Tenesmus Vomiting	Foreign	Wellbutrin Sr   Betamehtasone Na Po4 Trazodone Hydrochloride Azathioprine Diclofenac Sodium	PS   C C C C	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/20/01ISR Number: 3743818-0Report Type:Expedited (15-DaCompany Report #A0103160A  
Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Pneumonia	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG /			Health				
TWICE PER DAY			Professional				
/ ORAL				Nifedipine	C		

Date:06/20/01ISR Number: 3743832-5Report Type:Expedited (15-DaCompany Report #B0111329A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Foreign	Bupropion	PS	Glaxo Wellcome Inc	ORAL
150 MG/ORAL		Stress	Health	Hydrochloride			
			Professional				

Date:06/20/01ISR Number: 3743833-7Report Type:Expedited (15-DaCompany Report #A0150947A  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Body Temperature	Foreign	Bupropion	PS	Glaxo Wellcome Inc	ORAL
Intervention to		Increased		Hydrochloride			
150 MG/TWICE							
Prevent Permanent		Chest Pain					
PER DAY/ ORAL 18	DAY	Face Oedema		Atoravastin Calcium	C		
Impairment/Damage		Periorbital Oedema					
		Pharyngolaryngeal Pain					
		Urticaria					
		White Blood Cell Disorder					

Date:06/20/01ISR Number: 3743834-9Report Type:Expedited (15-DaCompany Report #A0143251A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/TWICE		Convulsion Loss Of Consciousness	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ORAL				Naproxen Omeprazole Aspirin Amlodipine Trazodone Medroxyprogesterone Ace. Bromazepam Sertraline Hydrochloride Fiorinal Percocet	C C C C C C C C C C C C		

Date:06/20/01ISR Number: 3743835-0Report Type:Expedited (15-DaCompany Report #A0151353A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required ORAL		Agranulocytosis	Foreign	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
Intervention to Prevent Permanent Impairment/Damage 2 MG/ORAL		Infection		Clonazepam (Formulation Unknown)	SS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/21/01ISR Number: 3743451-0Report Type:Expedited (15-DaCompany Report #A0110910A

Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death				Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Abdominal Pain					
Life-Threatening		Cardiac Arrest					
per day	15	DAY					
Disability		Idiopathic Thrombocytopenic Purpura		Pain Medication (Unspecified)	C		

Date:06/21/01ISR Number: 3743465-0Report Type:Expedited (15-DaCompany Report #A0151572A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other				Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice		Cardiovascular Disorder					
per day				Cardiovascular Medication	SS		

Date:06/21/01ISR Number: 3743469-8Report Type:Expedited (15-DaCompany Report #A0151751A

Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other				Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Ectopic Pregnancy					
per day							

Date:06/21/01ISR Number: 3744003-9Report Type:Expedited (15-DaCompany Report #A103444

Age:79 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required			Consumer	Zoloft	PS	Pfizer	
		Blood Pressure Increased					

Intervention to	Drug Ineffective		Pharmaceuticals Inc
50.00 MG			
Prevent Permanent	Ecchymosis		
TOTAL:DAILY			
Impairment/Damage	Haemorrhagic Stroke	Wellbutrin	SS ORAL
300.000			
	Head Injury		
TOTAL:BID:ORA			
	Headache		
L			
	Heart Rate Decreased	Coumadin	SS
	Heart Rate Increased	Effexor Sr	SS ORAL
300.00 MG			
	Hyperhidrosis		
TOTAL:BID:ORA			
	Laboratory Test Abnormal		
L			
	Medication Error	Toprol	C
	Overdose	Zebeta	C
	Syncope	Lanoxin	C
		Lasix	C
		Synthroid	C
		Vitamin E	C
		Estrogen	C

Date:06/22/01ISR Number: 3744090-8Report Type:Expedited (15-DaCompany Report #A0149803A  
Age:29 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Agitation
Initial or Prolonged	Alcoholism
Other	Catatonia
	Coma
	Convulsion
	Depression
	Hallucination

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Intentional Misuse Tachycardia Toxicologic Test Abnormal	Report Source	Product	Role	Manufacturer	Route
7.5G	See text	Vomiting	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
				Amphetamine	C		
				Cannabis	C		

Date:06/22/01ISR Number: 3744094-5Report Type:Expedited (15-DaCompany Report #A0152007A  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coronary Artery Disease		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
450MG	Per day	YR		Statins	C		

Date:06/22/01ISR Number: 3745460-4Report Type:Expedited (15-DaCompany Report #A0130455A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Congenital Anomaly		Asthenia	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL				Fluvoxamine Tablet (Fluvoxamine)	SS		ORAL
		Dysarthria Gait Disturbance					
		Hypothermia					
100 MG / PER							
DAY / ORAL							

Date:06/22/01ISR Number: 3745462-8Report Type:Expedited (15-DaCompany Report #A0114129A  
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Asthenia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /TWICE							
Other		Chest Pain	Health				
PER DAY /							



ORAL

Coordination Abnormal

Professional

Cough  
Cyanosis  
Disorientation  
Dizziness  
Fatigue  
Feeling Cold  
Feeling Hot  
Feeling Jittery  
Headache  
Hearing Impaired  
Insomnia  
Neck Pain  
Pallor  
Paraesthesia  
Pollakiuria  
Poverty Of Thought  
Content  
Sneezing  
Tinnitus  
Visual Disturbance

Diuretic

C

Date:06/22/01ISR Number: 3745463-XReport Type:Expedited (15-DaCompany Report #A0086224A

Age:52 YR Gender:Male I/FU:F

Outcome	PT	Report Source
Death	Myocardial Infarction	Foreign Literature

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Health Professional	Product	Role	Manufacturer	Route
150 MG /			Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY						
/ ORAL	2 DAY					

Date:06/22/01ISR Number: 3745495-1Report Type:Expedited (15-DaCompany Report #A114068  
 Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged	300.00 MG	Amnesia Cardio-Respiratory Arrest Coma	Foreign Consumer	Viagra	PS	Pfizer Agricultural Div	
TOTAL: BID: ORA		Convulsion		Bupropion	SS		ORAL
L		Drug Interaction					
0.40 MG		Fall		Tamsulosin	SS		ORAL
TOTAL: DAILY: O		Head Injury					
RAL		Tachycardia					
				Atorvastatin	C		
				Lorazepam	C		
				Aspirin	C		
				Dimethyl Sulfone	C		
				Ubidecarenone	C		
				Multivitamin	C		
				Fish Oil	C		
				Pentanic Acid	C		

Date:06/22/01ISR Number: 3749762-7Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #A0124951A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/UNK/		Convulsion					
ORAL		Epileptic Aura					
		Fatigue					
		Loss Of Consciousness					
		Malaise					
		Muscle Tightness					
		Neck Pain					
		Overdose					
		Speech Disorder					
		Syncope					

Date:06/22/01ISR Number: 3749763-9Report Type:Periodic Company Report #A0124945A  
Age:26 YR Gender:Female I/FU:F

Outcome	PT
Other	Arthralgia
	Candidiasis
	Decreased Activity
	Difficulty In Walking
	Erythema
	Face Oedema
	Insomnia
	Muscle Spasms
	Oedema Peripheral
	Pruritus

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Urticaria

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE		Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ORAL			Medroxyprogesterone Ace.	C		

Date:06/22/01ISR Number: 3749764-0Report Type:Periodic Company Report #A0123486A  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Other		Back Pain					
150 MG/ TWICE		Convulsion					
PER DAY/ ORAL		Dysphagia		One-A-Day Vitamins	C		
		Dyspnoea					
		Nausea					
		Speech Disorder					
		Throat Tightness					
		Tremor					

Date:06/22/01ISR Number: 3749765-2Report Type:Periodic Company Report #A0057814A  
 Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
Disability		Agitation					
150 MG/TWICE		Amnesia	Professional				
PER DAY/ ORAL		Arthralgia		Diazepam			
		Bronchitis		(Formulation Unknown)			
		Confusional State		(Diazepam)	SS		
		Cough		Ethanol (Formulation			
		Decreased Activity		Unknown) (Alcohol)	SS		
		Depression		Conjugated Estrogens	C		
		Difficulty In Walking		Amitriptyline Hcl	C		

Dry Mouth  
Dyspraxia  
Grand Mal Convulsion  
Hallucination  
Headache  
Nervousness  
Speech Disorder  
Tremor

Decongestant C  
Medroxyprogesterone  
Ace. C  
Nabumetone C  
Aspirin C

Date:06/22/01ISR Number: 3749766-4Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #A0145488A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Blood Pressure Increased					
PER DAY/ ORAL		Erythema					
		Flushing					
		Hypersensitivity					
		Hyperventilation					
		Paraesthesia					
		Pruritus					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/22/01ISR Number: 3749767-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0145142A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL	9 DAY	Difficulty In Walking	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Face Oedema Hypersensitivity Pain Throat Tightness Urticaria					

Date:06/22/01ISR Number: 3749768-8Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #A0145079A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ TWICE		Convulsion	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged PER DAY/ORAL							

Date:06/22/01ISR Number: 3749769-XReport Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #A0144963A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG/ TWICE		Dermatitis	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL		Hypersensitivity Pruritus Vomiting					

Date:06/22/01ISR Number: 3749770-6Report Type:Periodic  
 Age:27 YR Gender:Male I/FU:I

Company Report #A0142980A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Circulatory Collapse	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE	Convulsion					
PER DAY/ ORAL	Disorientation					
	Memory Impairment					
	Tongue Disorder					

Date:06/22/01ISR Number: 3749771-8Report Type:Periodic Company Report #A0142186A  
 Age:37 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Arthralgia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE	Burning Sensation	Professional				
PER DAY/ ORAL	Dry Skin					
	Dyspnoea					
	Fatigue					
	Myalgia					
	Pain					
	Pruritus					
	Rash Scaly					
	Serum Sickness					
	Swelling					
	Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/22/01ISR Number: 3749772-XReport Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0130468A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1 TABLET/ Initial or Prolonged TWICE PER DAY/ORAL	Anaphylactic Shock Dermatitis Dyspnoea	Health Professional Company Representative	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/22/01ISR Number: 3749773-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0130291A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ ORAL Initial or Prolonged	Dysgeusia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/22/01ISR Number: 3749774-3Report Type:Periodic  
Age:33 YR Gender:Male I/FU:I

Company Report #A0129542A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL	Bite Dizziness Grand Mal Convulsion Head Injury Loss Of Consciousness Tremor	Health Professional	Zyban Pravastatin Sodium Nicotine	PS C C	Glaxo Wellcome Inc	ORAL

Date:06/22/01ISR Number: 3749786-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0139615A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						



Other Convulsion Health Zyban PS Glaxo Wellcome Inc ORAL  
150 MG /  
TWICE PER DAY Professional  
/ ORAL Company  
Representative

Date:06/22/01ISR Number: 3749787-1Report Type:Periodic Company Report #A0138575A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Feeling Abnormal					
TWICE PER DAY		Muscle Twitching					
/ ORAL							

Date:06/22/01ISR Number: 3749788-3Report Type:Periodic Company Report #A0137592A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /		Road Traffic Accident					
TWICE PER DAY							
/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/22/01ISR Number: 3749789-5Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0137115A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG / TWICE PER DAY / ORAL		Blood Pressure Increased Confusional State Convulsion Disorientation Muscle Rigidity Pain Panic Reaction Rhinorrhoea Salivary Hypersecretion Tremor Urinary Incontinence Visual Acuity Reduced	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/22/01ISR Number: 3749790-1Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #A0135779A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Disability		Convulsion	Consumer	Zyban	PS	Glaxo Wellcome Inc	

Date:06/22/01ISR Number: 3749791-3Report Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #A0134397A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG / SINGLE DOSE / ORAL		Amnesia Grand Mal Convulsion Overdose	Health Professional Company Representative	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Semisodium Valproate Clonazepam	C C		

Olanzapine

C

Date:06/22/01ISR Number: 3749792-5Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #A0132271A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / ORAL	Asthenia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged	Convulsion		Oxygen	C		
	Difficulty In Walking		Salbutamol Sulphate	C		
	Dizziness		Ipratropium Bromide	C		
	Dyskinesia		Nebulizer	C		
	Tremor		Diazepam	C		
			Primidone	C		

Date:06/22/01ISR Number: 3749793-7Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #A0131564A

Outcome	PT
Hospitalization -	Dermatitis
Initial or Prolonged	Dyspepsia
	Dyspnoea
	Face Oedema
	Hypersensitivity
	Nausea

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Oedema Peripheral Pruritus Urticaria	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:06/22/01ISR Number: 3749794-9Report Type:Periodic Company Report #A0131413A  
Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
Other		Agranulocytosis	Professional				
150 MG /		Neutropenia	Company				
TWICE PER DAY			Representative				
/ ORAL							

Date:06/22/01ISR Number: 3749795-0Report Type:Periodic Company Report #A0131272A  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
Other		Convulsion	Professional				
150 MG /							
TWICE PER DAY							
/ ORAL							

Date:06/22/01ISR Number: 3749796-2Report Type:Periodic Company Report #A0128124A  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Life-Threatening Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL Disability Other Required Intervention to Prevent Permanent Impairment/Damage	Cardiac Failure Congestive  Chest Pain  Convulsion Myocardial Infarction	Health Professional	Bupropion Hydrochloride          Nicotine	PS          C	Glaxo Wellcome Inc	ORAL
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Date:06/22/01ISR Number: 3749797-4Report Type:Periodic Company Report #A0127489A  
 Age:80 YR Gender:Male I/FU:I

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150 MG/ ORAL Required Intervention to Prevent Permanent Impairment/Damage	Dermatitis  Hypersensitivity Pruritus Rash Macular Skin Hypopigmentation Urticaria	Health  Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/22/01ISR Number: 3749798-6Report Type:Periodic Company Report #A0127208A  
 Age:55 YR Gender:Male I/FU:I

Outcome Hospitalization - Initial or Prolonged	PT Chest Pain Depressed Level Of Consciousness
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Gastrooesophageal Reflux  
Disease

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ ORAL		Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Multivitamin	C		

Date:06/22/01ISR Number: 3749799-8Report Type:Periodic Company Report #A0127038A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Convulsion	Health  Professional Company Representative	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/22/01ISR Number: 3749800-1Report Type:Periodic Company Report #A0126954A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Convulsion	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:06/22/01ISR Number: 3749801-3Report Type:Periodic Company Report #A0126757A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG/ TWICE PER DAY/ ORAL	5 WK	Amnesia  Grand Mal Convulsion  Petit Mal Epilepsy	Health  Professional	Zyban  Prednisone Diazepam	PS  SS SS	Glaxo Wellcome Inc	ORAL

Date:06/22/01ISR Number: 3749802-5Report Type:Periodic Company Report #A0126289A  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Accident	Consumer	Zyban	PS	Glaxo Wellcome Inc	

Date:06/22/01ISR Number: 3749803-7Report Type:Periodic Company Report #A0126220A  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization - Initial or Prolonged		Confusional State Ecchymosis Excoriation Grand Mal Convulsion Loss Of Consciousness Weight Increased	Health Professional	Zyban Conjugated Estrogens	PS C	Glaxo Wellcome Inc	

Date:06/22/01ISR Number: 3749804-9Report Type:Periodic Company Report #A0126219A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL				Phenobarbitone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/22/01ISR Number: 3749805-0Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0125898A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL		Difficulty In Walking	Professional				
		Eyelid Oedema		Thyroxine Sodium	C		
		Joint Stiffness					
		Pruritus					
		Rash Erythematous					
		Swelling					
		Urticaria					

Date:06/22/01ISR Number: 3749806-2Report Type:Periodic  
Age:51 YR Gender:Male I/FU:I

Company Report #A0125812A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Bite	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
Initial or Prolonged		Convulsion					
PER DAY/ ORAL							
		Fall		Lisinopril	C		
		Head Injury		Atorvastatin Calcium	C		
		Loss Of Consciousness					
		Movement Disorder					
		Tongue Disorder					

Date:06/22/01ISR Number: 3749807-4Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #A0125320A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Depersonalisation	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
Initial or Prolonged		Dyspnoea					
PER DAY/ ORAL 1 WK							
		Hyperhidrosis					
		Malaise					
		Nervousness					



Pallor  
Syncope  
Tachycardia

Date:06/25/01ISR Number: 3745249-6Report Type:Expedited (15-DaCompany Report #A0137061A  
Age:17 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Sinus Tachycardia		Celexa	SS		ORAL
400MG per day						
Other	Suicide Attempt					
	Tremor					

Date:06/25/01ISR Number: 3745252-6Report Type:Expedited (15-DaCompany Report #A0149863A  
Age:60 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Other	Dry Mouth	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice						
	Dry Throat					
per day						
	Emphysema					
	Speech Disorder					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/25/01ISR Number: 3745254-XReport Type:Expedited (15-DaCompany Report #A0151556A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Valtrex	PS	Glaxo Wellcome	ORAL
		Drug Interaction		Wellbutrin	SS	Glaxo Wellcome	ORAL

Date:06/25/01ISR Number: 3746187-5Report Type:Expedited (15-DaCompany Report #A0150739A

Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bipolar Disorder	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
ORAL		Sexual Offence	Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	SS		ORAL
ORAL	2	MON					

Date:06/25/01ISR Number: 3746425-9Report Type:Expedited (15-DaCompany Report #A0151751A

Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ectopic Pregnancy	Foreign Study	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG, TWICE			Consumer				
PER DAY, ORAL							

Date:06/25/01ISR Number: 3746430-2Report Type:Expedited (15-DaCompany Report #B0109628A

Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardiac Arrest	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ORAL		Confusional State					
		Dysarthria					

Hallucination  
Overdose  
Tremor  
Vomiting

Date:06/25/01ISR Number: 3746431-4Report Type:Expedited (15-DaCompany Report #B0108775A  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Foreign	Bupropion			
Hospitalization - 150 MG, ORAL		Confusional State		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Dysarthria Hallucination Overdose Tremor Vomiting		Nifedipine	C		

Date:06/25/01ISR Number: 3746458-2Report Type:Expedited (15-DaCompany Report #A0110910A  
Age:33 YR Gender:Female I/FU:F

Outcome	PT
Death	Abdominal Pain Upper
Life-Threatening	Blood Disorder
Disability	Cardiac Arrest Idiopathic

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Thrombocytopenic Purpura  
Influenza

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG /		Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
TWICE PER DAY		Health				
/ ORAL		Professional				
			Pain Medication	C		

Date:06/25/01ISR Number: 3747190-1Report Type:Expedited (15-DaCompany Report #A0151572A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Cardiovascular Disorder	Professional				
150 MG/ TWICE			Company				
PER DAY/ ORAL			Representative				

Date:06/26/01ISR Number: 3746299-6Report Type:Expedited (15-DaCompany Report #A0145968A  
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxo Wellcome	ORAL
Hospitalization -		Asthenia					
150MG Per day 3 DAY		Decreased Appetite		Unknown			
Initial or Prolonged		Difficulty In Walking		Anti-Depressant	C		
		Dysphagia					
		Hypersensitivity					

Date:06/26/01ISR Number: 3746305-9Report Type:Expedited (15-DaCompany Report #B0100202A  
Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Hypertensive Crisis Health Zyban PS Glaxo Wellcome  
 150MG Twice  
 Initial or Prolonged Paraesthesia Professional  
 per day 8 DAY  
 Transient Ischaemic Aspirin C  
 150MG In the  
 morning Attack  
 Hrt C

Date:06/26/01ISR Number: 3746307-2Report Type:Expedited (15-DaCompany Report #B0109587A  
 Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 10.5G		Grand Mal Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged cumulative dose		Hypothermia Intentional Misuse Nausea Pyrexia Tachycardia Vomiting					

Date:06/26/01ISR Number: 3746311-4Report Type:Expedited (15-DaCompany Report #B0111705A  
 Age:51 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Depressed Level Of Consciousness Muscular Weakness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nystagmus

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
WK			Zyban	PS	Glaxo Wellcome	ORAL
			Proton Pump Inhibitor	C		

Date:06/26/01ISR Number: 3746313-8Report Type:Expedited (15-DaCompany Report #B0111849A  
Age:69 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest Circulatory Collapse		Zyban	PS	Glaxo Wellcome	

Date:06/26/01ISR Number: 3746314-XReport Type:Expedited (15-DaCompany Report #B0111850A  
Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Zyban	PS	Glaxo Wellcome	

Date:06/26/01ISR Number: 3746315-1Report Type:Expedited (15-DaCompany Report #B0111851A  
Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Fall		Zyban Sildenafil Alcohol	PS C C	Glaxo Wellcome	

Date:06/26/01ISR Number: 3746316-3Report Type:Expedited (15-DaCompany Report #B0111855A  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Angiopathy		Zyban	PS	Glaxo Wellcome	

Myocardial Infarction  
Tachycardia  
With Nerve Paralysis

Date:06/26/01ISR Number: 3746318-7Report Type:Expedited (15-DaCompany Report #B0111862A  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	7 DAY		Cerebral Artery Occlusion	Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged			Cerebral Artery Thrombosis Ischaemic Stroke Oedema				

Date:06/26/01ISR Number: 3746320-5Report Type:Expedited (15-DaCompany Report #B0111882A  
Age: Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Convulsion Hypoglycaemia Intentional Misuse Renal Failure Acute

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Rhabdomyolysis

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
6G cumulative dose		Consumer	Zyban	PS	Glaxo Wellcome	ORAL
UNKNOWN cumulative dose	20TAB		Co-Dydramol	SS	Glaxo Wellcome	
UNKNOWN cumulative dose	15TAB		Bezafibrate	SS		
UNKNOWN cumulative dose	15TAB		Gliclazide	SS		
RESPIRATORY (INHALATION)			Salbutamol	C	Glaxo Wellcome	
			Alcohol	C		

Date:06/26/01ISR Number: 3746321-7Report Type:Expedited (15-DaCompany Report #B0111884A  
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - Initial or Prolonged		Joint Effusion		Zyban	PS	Glaxo Wellcome	

Date:06/26/01ISR Number: 3746322-9Report Type:Expedited (15-DaCompany Report #B0111886A  
Age:40 YR Gender:Male I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Chest Pain		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Hyponatraemia					
per day	7	DAY		Atenolol	C		ORAL

Date:06/26/01ISR Number: 3746323-0Report Type:Expedited (15-DaCompany Report #B0111947A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Bundle Branch Block Right		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Chest Pain					
Initial or Prolonged		Dyspnoea		Zopiclone	C		
per day	7	Palpitations		Propiomazine	C		
		Pulmonary Embolism		Somadril	C		

Date:06/26/01ISR Number: 3746324-2Report Type:Expedited (15-DaCompany Report #B0111954A  
Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coma		Zyban	PS	Glaxo Wellcome	
150MG Per day		Haemorrhagic Stroke		Salbutamol	C	Glaxo Wellcome	
Hospitalization -				Diclofenac	C		
Initial or Prolonged							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/26/01ISR Number: 3748136-2Report Type:Expedited (15-DaCompany Report #A0149803A

Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	7.5 G/SEE	Agitation Catatonia	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Other TEXT/ORAL		Coma					
		Confusional State		Amphetamine	C		
		Convulsion		Cannabis	C		
		Depression					
		Gait Disturbance					
		Hallucination					
		Intentional Misuse					
		Sedation					
		Tachycardia					
		Vomiting					

Date:06/26/01ISR Number: 3748210-0Report Type:Direct

Company Report #

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hallucination		Wellbutrin	PS		

Date:06/26/01ISR Number: 3748455-XReport Type:Expedited (15-DaCompany Report #A0152007A

Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 450 MG / PER		Coronary Artery Disease	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
DAY / ORAL	YR		Professional				
				Statins	C		

Date:06/27/01ISR Number: 3747617-5Report Type:Expedited (15-DaCompany Report #B0109706A

Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Anxiety	Health	Zyban	PS	Glaxo Wellcome	ORAL
6 WK							
Hospitalization - Initial or Prolonged Disability		Haemorrhagic Stroke Ruptured Cerebral Aneurysm	Professional				
Other		Status Epilepticus					

Date:06/27/01ISR Number: 3747619-9Report Type:Expedited (15-DaCompany Report #B0111860A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Effect Decreased	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	11 DAY	Drug Interaction					
350MCG Per		Pregnancy On Oral		Micronor	SS		ORAL
day	412 DAY	Contraceptive					
500MCG Weekly				Cabergoline	C		ORAL

Date:06/27/01ISR Number: 3747620-5Report Type:Expedited (15-DaCompany Report #B0111950A  
Age:35 YR Gender:Male I/FU:I

Outcome  
Hospitalization -  
Initial or Prolonged  
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Disability

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Panic Attack		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/27/01ISR Number: 3747624-2Report Type:Expedited (15-DaCompany Report #B0112183A  
Age:49 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Twice		Cerebrovascular Accident		Zyban	PS	Glaxo Wellcome	ORAL
	per day		Facial Palsy					
	20MG At night				Atorvastatin	C		ORAL
	2.5MG Per day				Bendrofluazide	C	Glaxo Wellcome	ORAL

Date:06/27/01ISR Number: 3747625-4Report Type:Expedited (15-DaCompany Report #B0112188A  
Age:55 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day	12 DAY	Anxiety		Zyban	PS	Glaxo Wellcome	ORAL
			Chest Pain					
			Hyperhidrosis					
			Suicide Attempt					
			Visual Disturbance					

Date:06/27/01ISR Number: 3747626-6Report Type:Expedited (15-DaCompany Report #B0112285A  
Age:47 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day	4 DAY	Abdominal Pain		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged			Cerebellar Infarction					

Cerebrovascular Disorder  
Coordination Abnormal  
Dizziness

Date:06/27/01ISR Number: 3747627-8Report Type:Expedited (15-DaCompany Report #B0112291A  
Age:63 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death		Circulatory Collapse	Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day 2 DAY						
Hospitalization -		Renal Failure	Loperamide	C		ORAL
2MG Per day						
Initial or Prolonged			Allopurinol	C	Glaxo Wellcome	ORAL
300MG Per day						
			Meloxicam	C		ORAL
7.5MG Per day						
			Salbutamol	C	Glaxo Wellcome	
RESPIRATORY						
(INHALATION)	100MCG As					
required						
			Temazepam	C		ORAL
10MG Per day						

Date:06/27/01ISR Number: 3747628-XReport Type:Expedited (15-DaCompany Report #B0112293A  
Age:45 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Balance Disorder
Initial or Prolonged	Hypoaesthesia
Other	Malaise
	Monoparesis
	Tearfulness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Transient Ischaemic Attack				
Dose	Duration		Report Source	Product	Role	Manufacturer
150MG Per day	2 DAY			Zyban	PS	Glaxo Wellcome
						ORAL

Date:06/27/01ISR Number: 3749369-1Report Type:Expedited (15-DaCompany Report #A0145968A  
Age:54 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - 150 MG/ PER	Asthenia	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged DAY/ ORAL	Burning Sensation					
	Difficulty In Walking		Antidepressant	C		
	Dysphagia					
	Dyspnoea					
	Hypersensitivity					
	Oral Intake Reduced					

Date:06/27/01ISR Number: 3749726-3Report Type:Expedited (15-DaCompany Report #B0100202A  
Age:59 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - 150 MG/TWICE	Hypertensive Crisis	Foreign	Zyban	PS	Glaxo Wellcome Inc	
Initial or Prolonged PER	Transient Ischaemic	Health				
DAY/UNKNOWN	Attack	Professional				
			Aspirin	C		
			Hrt	C		

Date:06/27/01ISR Number: 3749732-9Report Type:Expedited (15-DaCompany Report #B0111851A  
Age:45 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					

Death	Fall	Foreign	Zyban	PS	Glaxo Wellcome Inc
150 MG					
		Health Professional	Sildenafil Citrate	C	
			Ethanol	C	

Date:06/27/01ISR Number: 3749733-0Report Type:Expedited (15-DaCompany Report #B0111947A  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY		Atrioventricular Block	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Chest Pain					
		Dyspnoea					
		Palpitations		Zopiclone	C		
		Pulmonary Embolism		Propiomazine	C		
				Carisoprodol	C		

Date:06/27/01ISR Number: 3749734-2Report Type:Expedited (15-DaCompany Report #B0111886A  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150 MG / TWICE PER DAY		Chest Pain	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Hyponatraemia					
				Atenolol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/27/01ISR Number: 3749735-4Report Type:Expedited (15-DaCompany Report #B0111882A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / ORAL		Convulsion	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Hypoglycaemia		Co-Dydramol Tablet (Co-Dydramol)	SS		
		Intentional Misuse		Bezafibrate Tablet (Bezafibrate)	SS		
		Renal Failure Acute		Gliclazide Tablet (Gliclazide)	SS		
		Rhabdomyolysis		Salbutamol Sulphate	C		
				Ethanol	C		

Date:06/27/01ISR Number: 3749736-6Report Type:Expedited (15-DaCompany Report #B0111855A

Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG		Cranial Nerve Disorder	Foreign	Zyban	PS	Glaxo Wellcome Inc	
		Diplopia					
		Myocardial Infarction					
		Tachycardia					

Date:06/27/01ISR Number: 3749737-8Report Type:Expedited (15-DaCompany Report #B0111705A

Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / ORAL		Neurological Symptom	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged			Health Professional	Proton Pump Inhibitors	C		

Date:06/27/01ISR Number: 3749738-XReport Type:Expedited (15-DaCompany Report #B0109587A

Age:21 YR Gender:Female I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / ORAL		Grand Mal Convulsion	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Hypothermia Intentional Misuse Nausea Pyrexia Tachycardia Vomiting	Health Professional				

Date:06/27/01ISR Number: 3749899-2Report Type:Expedited (15-DaCompany Report #B0111954A  
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG/ PER		Coma	Foreign	Zyban	PS	Glaxo Wellcome Inc	
Hospitalization - DAY		Haemorrhagic Stroke					
Initial or Prolonged		Headache		Salbutamol Sulphate Diclofenac	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/27/01ISR Number: 3749900-6Report Type:Expedited (15-DaCompany Report #B0111884A  
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Emphysema	Foreign	Bupropion			
Hospitalization - 150 MG		Joint Effusion	Consumer	Hydrochloride	PS	Glaxo Wellcome Inc	
Initial or Prolonged							

Date:06/27/01ISR Number: 3749901-8Report Type:Expedited (15-DaCompany Report #B0111862A  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebral Artery	Foreign	Bupropion			
Hospitalization - 150 MG/ORAL		Thrombosis		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Ischaemic Stroke Pulmonary Oedema					

Date:06/27/01ISR Number: 3749902-XReport Type:Expedited (15-DaCompany Report #B0111849A  
Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Foreign	Zyban	PS	Glaxo Wellcome Inc	
150 MG		Circulatory Collapse					

Date:06/27/01ISR Number: 3749903-1Report Type:Expedited (15-DaCompany Report #B0111850A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Foreign	Zyban	PS	Glaxo Wellcome Inc	
150 MG			Health Professional				

Date:06/28/01ISR Number: 3748535-9Report Type:Expedited (15-DaCompany Report #A0152031A  
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 400MG per day 4 MON	Angiopathy		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Convulsion		Beclovent	C	Glaxo Wellcome	
	Excessive Masturbation		Celexa	C		
	Loss Of Consciousness		Synthroid	C	Glaxo Wellcome	
	Sedation		Tetracycline	C		
	Speech Disorder		Neurontin	C		

Date:06/28/01ISR Number: 3748542-6Report Type:Expedited (15-DaCompany Report #B0110855A  
Age:31 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150MG	Abortion Induced		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Pregnancy					
Variable dose	Venous Thrombosis Limb					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/28/01ISR Number: 3748545-1Report Type:Expedited (15-DaCompany Report #B0111869A  
Age:58 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150MG	Duration Deep Vein Thrombosis		Zyban	PS	Glaxo Wellcome	ORAL
Variable dose 36 DAY						

Date:06/28/01ISR Number: 3748546-3Report Type:Expedited (15-DaCompany Report #B0111956A  
Age:47 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Per day 4 DAY Initial or Prolonged	Duration Cerebrovascular Accident Intestinal Gangrene		Zyban	PS	Glaxo Wellcome	

Date:06/28/01ISR Number: 3748549-9Report Type:Expedited (15-DaCompany Report #B0112406A  
Age:44 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death	Duration Death	Health Professional	Zyntabac	PS	Glaxo Wellcome	

Date:06/28/01ISR Number: 3748550-5Report Type:Expedited (15-DaCompany Report #B0112411A  
Age:54 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day 4 DAY 10MG per day	Duration Chest Pain		Zyban Amlodipine	PS C	Glaxo Wellcome	ORAL ORAL

Date:06/29/01ISR Number: 3749418-0Report Type:Expedited (15-DaCompany Report #A0141089A  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day	Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged		Fall		Alcohol	SS		
		Fracture		Flovent	C	Glaxo Wellcome	
		Stupor		Serevent	C	Glaxo Wellcome	
		Suicide Attempt		Atrovent	C	Glaxo Wellcome	
		Therapeutic Agent		Proventil	C	Glaxo Wellcome	
		Toxicity					

Date:06/29/01ISR Number: 3749424-6Report Type:Expedited (15-DaCompany Report #A0151751A  
Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged per day	150MG Twice 56 DAY	Ectopic Pregnancy Post Procedural Complication		Zyban	PS	Glaxo Wellcome	ORAL

Date:07/02/01ISR Number: 3750354-4Report Type:Expedited (15-DaCompany Report #B0097894A  
Age:58 YR Gender:Male I/FU:F

Outcome	PT
Disability	Asthenopia Condition Aggravated Dizziness Gastrointestinal

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Candidiasis				
		Insomnia				
		Pharyngolaryngeal Pain				
		Throat Tightness	Report Source	Product	Role	Manufacturer
Dose	Duration					Route
150MG Per day	15 DAY			Bupropion Hydrochloride	PS	Glaxo Wellcome
						ORAL

Date:07/02/01ISR Number: 3750355-6Report Type:Expedited (15-DaCompany Report #B0100339A  
Age:66 YR Gender:Female I/FU:F

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose		Anxiety		Zyban	PS	Glaxo Wellcome	
Hospitalization -	3 DAY						
Initial or Prolonged		Headache		Atorvastatin	C		
10MG per day							
2.5MG per day		Hypoaesthesia		Enalapril	C		
		Lethargy					
		Loss Of Consciousness					

Date:07/02/01ISR Number: 3750362-3Report Type:Expedited (15-DaCompany Report #B0112544A  
Age:49 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose		Migraine		Zyban	PS	Glaxo Wellcome	
Other		Subarachnoid Haemorrhage					

Date:07/02/01ISR Number: 3750364-7Report Type:Expedited (15-DaCompany Report #D0017705A  
Age:63 YR Gender:Female I/FU:F

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose		Chemical Poisoning		Zyban	PS	Glaxo Wellcome	ORAL
Death							
150MG Twice		Completed Suicide					
per day	82 DAY						
125MG Per day	YR	Depression Suicidal		L-Thyroxin 125	C	Glaxo Wellcome	ORAL

10MG Per day YR Baymycard C ORAL

Date:07/02/01ISR Number: 3750773-6Report Type:Direct Company Report #  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Wellbutrin	PS		
PT TOOK							
Other							
OVERDOSE ONCE	1 DAY			Benadryl	C		

Date:07/02/01ISR Number: 3750997-8Report Type:Expedited (15-DaCompany Report #A0141089A  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Arthralgia	Health	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							
Hospitalization -		Fall	Professional				
DAY / ORAL							
Initial or Prolonged		Fracture		Fluticasone			
		Intentional Misuse		Propionate	C		
		Restlessness		Salmeterol Xinafoate	C		
		Stupor		Ipratropium Bromide	C		
		Suicide Attempt		Salbutamol Sulphate	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/02/01ISR Number: 3751047-XReport Type:Expedited (15-DaCompany Report #B0110855A  
 Age:31 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150	Abortion Induced Complications Of Maternal	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
MG/VARIABLE DOSE/ ORAL	Exposure To Therapeutic Drugs Venous Thrombosis Limb	Professional				

Date:07/02/01ISR Number: 3751048-1Report Type:Expedited (15-DaCompany Report #B0111956A  
 Age:47 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged UNKNOWN	Cerebrovascular Accident Gait Disturbance	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
150 MG/ PER DAY	Intestinal Ischaemia					

Date:07/02/01ISR Number: 3751049-3Report Type:Expedited (15-DaCompany Report #B0112406A  
 Age:44 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death UNKNOWN	Death	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
150 MG		Professional				

Date:07/02/01ISR Number: 3751269-8Report Type:Expedited (15-DaCompany Report #A0152031A  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						



Hospitalization - ORAL	Angiopathy	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged	Convulsion		Beclomethasone			
	Loss Of Consciousness		Dipropion	C		
	Psychosexual Disorder		Citalopram			
	Speech Disorder		Hydrobromide	C		
			Thyroxine Sodium	C		
			Tetracycline	C		
			Gabapentin	C		

Date:07/02/01ISR Number: 3751398-9Report Type:Expedited (15-DaCompany Report #A0151556A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Valtrex	PS	Glaxo Wellcome Inc	ORAL
ORAL		Drug Interaction	Professional Company Representative	Bupropion Hydrochloride Tablet-Controlled Release (Bupropion Hydrochloride)	SS		ORAL
ORAL							

Date:07/02/01ISR Number: 3751432-6Report Type:Expedited (15-DaCompany Report #A0151751A

Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Ectopic Pregnancy Post Procedural Complication	Foreign Study Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/02/01ISR Number: 3751635-0Report Type:Expedited (15-DaCompany Report #B0112188A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Foreign	Bupropion			
300 MG/PER		Chest Pain		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
DAY/ORAL		Hyperhidrosis					
		Suicide Attempt					
		Visual Disturbance					

Date:07/02/01ISR Number: 3751639-8Report Type:Expedited (15-DaCompany Report #B0112291A  
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Circulatory Collapse	Foreign	Bupropion			
Hospitalization -		Malaise		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER							
Initial or Prolonged		Renal Failure					
DAY/ORAL							
				Loperamide			
				Hydrochloride	C		
				Allopurinol	C		
				Meloxicam	C		
				Salbutamol Sulphate	C		
				Temazepam	C		

Date:07/02/01ISR Number: 3751646-5Report Type:Expedited (15-DaCompany Report #B0112293A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Balance Disorder	Foreign	Bupropion			
Initial or Prolonged		Hypoaesthesia		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER							
Required		Malaise					
DAY/ORAL							
Intervention to		Tearfulness					
Prevent Permanent		Transient Ischaemic					
Impairment/Damage		Attack					

Date:07/02/01ISR Number: 3751650-7Report Type:Expedited (15-DaCompany Report #B0112285A  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/PER		Abdominal Pain Cerebellar Infarction	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

DAY

Date:07/02/01ISR Number: 3751654-4Report Type:Expedited (15-DaCompany Report #B0111950A  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/PER		Panic Attack	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Disability  
DAY/ORAL

Date:07/02/01ISR Number: 3751703-3Report Type:Expedited (15-DaCompany Report #B0109706A  
Age:31 YR Gender:Female I/FU:F

Outcome  
Life-Threatening  
Hospitalization -  
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Initial or Prolonged  
Disability  
Required

Intervention to Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Prevent Permanent Impairment/Damage 150 MG / ORAL 6 WK	Haemorrhagic Stroke Ruptured Cerebral Aneurysm Status Epilepticus	Foreign Health Professional Company Representative	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:07/02/01ISR Number: 3751741-0Report Type:Expedited (15-DaCompany Report #A0137061A  
Age:17 YR Gender:Female I/FU:F

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL	Convulsion	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged Other ORAL	Sinus Tachycardia Suicide Attempt Tremor		Citalopram Hydrobromide Tablet (Citalopram Hydrobromide)	SS		ORAL

Date:07/02/01ISR Number: 3751743-4Report Type:Expedited (15-DaCompany Report #A0149863A  
Age:60 YR Gender:Female I/FU:F

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG / TWICE PER DAY / ORAL	Dry Mouth Dry Throat Emphysema Speech Disorder	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:07/02/01ISR Number: 3751771-9Report Type:Expedited (15-DaCompany Report #B0111860A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent TWICE PER DAY Impairment/Damage / ORAL	150 MG /	Drug Effect Decreased Drug Interaction Pregnancy	Foreign Other	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
350 MCG / PER DAY / ORAL				Norethisterone (Formulation Unknown) (Norethindrone)	SS		ORAL
				Cabergoline	C		

Date:07/02/01ISR Number: 3751772-0Report Type:Expedited (15-DaCompany Report #B0112183A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability TWICE PER DAY / ORAL	150 MG /	Cerebrovascular Accident	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Atorvastatin Calcium	C		
				Bendrofluazide	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/02/01ISR Number: 3751804-XReport Type:Expedited (15-DaCompany Report #B0112411A  
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to prevent Permanent Impairment/Damage		Chest Pain	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
				Amlodipine	C		

Date:07/02/01ISR Number: 3751805-1Report Type:Expedited (15-DaCompany Report #B0111869A  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Deep Vein Thrombosis	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG (VARIABLE DOSE)							

Date:07/03/01ISR Number: 3750964-4Report Type:Expedited (15-DaCompany Report #A0151572A  
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Phlebitis	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
Other 800MG As required		Pulmonary Embolism	Professional	Ibuprofen	C		ORAL

Date:07/03/01ISR Number: 3750965-6Report Type:Expedited (15-DaCompany Report #A0152843A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Missed		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Complications Of Maternal					
per day		Exposure To Therapeutic Drugs					

Date:07/03/01ISR Number: 3750975-9Report Type:Expedited (15-DaCompany Report #B0112964A  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Anaphylactic Reaction Ventricular Tachycardia		Zyban	PS	Glaxo Wellcome	

Date:07/03/01ISR Number: 3750976-0Report Type:Expedited (15-DaCompany Report #B0112965A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alanine Aminotransferase Increased Blood Bilirubin Increased Liver Function Test Abnormal Malaise Red Blood Cell Sedimentation Rate Increased		Zyban	PS	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/05/01ISR Number: 3751488-0Report Type:Expedited (15-DaCompany Report #A0153272A  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Atrioventricular Block		Wellbutrin	PS	Glaxo Wellcome	ORAL
75MG Twice		Complete					
per day							
		Bundle Branch Block Right		Relafen	SS	Glaxo Wellcome	ORAL
500MG Twice		Drug Interaction					
per day	14	DAY					
		Heart Rate Irregular					
		Medication Error					
		Ventricular Extrasystoles					

Date:07/05/01ISR Number: 3751514-9Report Type:Expedited (15-DaCompany Report #B0113104A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Abdominal Pain Upper		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Condition Aggravated					
Initial or Prolonged							
per day	42	DAY					
		Headache		Tylox	C		ORAL
		Palpitations		Hydroxyzine	C		ORAL
100MG per day							

Date:07/05/01ISR Number: 3751515-0Report Type:Expedited (15-DaCompany Report #B0113105A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Arthritis		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day	7	DAY					
		Insomnia		Diclofenac	C		ORAL
75MG Per day							
				Kliofem	C		ORAL
				Omeprazole	C		ORAL
20MG Per day							



Date:07/05/01ISR Number: 3752859-9Report Type:Expedited (15-DaCompany Report #A0151572A  
Age:48 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150MG (TWICE	Cardiovascular Disorder	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged PER DAY) ORAL	Phlebitis	Professional				
Other	Pulmonary Embolism	Company Representative	Ibuprofen	C		

Date:07/05/01ISR Number: 3753414-7Report Type:Periodic Company Report #254104  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Blood Alkaline Phosphatase Increased	Consumer	Clonopin	PS	Hoffmann La Roche Inc	ORAL
1 MG 3 PER DAY ORAL	Gamma-Glutamyltransferase Increased		Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
1MG 3PER DAY ORAL						

Date:07/05/01ISR Number: 3753490-1Report Type:Expedited (15-DaCompany Report #EMADSS2001003946  
Age:33 YR Gender:Female I/FU:I

Outcome	PT
Other	Drug Interaction Pregnancy On Oral

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Contraceptive

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
350 MCG, DAILY, ORAL		Foreign Health Professional	Micronor	PS	Rw Johnson Pharmaceutical Research Institute Div Ortho Pharm	ORAL
150 MG, 2 IN 1 DAILY, ORAL			Zyban (Amfebutamone Hydrochloride)	SS		ORAL
			Dostinex	C		

Date:07/05/01ISR Number: 3753534-7Report Type:Expedited (15-DaCompany Report #A0152843A  
Age:37 YR Gender:Female I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Abortion Missed Blighted Ovum Complications Of Maternal Exposure To Therapeutic Drugs	Foreign Study Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:07/05/01ISR Number: 3753629-8Report Type:Expedited (15-DaCompany Report #B0112964A  
Age:43 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG		Anaphylactic Reaction Ventricular Tachycardia	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:07/05/01ISR Number: 3753630-4Report Type:Expedited (15-DaCompany Report #B0112965A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alanine Aminotransferase Increased	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
150 MG		Blood Bilirubin Increased Malaise Red Blood Cell Sedimentation Rate Increased	Professional				

Date:07/06/01ISR Number: 3753122-2Report Type:Expedited (15-DaCompany Report #A0149363A  
Age:14 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Attention
Initial or Prolonged	Deficit/Hyperactivity
Other	Disorder Blood Pressure Increased Condition Aggravated Drug Ineffective Electroencephalogram Abnormal Grand Mal Convulsion Hallucination

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Intentional Misuse Irritability Loss Of Consciousness				Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
Memory Impairment	150MG Twice per day			Professional				
Sedation					Wellbutrin	SS	Glaxo Wellcome	
Tachycardia					Tylenol	SS		
					Imodium	SS		
	10MG Per day				Paxil	C	Glaxo Wellcome	ORAL
					Tetracycline	C		
					Retin-A	C		
TOPICAL								

Date:07/06/01ISR Number: 3753129-5Report Type:Expedited (15-DaCompany Report #B0107040A  
Age:48 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	5MG At night		Choking	Health	Zyban	PS	Glaxo Wellcome	ORAL
			Coronary Artery Disease	Professional	Nitrazepam	C		ORAL
	200MG As required		Cyanosis		Meptazinol	C		ORAL
			Loss Of Consciousness					
			Myocardial Infarction		Salbutamol	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)		200MCG As						
required								

Date:07/06/01ISR Number: 3753935-7Report Type:Expedited (15-DaCompany Report #2001011567-1  
Age:50 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Atrioventricular Block	Health	Relafen	PS	Smithkline Beecham	

500 Complete Professional Pharmaceuticals ORAL  
 Ventricular Arrhythmia  
 MILLIGRAMS  
 1.0 AS NEEDED  
 ORAL 14 DAY  
 Wellbutrin  
 (Amfebutamone  
 Hydrochloride) SS ORAL  
 75 MILLIGRAMS  
 2.0 DAILY  
 ORAL

Date:07/06/01ISR Number: 3754397-6Report Type:Expedited (15-DaCompany Report #D0017705A  
 Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Depression Suicidal	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY / ORAL				Thyroxine Sodium Nisoldipine	C C		

Date:07/06/01ISR Number: 3754398-8Report Type:Expedited (15-DaCompany Report #B0112544A  
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Migraine Subarachnoid Haemorrhage	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/06/01ISR Number: 3754400-3Report Type:Expedited (15-DaCompany Report #B0097894A

Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG/ PER DAY / ORAL	Asthenopia Discomfort  Dizziness  Gastrointestinal Candidiasis Insomnia Oral Candidiasis Throat Tightness	Foreign Study  Health  Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:07/06/01ISR Number: 3754401-5Report Type:Expedited (15-DaCompany Report #B0100339A

Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged AS DIRECTED		Anxiety Asthenia  Fall Headache Hyperventilation Lethargy Loss Of Consciousness Malaise Movement Disorder Paraesthesia Syncope Tremor Vision Blurred	Foreign Health  Professional	Bupropion Hydrochloride  Atorvastatin Calcium Enalapril	PS  C C	Glaxo Wellcome Inc	

Date:07/09/01ISR Number: 3753829-7Report Type:Expedited (15-DaCompany Report #A0153280A

Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Alcoholism		Wellbutrin	PS	Glaxo Wellcome	ORAL

Initial or Prolonged  
per day 1 YR

Amnesia

Drug Abuser  
Grand Mal Convulsion  
Medication Error  
Toxicologic Test Abnormal

Ethanol SS  
No Concurrent  
Medication C

Date:07/09/01ISR Number: 3753830-3Report Type:Expedited (15-DaCompany Report #B0099253A

Age:33 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 11 DAY Hospitalization - Initial or Prolonged Disability	C-Reactive Protein Increased Cerebrovascular Accident Precerebral Artery Occlusion Urticaria		Zyban Oral Contraceptive	PS C	Glaxo Wellcome	ORAL ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/09/01ISR Number: 3753834-0Report Type:Expedited (15-DaCompany Report #B0113284A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coronary Artery Disease Overdose Sudden Death		Zyban	PS	Glaxo Wellcome	

Date:07/09/01ISR Number: 3753835-2Report Type:Expedited (15-DaCompany Report #B0113566A  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day 35 DAY	Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day				Aspirin	C		ORAL
150MG Per day				Bendrofluazide	C	Glaxo Wellcome	ORAL
2.5MG Per day				Amlodipine	C		ORAL
10MG Per day				Simvastatin	C		ORAL
10MG At night				Enalapril	C		ORAL
10MG At night							

Date:07/09/01ISR Number: 3753836-4Report Type:Expedited (15-DaCompany Report #B0113572A  
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Twice per day 28 DAY	Atrial Fibrillation		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice per day				Simvastatin	C		ORAL
10MG Per day				Amlodipine	C		ORAL
5MG per day				Indomethacin	C		ORAL
50MG per day							



Date:07/09/01ISR Number: 3753837-6Report Type:Expedited (15-DaCompany Report #D0018145A  
 Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Haemorrhagic Stroke Hypertension		Zyban Diclofenac	PS C	Glaxo Wellcome	ORAL
UNKNOWN							

Date:07/09/01ISR Number: 3754727-5Report Type:Expedited (15-DaCompany Report #A0149363A  
 Age:14 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER Required DAY/ ORAL		Blood Pressure Increased Disturbance In Attention Drug Ineffective	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Intervention to Prevent Permanent Impairment/Damage		Electroencephalogram Abnormal Grand Mal Convulsion Hallucination Intentional Misuse Irritability Memory Impairment Sedation Tachycardia		Wellbutrin Tablet (Bupropion Hydrochloride) Paracetamol (Formulation Unknown) (Acetaminophen) Loperamide Hydrochloride Tablet Paroxetine Hydrochloride Tetracycline Tretinoin	SS    SS  SS C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/09/01ISR Number: 3754815-3Report Type:Expedited (15-DaCompany Report #A0153272A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Atrioventricular Block	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
75 MG/TWICE		Complete	Professional				
PER DAY/ORAL							
		Bundle Branch Block Bilateral		Nabumetone ( Nabumetone)	SS		ORAL
500 MG/TWICE							
PER DAY/ORAL		Drug Interaction					
		Ventricular Extrasystoles					

Date:07/09/01ISR Number: 3754868-2Report Type:Direct Company Report #  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Disability		Abnormal Behaviour Alcoholism		Zoloft 100 Mgs Roerig, J.B.	PS	Roerig, J.B.	ORAL
2 PER DAY		Delinquency					
ORAL		Drug Abuser Imprisonment		Wellbutrin 150 Mgs Glaxo Well	SS	Glaxo Well	ORAL
1 PER DAY		Obsessive-Compulsive Disorder Paraphilia Suicide Attempt Theft Thinking Abnormal					
ORAL							

Date:07/09/01ISR Number: 3754914-6Report Type:Expedited (15-DaCompany Report #2001058760US  
Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Headache	Health	Solu-Medrol	PS	Pharmacia And Upjohn	

Initial or Prolonged	Pruritus	Professional		Co	
INTRAVENOUS	125 MG, QID,				
Other	Urticaria				
IV					
			Hydroxyzine	SS	ORAL
ORAL					
			Prednisone	SS	ORAL
ORAL					
			Wellbutrin (Amfebutamone Hydrochloride)	SS	ORAL
150 MG, BID					
ORAL					
			Ortho Tri-Cyclen	C	
			Propranolol	C	

Date:07/09/01ISR Number: 3755450-3Report Type:Expedited (15-DaCompany Report #B0107040A  
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Choking	Foreign	Bupropion			
		Coronary Artery Disease	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL							
		Cyanosis	Professional	Nitrazepam	C		
		Loss Of Consciousness		Meptazinol	C		
		Myocardial Infarction		Salbutamol Sulphate	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/09/01ISR Number: 3755531-4Report Type:Expedited (15-DaCompany Report #B0113104A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ TWICE Initial or Prolonged PER DAY/ ORAL		Abdominal Pain Upper	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Condition Aggravated					
		Headache		Tylox	C		
		Palpitations		Hydroxyzine	C		

Date:07/09/01ISR Number: 3755533-8Report Type:Expedited (15-DaCompany Report #B0113105A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 300 MG/ PER DAY/ ORAL		Arthritis	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Insomnia					
				Diclofenac	C		
				Kliefem	C		
				Omeprazole	C		

Date:07/10/01ISR Number: 3754634-8Report Type:Expedited (15-DaCompany Report #A0152291A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day 265 DAY 1TAB Per day		Blood Cholesterol Increased		Bupropion	PS	Glaxo Wellcome	ORAL
		Cholelithiasis		Orthotricyclen	C		ORAL
		Weight Decreased					

Date:07/10/01ISR Number: 3754642-7Report Type:Expedited (15-DaCompany Report #B0111329A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction		Zyntabac	PS	Glaxo Wellcome	ORAL

Date:07/10/01ISR Number: 3754645-2Report Type:Expedited (15-DaCompany Report #B0112964A  
Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	19 DAY	Anaphylactic Reaction Ventricular Tachycardia		Zyban	PS	Glaxo Wellcome	ORAL

Date:07/10/01ISR Number: 3754649-XReport Type:Expedited (15-DaCompany Report #B0113567A  
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Twice per day	5 DAY	Anxiety Dizziness Panic Attack		Zyban	PS	Glaxo Wellcome	ORAL

Date:07/10/01ISR Number: 3754651-8Report Type:Expedited (15-DaCompany Report #D0018424A  
Age:60 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged Other	Atrophy Convulsion Delirium

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Malaise					
		Nervous System Disorder					
			Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	1 DAY			Hormone-Plaster	SS		
UNKNOWN							

Date:07/10/01ISR Number: 3756647-9Report Type:Expedited (15-DaCompany Report #A103444  
 Age:79 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Blood Pressure Increased	Consumer	Zoloft Tablets	PS	Pfizer	
Intervention to		Drug Ineffective				Pharmaceuticals Inc	
50.00 MG							
Prevent Permanent		Ecchymosis					
TOTAL:DAILY							
Impairment/Damage		Fall		Wellbutrin	SS		ORAL
300.00 MG							
TOTAL:PID:ORA		Haemorrhagic Stroke					
L		Head Injury					
		Headache		Coumadin	SS		
300.00 MG		Heart Rate Decreased		Effexor Sr	SS		ORAL
TOTAL:PID:ORA		Heart Rate Increased					
L		Hyperhidrosis					
		Medication Error		Toprol	C		
		Overdose		Zebeta	C		
		Syncope		Lanoxin	C		
				Lasix	C		
				Synthroid	C		
				Vitamin E	C		
				Estrogen	C		

Date:07/11/01ISR Number: 3755614-9Report Type:Expedited (15-DaCompany Report #263690  
 Age:41 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Abnormal Dreams		Xenical	PS	Roche	
WITH MEALS		181 DAY	Bone Disorder		Didronel	C		
			Bone Pain		Estrace	C		
			Condition Aggravated		Zestril	C		
			Drug Ineffective		Naprosyn Ec	I	Roche	
			Drug Interaction		Keflex	I		
			Food Craving		Ceftin	I		
			Haemophilus Infection		Augmentin	I		
			Hypoaesthesia		Wellbutrin	I		
			Neuralgia					
			Osteomyelitis					
			Osteonecrosis					
			Pain In Jaw					
			Poor Peripheral Circulation					

Date:07/11/01ISR Number: 3755628-9Report Type:Expedited (15-DaCompany Report #A0153345A  
Age:31 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Complications Of Maternal Exposure To Therapeutic Drugs Premature Baby Premature Labour Premature Rupture Of

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Membranes

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
250MG per day	139 DAY		Bupropion	PS	Glaxo Wellcome	ORAL

Date:07/11/01ISR Number: 3755633-2Report Type:Expedited (15-DaCompany Report #B0110444A  
Age:67 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	7 DAY	Arrhythmia		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - RESPIRATORY		Coma		Fluticasone	C	Glaxo Wellcome	
Initial or Prolonged (INHALATION)		Malaise					
RESPIRATORY		Nausea		Salmeterol	C	Glaxo Wellcome	
(INHALATION)		Pulmonary Embolism					
RESPIRATORY		Syncope		Beclomethasone	C	Glaxo Wellcome	
(INHALATION)	6PUFF per day			Salbutamol	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)	100UG Six						
times per day				Spirolactone	C		ORAL
75MG Twice				Losartan	C		ORAL
per day				Frusemide	C		ORAL

Date:07/11/01ISR Number: 3755643-5Report Type:Expedited (15-DaCompany Report #B0113169A  
Age:65 YR Gender:Male I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Decreased Appetite		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Hospitalization -		Dry Throat					
per day							
Initial or Prolonged		Dyspnoea Oedema Peripheral Retching					

Date:07/11/01ISR Number: 3755644-7Report Type:Expedited (15-DaCompany Report #B0113282A  
Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Malaise		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Per day	3 DAY	Myocardial Infarction					

Date:07/11/01ISR Number: 3756507-3Report Type:Expedited (15-DaCompany Report #TEST00201002994  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Arthralgia Dermatitis	Health Professional	Androgel	PS	Unimed Pharmaceuticals Inc	
5 G QD TD		Serum Sickness Urticaria	Company Representative	Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL
DAILY PO				Growth Hormone (Human Growth Hormone, Recombinant) Steroid	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/12/01ISR Number: 3756349-9Report Type:Expedited (15-DaCompany Report #B0100762A  
Age:19 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 22 DAY	Amnesia		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - 9.6G Single	Coma		Brufen	SS		ORAL
Initial or Prolonged dose	Confusional State					
Other	Coordination Abnormal Disorientation Grand Mal Convulsion Headache Loss Of Consciousness Overdose Suicide Attempt		Alcohol Alcohol Cannabis	SS C C		ORAL

Date:07/12/01ISR Number: 3756351-7Report Type:Expedited (15-DaCompany Report #B0104336A  
Age:34 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death 150MG Per day	Completed Suicide		Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY  (INHALATION)			Salbutamol	C	Glaxo Wellcome	

Date:07/12/01ISR Number: 3756353-0Report Type:Expedited (15-DaCompany Report #B0109122A  
Age:41 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150MG Per day 3 DAY	Alanine Aminotransferase Increased Alpha 1 Foetoprotein Increased Aspartate Aminotransferase	Health Professional	Zyntabac	PS	Glaxo Wellcome	ORAL

Increased  
 Carcinoembryonic Antigen  
 Increased  
 Gamma-Glutamyltransferase  
 Increased

Date:07/12/01ISR Number: 3756355-4Report Type:Expedited (15-DaCompany Report #B0111855A  
 Age:56 YR Gender:Female I/FU:F

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice	Blood Lactate		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day 7 WK	Dehydrogenase Increased					
Other 20MG Per day	Cardiac Disorder		Citalopram	C		
300MG Per day	Cranial Nerve Disorder		Thiamine	C		
.25MG Twice	Diplopia		Alprazolam	C		
per day	Hypercholesterolaemia					
2U Per day	Myocardial Infarction		Fluticasone	C	Glaxo Wellcome	
2U Per day	Nystagmus		Salmeterol	C	Glaxo Wellcome	
	Pulmonary Function Test Decreased Supraventricular Extrasystoles Tachycardia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/12/01ISR Number: 3756363-3Report Type:Expedited (15-DaCompany Report #B0113725A  
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day	5 DAY	Hepatitis	Zyban	PS	Glaxo Wellcome	ORAL
Pericardial Effusion							

Date:07/12/01ISR Number: 3756364-5Report Type:Expedited (15-DaCompany Report #B0114026A  
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG As	Initial or Prolonged	Fracture	Zyban	PS	Glaxo Wellcome	
directed							
Insomnia							
Palpitations							
Tremor							

Date:07/12/01ISR Number: 3757002-8Report Type:Expedited (15-DaCompany Report #A0152291A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG (TWICE	Initial or Prolonged	Blood Cholesterol	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
PER DAY) ORAL							
Increased							
Health							
Cholelithiasis							
Weight Decreased							
Professional							
Ortho Tri-Cyclen							
C							

Date:07/12/01ISR Number: 3757004-1Report Type:Expedited (15-DaCompany Report #A0153280A  
Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150 MG (TWICE		Amnesia	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

Initial or Prolonged Grand Mal Convulsion  
PER DAY) ORAL  
Intentional Misuse  
Suicide Attempt  
Toxicologic Test Abnormal

Professional  
Company  
Representative

Ethanol (Alcohol) SS

Date:07/12/01ISR Number: 3757029-6Report Type:Direct  
Age:48 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG BID	1 MON	Hearing Impaired Hypoaesthesia		Zyban (150mg)	PS		

Date:07/12/01ISR Number: 3757179-4Report Type:Expedited (15-DaCompany Report #B0113572A  
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150 MG TWICE PER DAY ORAL		Atrial Fibrillation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
				Simvastatin	C		
				Amlodipine	C		
				Indomethacin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/12/01ISR Number: 3757180-0Report Type:Expedited (15-DaCompany Report #B0113566A  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG PER	Angina Pectoris	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
DAY ORAL				Aspirin	C		
				Bendrofluazide	C		
				Amlodipine	C		
				Simvastatin	C		
				Enalapril	C		

Date:07/12/01ISR Number: 3757181-2Report Type:Expedited (15-DaCompany Report #D0018145A  
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG ORAL	Haemorrhagic Stroke	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Hypertension		Diclofenac	C		

Date:07/12/01ISR Number: 3757182-4Report Type:Expedited (15-DaCompany Report #B0099253A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG ORAL	C-Reactive Protein	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - Initial or Prolonged Disability		Increased Carotid Artery Stenosis Cerebrovascular Accident Precerebral Artery Occlusion Urticaria	Health Professional	Oral Contraceptive	C		

Date:07/12/01ISR Number: 3757189-7Report Type:Expedited (15-DaCompany Report #B0113284A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coronary Artery Disease Overdose Sudden Death	Foreign Consumer	Zyban	PS	Glaxo Wellcome Inc	

Date:07/12/01ISR Number: 3757190-3Report Type:Expedited (15-DaCompany Report #B0113567A  
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150 MG / TWICE PER DAY / ORAL		Anxiety  Dizziness  Panic Attack	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:07/12/01ISR Number: 3757192-7Report Type:Expedited (15-DaCompany Report #D0018424A  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / PER Initial or Prolonged DAY / ORAL 1 DAY Other		Cerebral Atrophy  Convulsion  Delirium Malaise	Foreign  Health  Professional	Zyban   Hormones (Formulation Unknown)	PS   SS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/12/01ISR Number: 3757224-6Report Type:Expedited (15-DaCompany Report #B0111329A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG / ORAL	Myocardial Infarction	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:07/12/01ISR Number: 3757225-8Report Type:Expedited (15-DaCompany Report #B0112964A  
Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG / TWICE PER DAY / ORAL	Anaphylactic Reaction Ventricular Tachycardia	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:07/12/01ISR Number: 3757414-2Report Type:Expedited (15-DaCompany Report #A0152528A  
Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL	3 DAY	Angina Pectoris	Foreign	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Hypokalaemia	Literature Health Professional	Pseudoephedrine (Formulation Unknown)	SS		
Other		Myocardial Infarction		(Pseudoephedrine)	SS		
		Myocardial Ischaemia		Erythromycin (Formulation Unknown)	SS		
		Respiratory Alkalosis		(Erythromycin)	SS		
		Tachypnoea		Nicotine (Formulation Unknown) (Nicotine)	SS		



Date:07/13/01ISR Number: 3757131-9Report Type:Direct  
Age:57 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis		Zyban 150mg Catalytica/Glaxo Wellcome	PS	Catalytica/Glaxo Wellcome	ORAL
150MG BID							
ORAL				Glucophage Isordil Nicoderm Micronase	C C C C		

Date:07/13/01ISR Number: 3757330-6Report Type:Expedited (15-DaCompany Report #A0138561A  
Age:53 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Alcoholism
Initial or Prolonged	Cognitive Disorder
Disability	Dementia
Other	Disturbance In Attention Injury Memory Impairment

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Road Traffic Accident Sleep Apnoea Syndrome Social Avoidant Behaviour	Wellbutrin	PS	Glaxo Wellcome	ORAL
20MG Per day			Alcohol Celexa	SS C		ORAL

Date:07/13/01ISR Number: 3757331-8Report Type:Expedited (15-DaCompany Report #A0149272A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 71 DAY		Haemoptysis		Zyban	PS	Glaxo Wellcome	ORAL
TOPICAL		Lung Disorder Neoplasm		Prevacid Estrogel	C C		

Date:07/13/01ISR Number: 3758668-9Report Type:Expedited (15-DaCompany Report #B0113282A  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG/ PER DAY/ ORAL		Malaise	Foreign	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
		Myocardial Infarction	Literature				

Date:07/13/01ISR Number: 3758671-9Report Type:Expedited (15-DaCompany Report #B0113169A  
Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 150 MG/ TWICE		Decreased Appetite Dry Throat	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Initial or Prolonged      Dyspnoea      Professional  
PER DAY/ ORAL

Oedema Peripheral  
Retching

Date:07/13/01ISR Number: 3758753-1Report Type:Expedited (15-DaCompany Report #263690  
Age:41 YR      Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abnormal Dreams	Consumer	Xenical	PS	Hlr Technology	ORAL
120 MG 3 PER		Angiopathy					
DAY ORAL		Bone Pain		Naprosyn Ec			
		Condition Aggravated		(Naproxen)	SS		ORAL
1 PER PRN		Drug Ineffective					
ORAL		Drug Interaction		Keflex (Cephalexin)	SS		ORAL
4 GRAM DAILY		Food Craving					
ORAL		Haemophilus Infection		Ceftin (Cefuroxime			
		Hypoaesthesia		Axetil)	SS		ORAL
ORAL		Osteomyelitis		Augmentin			
		Osteonecrosis		(*Amoxicillin/*Amoxi			
		Pain In Jaw		cillin			
375MG 3 PER				Sodium/*Clavulanate			
DAY ORAL				Potassium)	SS		ORAL
				Wellbutrin			
				(Bupropion			
				Hydrochloride)	SS		ORAL
ORAL				Zestril (Lisinopril)	C		
				Didronel (Etidronate			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Disodium) C  
 Estrace (Estradiol) C

Date:07/13/01ISR Number: 3758804-4Report Type:Expedited (15-DaCompany Report #HQ3100411JUL2001  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Complications Of Maternal Exposure To Therapeutic	Health Professional	Effexor	PS	Wyeth Ayerst Laboratories	ORAL
ORAL	Drugs Convulsion Neonatal		Remeron (Mirtazapine)	SS		ORAL
ORAL			Wellbutrin	SS		

Date:07/13/01ISR Number: 3758905-0Report Type:Expedited (15-DaCompany Report #B0110444A  
 Age:67 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death Hospitalization - 150 MG / ORAL Initial or Prolonged	Arrhythmia Coma	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
	Malaise Nausea Pulmonary Embolism Syncope		Fluticasone Propionate Salmeterol Xinafoate Beclomethasone Dipropion Salbutamol Sulphate Spironolactone Losartan Potassium Frusemide	C C C C C C C C		

Date:07/13/01ISR Number: 3758940-2Report Type:Expedited (15-DaCompany Report #A0153345A  
 Age:31 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - TWICE PER	Complications Of Maternal	Foreign	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL

Initial or Prolonged  
DAY/ ORAL

Exposure To Therapeutic  
Drugs  
Pregnancy  
Premature Baby  
Premature Rupture Of  
Membranes

Study  
Health  
Professional

Date:07/16/01ISR Number: 3758460-5Report Type:Expedited (15-DaCompany Report #B0111853A  
Age:52 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice Initial or Prolonged per day 16 WK Disability	Carotid Artery Thrombosis Cerebrovascular Accident		Zyban	PS	Glaxo Wellcome	
	Supraventricular Extrasystoles Ventricular Extrasystoles Ventricular Hypertrophy		Alprazolam Sertraline Premelle Alcohol	C C C C		ORAL  ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/16/01ISR Number: 3758465-4Report Type:Expedited (15-DaCompany Report #B0114027A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Agitation		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Other		Benign Breast Neoplasm					
per day		Emotional Disorder		Mebeverine	C		ORAL

Date:07/16/01ISR Number: 3758466-6Report Type:Expedited (15-DaCompany Report #B0114185A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour		Zyban	PS	Glaxo Wellcome	
		Dementia					
		Mental Disorder					

Date:07/16/01ISR Number: 3759442-XReport Type:Expedited (15-DaCompany Report #A103444  
Age:79 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Blood Pressure Increased	Consumer	Zoloft	PS	Pfizer	
Intervention to		Blood Thyroid Stimulating				Pharmaceuticals Inc	
50.00 MG							
Prevent Permanent		Hormone Decreased					
TOTAL:DAILY							
Impairment/Damage		Drug Ineffective		Wellbutrin	SS		ORAL
300.000 MG							
		Ecchymosis					
TOTAL:BID:ORA							
		Epistaxis					
L							
		Fall		Coumadin	SS		
		Haemorrhagic Stroke		Effexor Sr	SS		ORAL
300.00							
		Head Injury					
TOTAL:BID:ORA							
		Headache					
L							

Heart Rate Decreased  
 Heart Rate Increased  
 Hyperhidrosis  
 Medication Error  
 Mood Swings  
 Movement Disorder  
 Overdose  
 Parkinson'S Disease  
 Staring  
 Syncope  
 Thyroid Function Test  
 Abnormal

Toprol C  
 Zebeta C  
 Lanoxin C  
 Lasix C  
 Synthroid C  
 Vitamin E C  
 Estrogen C

Date:07/16/01ISR Number: 3759478-9Report Type:Expedited (15-DaCompany Report #A0149272A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Haemoptysis Lung Disorder	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
ORAL		Neoplasm	Professional	Lansoprazole Oestradiol	C C		

Date:07/16/01ISR Number: 3759788-5Report Type:Direct Company Report #  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash Pruritic		Zyban 150mg Catalytica/Glaxowell			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

150 MG / BID/  
ORAL

Product	Role	Manufacturer	Route
come	PS	Catalytica/Glaxowell come	ORAL

Nicoderm

Product	Role	Manufacturer	Route
Nicoderm	C		

Date:07/16/01ISR Number: 3759829-5Report Type:Expedited (15-DaCompany Report #A0138561A  
Age:53 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG TWICE Initial or Prolonged PER DAY ORAL	Alcoholism	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Disability Required	Dementia	Professional				
Intervention to Prevent Permanent Impairment/Damage	Memory Impairment Personality Change Road Traffic Accident Sleep Apnoea Syndrome Social Avoidant Behaviour Spinal Cord Injury	Company Representative	Ethanol (Formulation Unknown) (Alcohol) Citalopram Hydrobromide	SS C		

Date:07/16/01ISR Number: 3760012-8Report Type:Expedited (15-DaCompany Report #B0113725A  
Age:67 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150 MG / PER DAY / ORAL	Hepatitis	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
	Pericardial Effusion					

Date:07/16/01ISR Number: 3760028-1Report Type:Expedited (15-DaCompany Report #B0100762A  
Age:19 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening ORAL	Amnesia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Hospitalization - 9.6 G/SINGLE	Confusional State	Literature	Ibuprofen Otc Tablet	SS		ORAL



Initial or Prolonged DOSE/ORAL Required Intervention to ORAL Prevent Permanent Impairment/Damage	Coordination Abnormal  Disorientation Grand Mal Convulsion  Headache Intentional Misuse Loss Of Consciousness Suicide Attempt	Health  Professional	Ethanol (Formulation Unknown ) (Alcohol)  Cannabis Ethanol	SS  C C	ORAL
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Date:07/16/01ISR Number: 3760032-3Report Type:Expedited (15-DaCompany Report #B0111855A  
Age:56 YR Gender:Female I/FU:F

Outcome Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage	PT Angiopathy Arterial Stenosis Blood Lactate Dehydrogenase Increased Cardiomyopathy Diplopia Echography Abnormal Electrocardiogram Ambulatory Abnormal Electrocardiogram Q Waves Electrocardiogram St Segment Elevation Eye Disorder Hypercholesterolaemia Hypokinesia
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL		Myocardial Infarction Myocardial Ischaemia Nystagmus Pulmonary Function Test Abnormal Supraventricular Extrasystoles Tachycardia Vith Nerve Paralysis	Foreign Health Professional	Zyban  Citalopram Thiamine Alprazolam Fluticasone Propionate Salmeterol Xinafoate	PS  C C C C C	Glaxo Wellcome Inc	ORAL

Date:07/16/01ISR Number: 3760034-7Report Type:Expedited (15-DaCompany Report #B0109122A  
Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG/PER DAY/ORAL		Alanine Aminotransferase Increased Alpha 1 Foetoprotein Increased Aspartate Aminotransferase Increased Gamma-Glutamyltransferase Increased Hepatitis A Liver Function Test Abnormal	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:07/16/01ISR Number: 3760077-3Report Type:Expedited (15-DaCompany Report #B0104336A  
Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG / PER		Completed Suicide	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL

DAY / ORAL

Health

Professional

Salbutamol Sulphate C

Date:07/16/01ISR Number: 3760625-3Report Type:Expedited (15-DaCompany Report #B0114026A

Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / AS		Fracture	Foreign	Zyban	PS	Glaxo Wellcome Inc	
Initial or Prolonged DIRECTED		Insomnia	Health				
		Palpitations Surgery Tremor	Professional				

Date:07/17/01ISR Number: 3759265-1Report Type:Expedited (15-DaCompany Report #A0154036A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Wellbutrin Paxil	PS SS	Glaxo Wellcome Glaxo Wellcome	ORAL ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/17/01ISR Number: 3759267-5Report Type:Expedited (15-DaCompany Report #A0154344A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Bupropion	PS	Glaxo Wellcome	ORAL
300MG Per day		Complications Of Maternal Exposure To Therapeutic Drugs					

Date:07/17/01ISR Number: 3759803-9Report Type:Direct Company Report #  
Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blepharospasm		Burproprion (75mg)	PS		ORAL
75MG Q AM PO		Muscle Spasms					

Date:07/17/01ISR Number: 3760205-XReport Type:Periodic Company Report #HQ2500719OCT2000  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Urticaria	Consumer	Advil	PS	Whitehall Laboratories Inc Div American Home Products Corp	ORAL

THREE TO SIX

TABLETS ONCE

DAILY, ORAL

150 MG 1X PER

1 DAY, ORAL

				Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL
--	--	--	--	---	----	--	------

				Unspecified Estrogen Replacement Therapy (Unspecified			
--	--	--	--	---	--	--	--

Estrogen Replacement  
Therapy ) C  
Atarax (Hydroxyzine  
Hydrochloride) C

Date:07/18/01ISR Number: 3760399-6Report Type:Expedited (15-DaCompany Report #A0154486A  
Age:81 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 3 MON			Brain Neoplasm Benign	Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged			Depression Psychotic Disorder	Procardia	C		

Date:07/19/01ISR Number: 3761213-5Report Type:Expedited (15-DaCompany Report #B0094547A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Twice per day			Completed Suicide	Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/19/01  
 ISR Number: 3761705-9  
 Report Type:Expedited (15-DaCompany Report #A0154344A  
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Study	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
300 MG / PER		Complications Of Maternal	Health				
DAY / ORAL		Exposure To Therapeutic	Professional				
		Drugs					

Date:07/19/01  
 ISR Number: 3761852-1  
 Report Type:Expedited (15-DaCompany Report #A0154036A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
ORAL			Consumer	Paroxetine			
				Hydrochloried			
				(Paroxetine			
				Hydrochloride)	SS		ORAL
ORAL							

Date:07/20/01  
 ISR Number: 3762278-7  
 Report Type:Direct  
 Age:71 YR Gender:Female I/FU:I  
 Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Abdominal Pain Upper		Relafin 200mg	PS		ORAL
2, 2 ORAL		Anxiety		Wellbutrin Sr 150 Mg			
		Cough		Catalytica	SS		ORAL
1 /2/ ORAL		Diarrhoea					
		Dry Skin					
		Flushing					
		Insomnia					
		Movement Disorder					
		Oedema Peripheral					
		Saliva Altered					

Date:07/20/01ISR Number: 3762405-1Report Type:Expedited (15-DaCompany Report #A0154486A  
Age:81 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - ORAL	Anxiety	Consumer	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged	Brain Neoplasm Benign Depression Psychotic Disorder		Nifedipine	C		

Date:07/23/01ISR Number: 3762477-4Report Type:Expedited (15-DaCompany Report #A0055937A  
Age:55 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Agitation
Initial or Prolonged	Agoraphobia
Disability	Alopecia Anaphylactic Shock Anuria Autonomic Nervous System Imbalance Back Pain Balance Disorder Bipolar Disorder Blood Cholesterol

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	12 DAY	Increased Blood Glucose Increased Blood Pressure Fluctuation	Zyban	PS	Glaxo Wellcome	ORAL
1.25MG Per day		Blood Pressure Increased Bradycardia Cognitive Disorder Convulsion Cystocele Delusion Depersonalisation Disorientation Disturbance In Attention Drug Hypersensitivity Dyspepsia Dyspnoea Dysthymic Disorder Dysuria Fatigue Haematuria Hallucination, Auditory Hallucination, Olfactory Hyperventilation Hypomania Intervertebral Disc Degeneration Labile Blood Pressure Mania Memory Impairment Mental Impairment Micturition Urgency Mitral Valve Incompetence Mood Swings Obsessive-Compulsive Disorder Oedema Palpitations Paranoia Psychomotor Hyperactivity Renal Disorder Renal Impairment Sciatica Speech Disorder Spinal Osteoarthritis	Premarin  Vitamin Asa Melatonin	C  C C		



Tachycardia  
Tinnitus  
Transient Ischaemic  
Attack  
Urethral Stricture  
Urinary Incontinence  
Ventricular Extrasystoles  
Weight Increased

Date:07/23/01ISR Number: 3762479-8Report Type:Expedited (15-DaCompany Report #A0144968A  
Age:61 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice Initial or Prolonged per day	Gastrointestinal Haemorrhage Ulcer		Zyban Aspirin Coumadin	PS SS C	Glaxo Wellcome Glaxo Wellcome	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/23/01ISR Number: 3762480-4Report Type:Expedited (15-DaCompany Report #A0150198A

Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day 2 DAY	Facial Palsy		Zyban	PS	Glaxo Wellcome	ORAL
				Clonidine	C		
				Estradiol	C		
				Aspirin	C		

Date:07/23/01ISR Number: 3762490-7Report Type:Expedited (15-DaCompany Report #B0112965A

Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Alanine Aminotransferase Increased Arthralgia Blood Bilirubin Increased Jaundice Lethargy Malaise Myalgia Red Blood Cell Sedimentation Rate Increased		Zyban	PS	Glaxo Wellcome	

Date:07/23/01ISR Number: 3762493-2Report Type:Expedited (15-DaCompany Report #B0113971A

Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	Sudden Death		Bupropion	PS	Glaxo Wellcome	ORAL
				Dihydrocodeine	C		ORAL
				Prothiaden	C		
UNKNOWN				Imipramine	C		

UNKNOWN

Date:07/23/01ISR Number: 3762498-1Report Type:Expedited (15-DaCompany Report #B0114445A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Difficulty In Walking		Zyban	PS	Glaxo Wellcome	
64 DAY		Dyspnoea					
		Fatigue					
		Feeling Abnormal					
		Headache					
		Insomnia					
		Pain In Extremity					
		Paralysis					
		Polymyalgia Rheumatica					
		Sedation					
		Tremor					

Date:07/23/01ISR Number: 3762499-3Report Type:Expedited (15-DaCompany Report #B0114486A  
Age:66 YR Gender:Female I/FU:I

Outcome	PT
Disability	Abdominal Pain Upper
Other	Blister
	Feeling Hot
	Headache

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	13 DAY	Hyperhidrosis Nausea Pain Tremor		Zyban	PS	Glaxo Wellcome	ORAL
135MG Three times per day				Thyroxine Mebeverine	C C	Glaxo Wellcome	ORAL ORAL

Date:07/23/01ISR Number: 3762500-7Report Type:Expedited (15-DaCompany Report #B0114491A  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Duodenal Ulcer Perforation		Zyban Trimipramine	PS C	Glaxo Wellcome	ORAL
UNKNOWN				Theophylline	C		
UNKNOWN				Aspirin	C		ORAL

Date:07/23/01ISR Number: 3762502-0Report Type:Expedited (15-DaCompany Report #B0114651A  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Ischaemia Sudden Death		Zyban	PS	Glaxo Wellcome	ORAL

Date:07/23/01ISR Number: 3762503-2Report Type:Expedited (15-DaCompany Report #B0114664A  
Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Twice per day	11 DAY	Anorexia Diplopia		Zyban	PS	Glaxo Wellcome	ORAL

100MG Per day 6 DAY      Hyperhidrosis      Paludrine      SS      ORAL  
 Nausea  
 Palpitations  
 Vision Blurred

Date:07/23/01ISR Number: 3763712-9Report Type:Expedited (15-DaCompany Report #EMADSS2001003946  
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction Unintended Pregnancy	Foreign Health Professional	Micronor	PS	Rw Johnson Pharmaceutical Research Institute Div Ortho Pharm	ORAL
350 MCG,							
DAILY, ORAL							
150 MG, 2 IN				Zyban (Amfebutamone Hydrochloride)	SS		ORAL
1 DAILY, ORAL				Dostinex (Cabergoline)	C		

Date:07/23/01ISR Number: 3763772-5Report Type:Expedited (15-DaCompany Report #B0094547A  
 Age: Gender:Male I/FU:F

Outcome	PT	Report Source
Death	Completed Suicide	Foreign Health

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Professional  
Company  
Representative

Dose	Duration	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY/ ORAL		Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:07/24/01ISR Number: 3763660-4Report Type:Expedited (15-DaCompany Report #A0154610A  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300MG Per day		Abortion Spontaneous		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Complications Of Maternal Exposure To Therapeutic Drugs					

Date:07/24/01ISR Number: 3763662-8Report Type:Expedited (15-DaCompany Report #A0154846A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day		Abortion Spontaneous		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Complications Of Maternal Exposure To Therapeutic Drugs					

Date:07/24/01ISR Number: 3763666-5Report Type:Expedited (15-DaCompany Report #B0110446A  
Age:79 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Atrial Flutter	Health	Zyban	PS	Glaxo Wellcome	ORAL

Initial or Prolonged      Dyspnoea      Professional  
per day      47      DAY

Date:07/24/01ISR Number: 3763667-7Report Type:Expedited (15-DaCompany Report #B0110883A  
Age:60 YR      Gender:Female      I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiomyopathy	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
150MG See text	10 DAY	Left Ventricular Failure					
50MG Unknown		Pulmonary Oedema		Amitriptyline	C		ORAL
5MG Unknown				Nitrazepam	C		ORAL
2MG Unknown				Loperamide	C		ORAL
				Co-Codamol	C		ORAL

Date:07/24/01ISR Number: 3763674-4Report Type:Expedited (15-DaCompany Report #B0114448A  
Age:44 YR      Gender:Male      I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Hepatic Congestion	Consumer	Zyntabac	PS	Glaxo Wellcome	ORAL
55 DAY		Hepatomegaly					
		Myocardial Infarction					
		Pulmonary Haemorrhage					
		Pulmonary Oedema					
		Thyroid Neoplasm					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/24/01ISR Number: 3763676-8Report Type:Expedited (15-DaCompany Report #B0114827A  
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Nystagmus		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	13	DAY					
		Vertigo					
RESPIRATORY				Fluticasone	C	Glaxo Wellcome	
(INHALATION)							
RESPIRATORY				Salbutamol	C	Glaxo Wellcome	
(INHALATION)							
				Tramadol	C		
				Lansoprazole	C		ORAL
80MG Per day				Frusemide	C		ORAL
20MG Per day				Tamoxifen	C		ORAL

Date:07/24/01ISR Number: 3763677-XReport Type:Expedited (15-DaCompany Report #B0114829A  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Hypothyroidism		Zyban	PS	Glaxo Wellcome	ORAL
1	DAY			Hepatitis A Immunization	C		
1	DAY			Salmonella Typhi Vaccine	C		
				Salbutamol	C	Glaxo Wellcome	

Date:07/24/01ISR Number: 3763678-1Report Type:Expedited (15-DaCompany Report #B0114982A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Hospitalization - 150MG Initial or Prolonged Variable dose	Abdominal Hernia	Health	Zyntabac	PS	Glaxo Wellcome	ORAL
		Professional				

Date:07/24/01ISR Number: 3763679-3Report Type:Expedited (15-DaCompany Report #B0115003A  
Age:50 YR Gender:Male I/FU:F

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG Per day  360MG Per day	Hypertensive Crisis		Zyban Irbesartan  Diltiazem  Calcium Carbonate Hydrochlorothiazide Vitamins Frusemide	PS C  C  C C C C	Glaxo Wellcome	

Date:07/24/01ISR Number: 3763680-XReport Type:Expedited (15-DaCompany Report #B0115005A  
Age:73 YR Gender:Female I/FU:F

Outcome Dose Other Duration	PT	Report Source	Product	Role	Manufacturer	Route
38 DAY	Asthenia  Blood Pressure Increased Convulsion Dysgeusia Fatigue Loss Of Consciousness Weight Decreased		Zyban	PS	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/24/01ISR Number: 3763681-1Report Type:Expedited (15-DaCompany Report #B0115008A  
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Physical Assault		Zyban	PS	Glaxo Wellcome	

Date:07/25/01ISR Number: 3764352-8Report Type:Expedited (15-DaCompany Report #A0124641A  
Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Arrhythmia	Health	Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
150MG Twice		Dehydration	Professional				
per day	6	MON		Alcohol	SS		
		Heat Stroke					
		Sudden Death					

Date:07/25/01ISR Number: 3764356-5Report Type:Expedited (15-DaCompany Report #A0152007A  
Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coronary Artery Disease	Health	Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
450MG Per day			Professional	Statins	C		

Date:07/25/01ISR Number: 3764365-6Report Type:Expedited (15-DaCompany Report #B0094547A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Blood Alcohol Increased	Health	Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Completed Suicide	Professional				
per day				Alcohol	SS		

Date:07/25/01ISR Number: 3764366-8Report Type:Expedited (15-DaCompany Report #B0094689A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Agitation		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Other		Insomnia					
per day	12 DAY						
		Restlessness					
		Weight Decreased					

Date:07/25/01ISR Number: 3764367-XReport Type:Expedited (15-DaCompany Report #B0103442A  
Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Atherosclerosis	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
		Dermatitis					
per day							
		Myocardial Infarction		Prednisolone	C		ORAL
40MG Per day	6 DAY						
				Nitrazepam	C		ORAL
5MG Per day							

Date:07/25/01ISR Number: 3764375-9Report Type:Expedited (15-DaCompany Report #B0114925A  
Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Deformity Of Orbit
Initial or Prolonged	Diplopia
Other	Ecchymosis
	Electroencephalogram

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Abnormal Facial Bones Fracture Sedation Wrist Fracture	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Zyban	PS	Glaxo Wellcome	ORAL

Date:07/25/01ISR Number: 3764389-9Report Type:Direct  
Age:17 YR Gender:Female I/FU:I Company Report #

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration			Zyban	PS		ORAL
Dose		Aggression					
Life-Threatening							
BID ORAL		Hyperthyroidism					
Hospitalization -		Impulsive Behaviour					
Initial or Prolonged		Psychotic Disorder					

Date:07/26/01ISR Number: 3765377-9Report Type:Expedited (15-DaCompany Report #A0125519A  
Age:49 YR Gender:Female I/FU:F

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose		Anxiety	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
Death		Arrhythmia	Professional	Acular	C		
Disability				Vancenase Aq	C	Glaxo Wellcome	NASAL
INTRAOcular		Coronary Artery Disease					
		Fibrosis					
		Hepatic Steatosis					
		Ill-Defined Disorder					
		Injury					
		Pain					
		Pancreatic Disorder					
		Pulmonary Congestion					
		Pulmonary Oedema					

Date:07/26/01ISR Number: 3765379-2Report Type:Expedited (15-DaCompany Report #A0149272A  
Age:54 YR Gender:Female I/FU:F

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose							

Other 150MG Twice per day	81 DAY	Haemoptysis Lung Disorder Neoplasm	Health Professional	Zyban	PS	Glaxo Wellcome	ORAL
30MG Per day				Prevacid	C		
TOPICAL				Estrogel	C		

Date:07/26/01ISR Number: 3765392-5Report Type:Expedited (15-DaCompany Report #B0099502A  
Age:53 YR Gender:Female I/FU:F

Outcome Dose Death 150MG Twice per day	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Haemorrhagic Stroke	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
		Ruptured Cerebral Aneurysm		Estracombi	C		

TRANSDERMAL

Date:07/26/01ISR Number: 3765393-7Report Type:Expedited (15-DaCompany Report #B0101345A  
Age:56 YR Gender:Male I/FU:F

Outcome Dose Death 150MG Per day	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Coronary Artery Disease	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
		Respiratory Disorder		Combivent	C		

RESPIRATORY  
( INHALATION)

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/26/01ISR Number: 3765394-9Report Type:Expedited (15-DaCompany Report #B0101353A

Age:67 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Cerebrovascular Accident	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice	Diabetes Mellitus					
per day						
UNKNOWN	75MG per day		Aspirin	C		
UNKNOWN	2PUFF Four		Salbutamol	C	Glaxo Wellcome	
times per day						
UNKNOWN	250MCG Twice		Fluticasone	C	Glaxo Wellcome	
per day						
UNKNOWN	75MG Twice		Diclofenac	C		
per day						
UNKNOWN	50MCG Twice		Salmeterol	C	Glaxo Wellcome	
per day						
UNKNOWN	200MCG Twice		Oxitropium	C		
per day						
UNKNOWN	10MG Per day		Imipramine	C		
UNKNOWN	40MG Per day		Frusemide	C		

Date:07/26/01ISR Number: 3765395-0Report Type:Expedited (15-DaCompany Report #B0101420A

Age:48 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Coronary Artery Disease		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice	Myocardial Infarction					
per day						

850MG Three

Metformin C

times per day

Human Insulin C  
Simvastatin C

20MG Per day

Date:07/26/01ISR Number: 3765396-2Report Type:Expedited (15-DaCompany Report #B0104495A  
Age:65 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300MG Per day	22 DAY	Blister	Zyban	PS	Glaxo Wellcome	ORAL
			Dermatitis	Cefaclor	C		ORAL
			Myocardial Infarction				

Date:07/26/01ISR Number: 3765398-6Report Type:Expedited (15-DaCompany Report #B0107823A  
Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Unknown	1 WK	Decreased Appetite	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged			Depression				
			Flat Affect				
			Intentional Misuse				

Date:07/26/01ISR Number: 3765399-8Report Type:Expedited (15-DaCompany Report #B0107824A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	10 DAY		Hypothermia	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged			Mood Altered	Aspirin	SS		
UNKNOWN			Social Avoidant Behaviour				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/26/01ISR Number: 3765400-1Report Type:Expedited (15-DaCompany Report #B0111133A  
Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Single	Lower Respiratory Tract		Zyban	PS	Glaxo Wellcome	
Hospitalization -	dose 1 DAY	Infection					
Initial or Prolonged	300MG per day	Malaise		Loperamide	C		
		Renal Failure		Allopurinol	C	Glaxo Wellcome	ORAL
		Urinary Tract Infection		Co-Proxamol	C		ORAL
		Vomiting		Meloxicam	C		ORAL
				Salbutamol	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)	200MCG Twice						
per day							

Date:07/26/01ISR Number: 3765401-3Report Type:Expedited (15-DaCompany Report #B0111862A  
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	7 DAY	Cardiac Failure		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -	Initial or Prolonged	Cerebral Artery Occlusion					
		Cheyne-Stokes Respiration					
		Crepitations					
		Hemiparesis					
		Ischaemic Stroke					
		Pulmonary Oedema					
		Ventricular Hypertrophy					

Date:07/26/01ISR Number: 3765423-2Report Type:Expedited (15-DaCompany Report #B0115263A  
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG	Angina Pectoris	Consumer	Zyban	PS	Glaxo Wellcome	ORAL



Initial or Prolonged Dyspepsia

Variable dose 6 DAY

Hypocalcaemia

UNKNOWN 2.5MG Per day

20MG Per day

75MG Twice

per day

10MG At night

20MG At night

90MG Twice

per day

UNKNOWN

25MG Per day

20MG Per day

2.5MG Per day

Bendrofluazide SS Glaxo Wellcome

Tamoxifen C ORAL

Aspirin C ORAL

Amitriptyline C ORAL

Simvastatin C ORAL

Diltiazem C ORAL

Glyceryl Trinitrate C Glaxo Wellcome

Atenolol C ORAL

Enalapril C ORAL

Bendrofluazide C Glaxo Wellcome ORAL

Date:07/26/01ISR Number: 3766258-7Report Type:Expedited (15-DaCompany Report #B0114651A

Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coronary Artery Disease Sudden Death	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG	ORAL						

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/26/01ISR Number: 3766317-9Report Type:Expedited (15-DaCompany Report #B0115005A  
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Foreign	Bupropion			
		Blood Pressure Increased		Hydrochloride	PS	Glaxo Wellcome Inc	
UNKNOWN	150 MG	Convulsion					
		Dysgeusia					
		Fatigue					
		Syncope					
		Weight Decreased					

Date:07/26/01ISR Number: 3766318-0Report Type:Expedited (15-DaCompany Report #B0114448A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Myocardial Infarction	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL		Pulmonary Haemorrhage					
		Pulmonary Oedema					
		Stress					

Date:07/26/01ISR Number: 3766319-2Report Type:Expedited (15-DaCompany Report #B0115008A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression	Foreign	Bupropion			
		Physical Assault	Literature	Hydrochloride	PS	Glaxo Wellcome Inc	
UNKNOWN	150 MG		Consumer				

Date:07/26/01ISR Number: 3766320-9Report Type:Expedited (15-DaCompany Report #B0114982A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Gastrointestinal Disorder	Health	Bupropion			

Initial or Prolonged      Hernia      Professional      Hydrochloride      PS      Glaxo Wellcome Inc      ORAL  
150

MG/VARIABLE

DOSE/ORAL

Date:07/26/01ISR Number: 3766321-0Report Type:Expedited (15-DaCompany Report #B0114829A

Age:50 YR      Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Hypothyroidism	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/ORAL				Hepatitis A Vaccine	C		
				Typhoid Vaccine	C		
				Salbutamol Sulphate	C		

Date:07/26/01ISR Number: 3766348-9Report Type:Expedited (15-DaCompany Report #B0114445A

Age:      Gender:Female      I/FU:I

Outcome	PT
Other	Arthralgia Decreased Activity Difficulty In Walking Dyspnoea Fatigue

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG		Headache Insomnia Malaise Nausea	Foreign	Zyban	PS	Glaxo Wellcome Inc	
		Pain In Extremity Polymyalgia Rheumatica Red Blood Cell Sedimentation Rate Increased Sedation Tremor	Consumer				

Date:07/26/01ISR Number: 3766361-1Report Type:Expedited (15-DaCompany Report #B0112965A  
Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Alanine Aminotransferase Increased	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
150 MG		Arthralgia Jaundice Lethargy Liver Function Test Abnormal Malaise Myalgia Red Blood Cell Sedimentation Rate Increased	Professional				

Date:07/26/01ISR Number: 3766420-3Report Type:Expedited (15-DaCompany Report #HQ3592224JUL2001  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion Drug Interaction	Health Professional	Effexor	PS	Wyeth Ayerst Laboratories	ORAL
150 MG 1 X			Other				
PER 1 DAY							

ORAL

Bupropion  
(Amfebutamone)

SS

ORAL

300 MG 1 X

PER 1 DAY

ORAL

Alprazolam  
(Alprazolam)

C

Date:07/26/01ISR Number: 3766427-6Report Type:Expedited (15-DaCompany Report #HQ3605224JUL2001  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction Serotonin Syndrome	Health Professional	Effexor Xr	PS	Wyeth Ayerst Laboratories	ORAL

420 MG 1X PER

1 DAY ORAL

Zyban (Amfebutamone  
Hydrochloride)

SS

ORAL

150 MG 1X PER

1 DAY ORAL 6 DAY

Epilim (Valproate  
Sodium)

C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/26/01ISR Number: 3766445-8Report Type:Expedited (15-DaCompany Report #B0114827A  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required							
150 MG/TWICE							
Intervention to							
PER DAY/ORAL							
Prevent Permanent Impairment/Damage							
		Nervous System Disorder	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Nystagmus					
		Vertigo		Fluticasone Propionate	C		
				Salbutamol Sulphate	C		
				Tramadol Hydrochloride	C		
				Lansoprazole	C		
				Frusemide	C		
				Tamoxifen	C		

Date:07/26/01ISR Number: 3766452-5Report Type:Expedited (15-DaCompany Report #B0110883A  
Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death							
150 MG/ORAL							
		Cardiomyopathy	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Left Ventricular Failure		Amitriptyline	C		
		Pulmonary Oedema		Nitrazepam	C		
		Sudden Death		Loperamide-Hydrochloride	C		
				Co-Codamol	C		

Date:07/26/01ISR Number: 3766465-3Report Type:Expedited (15-DaCompany Report #B0110446A  
Age:79 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged							
150 MG /							
		Atrial Flutter	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
		Dyspnoea	Professional				

TWICE PER DAY

/ ORAL

Date:07/26/01ISR Number: 3766467-7Report Type:Expedited (15-DaCompany Report #B0115003A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG		Hypertensive Crisis	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
				Irbesartan	C		
				Diltiazem	C		
				Calcium Carbonate	C		
				Hydrochlorothiazide	C		
				Multivitamin	C		
				Frusemide	C		

Date:07/26/01ISR Number: 3766468-9Report Type:Expedited (15-DaCompany Report #A0144968A  
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY		Gastrointestinal	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
		Haemorrhage	Health				
		Ulcer	Professional				
/ ORAL				Aspirin (Aspirin)	SS		
				Warfarin Sodium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/26/01ISR Number: 3766470-7Report Type:Expedited (15-DaCompany Report #B0114486A  
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Abdominal Pain Upper	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER Required DAY / ORAL		Blister					
Intervention to Prevent Permanent Impairment/Damage		Feeling Hot Headache Nausea Night Sweats Pain Tremor		Thyroxine Sodium Mebeverine	C C		

Date:07/26/01ISR Number: 3766471-9Report Type:Expedited (15-DaCompany Report #B0114664A  
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Anorexia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / TWICE PER DAY / ORAL		Diplopia Hyperhidrosis	Health Professional				
		Nausea Palpitations Vision Blurred		Proguanil Hydrochloride (Formulation Unknown)	SS		ORAL
100 MG / PER DAY / ORAL							

Date:07/26/01ISR Number: 3766472-0Report Type:Expedited (15-DaCompany Report #B0113971A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							



DAY / ORAL

Dihydrocodeine C  
Dothiepin  
Hydrochloride C  
Imipramine C

Date:07/26/01ISR Number: 3766473-2Report Type:Expedited (15-DaCompany Report #B0114491A  
Age:62 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 150 MG / ORAL	Duration Duodenal Ulcer	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Intervention to Prevent Permanent Impairment/Damage	Perforation		Trimipramine Theophylline Aspirin	C C C		

Date:07/26/01ISR Number: 3766530-0Report Type:Expedited (15-DaCompany Report #A0154610A  
Age:28 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG / PER	Duration Abortion Spontaneous	Study	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
DAY / ORAL	Complications Of Maternal Exposure To Therapeutic Drugs Pregnancy	Health Professional				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/26/01ISR Number: 3766586-5Report Type:Expedited (15-DaCompany Report #A0154846A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Study	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER		Complications Of Maternal	Health				
DAY/ORAL		Exposure To Therapeutic	Professional				
		Drugs					

Date:07/26/01ISR Number: 3766595-6Report Type:Expedited (15-DaCompany Report #A0150198A  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Facial Palsy	Health	Bupropion			
Intervention to			Professional	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER							
Prevent Permanent				Clonidine	C		
DAY/ ORAL				Oestradiol	C		
Impairment/Damage				Aspirin	C		

Date:07/26/01ISR Number: 3766602-0Report Type:Expedited (15-DaCompany Report #A0055937A  
Age:55 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Agoraphobia
Disability	Alopecia
	Anxiety
	Asthenia
	Autonomic Nervous System
	Imbalance
	Blood Glucose Increased
	Blood Pressure Increased
	Blood Triglycerides
	Increased
	Bradycardia
	Chills

Cognitive Disorder  
Confusional State  
Convulsion  
Cystocele  
Decreased Activity  
Diplopia  
Dizziness  
Dry Mouth  
Dysarthria  
Dyspepsia  
Dysthymic Disorder  
Enuresis  
Euphoric Mood  
Faeces Discoloured  
Fatigue  
Fear  
Feeling Abnormal  
Feeling Hot And Cold  
Fluid Retention  
Headache  
Hyperhidrosis  
Hypoaesthesia  
Insomnia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY / ORAL	Intervertebral Disc Protrusion Muscle Rigidity Muscle Spasms Muscle Twitching Musculoskeletal Stiffness	Health Professional	Bupropion Hydrochloride	PS	Eisai Inc	ORAL
	Neck Pain		Conjugated Estrogens	C		
	Nervousness		Vitamin	C		
	Panic Attack		Aspirin	C		
	Paraesthesia		Melatonin	C		
	Sciatica					
	Spinal X-Ray Abnormal					
	Stress					
	Throat Tightness					
	Tinnitus					
	Tongue Oedema					
	Transient Ischaemic Attack					
	Tremor					

Date:07/27/01ISR Number: 3766111-9Report Type:Expedited (15-DaCompany Report #B0100684A  
Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Zyban Lovan Alcohol	PS C C	Glaxo Wellcome	

Date:07/27/01ISR Number: 3766112-0Report Type:Expedited (15-DaCompany Report #B0100755A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Zyban	PS	Glaxo Wellcome	

Date:07/27/01ISR Number: 3766113-2Report Type:Expedited (15-DaCompany Report #B0101552A  
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bronchopneumonia Cardiac Arrest Coronary Artery Disease Myocardial Infarction Myocardial Ischaemia		Zyban	PS	Glaxo Wellcome	ORAL

Date:07/27/01ISR Number: 3766685-8Report Type:Expedited (15-DaCompany Report #B0094689A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150 MG / Required TWICE PER DAY Intervention to / ORAL Prevent Permanent Impairment/Damage	12 DAY	Agitation Insomnia Restlessness Weight Decreased	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/27/01ISR Number: 3766725-6Report Type:Expedited (15-DaCompany Report #A0124641A  
 Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG TWICE	Alcoholism	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
	PER DAY ORAL	Arrhythmia	Health				
		Dehydration	Professional				
		Heat Stroke					
		Sudden Death					

Date:07/27/01ISR Number: 3766727-XReport Type:Expedited (15-DaCompany Report #B0094547A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG TWICE	Blood Alcohol Increased Completed Suicide	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
	PER DAY ORAL		Professional				
			Company Representative				

Date:07/27/01ISR Number: 3766738-4Report Type:Expedited (15-DaCompany Report #B0103442A  
 Age:5 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG/TWICE	Atherosclerosis Dermatitis Myocardial Infarction	Foreign	Bupropion Hydrochloride (Zyban Tablet - Zyban	PS	Glaxo Wellcome Inc	ORAL
	PER DAY/ORAL			Prednisolone	C		
				Nitrazepam	C		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other ORAL		Ecchymosis Electroencephalogram Abnormal  Facial Bones Fracture Sedation Wrist Fracture	Foreign Health Professional  Company Representative	Bupropion Hydrochloride (Zyban Tablet-Zyban)	PS	Glaxo Wellcome Inc	ORAL

Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other  450 MG/PER DAY/ORAL	1 YR	Coronary Artery Disease	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bu7propion Hydrochloride)  Statins	PS  C	Glaxo Wellcome Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/27/01ISR Number: 3767234-0Report Type:Expedited (15-DaCompany Report #HQ3535223JUL2001

Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG 1 X	Concussion Drug Interaction	Health Professional	Effexor Xr	PS	Wyeth Ayerst Laboratories	ORAL
Other PER 1 DAY		Grand Mal Convulsion	Other	Zyban (Amfebutamone Hydrochloride, ) Unspecified Antianxiety Agent Valium (Diazepam)	SS C C		

Date:07/27/01ISR Number: 3767283-2Report Type:Expedited (15-DaCompany Report #HQ3601124JUL2001

Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 75 MG 1 X PER 1 DAY ORAL		Drug Interaction Feeling Hot	Health Professional	Effexor Xr	PS	Wyeth Ayerst Laboratories	ORAL
150 MG 1 X PER 1 DAY, 300 MG 1 X PER 1 DAY ORAL		Insomnia Malaise Serotonin Syndrome Tachycardia	Other	Zyban (Amfebutamone Hydrochloride)	SS		ORAL

Date:07/27/01ISR Number: 3767435-1Report Type:Periodic Company Report #A112580

Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Other	Libido Decreased	Consumer	Tikosyn	PS	Pfizer Pharmaceuticals Production Corp Ltd	ORAL
1000.00 MCG						
TOTAL BID						
ORAL						
300.00 MG			Wellbutrin	SS		ORAL
TOTAL BID						
ORAL						
100.00 MG			Lopressor	SS		ORAL
TOTAL BID						
ORAL						
			Vitamins	C		

Date:07/30/01ISR Number: 3766791-8Report Type:Expedited (15-DaCompany Report #A0150739A  
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Psychotic Disorder		Zyban	SS	Glaxo Wellcome	ORAL
2 MON		Sexual Offence		Prevacid	C		ORAL

Date:07/30/01ISR Number: 3766798-0Report Type:Expedited (15-DaCompany Report #B0089068A  
Age:32 YR Gender:Male I/FU:F

Outcome	PT
Death	Brain Oedema
	Cerebral Infarction



Death 150MG Per day	Coronary Artery Disease	Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY (INHALATION)	Mitral Valve Disease	Salbutamol	C	Glaxo Wellcome	
required	Myocardial Infarction				
100MCG per day	2PUFF As Ventricular Hypertrophy	Thyroxine	C	Glaxo Wellcome	ORAL
4G Per day		Glyceryl Trinitrate	C	Glaxo Wellcome	ORAL
20MG Per day		Candesartan	C		
20MG Per day		Fluoxetine	C		
40MG Per day		Omeprazole	C		
1TAB Per day		Frusemide	C		
		Didronel	C		ORAL

Date:07/30/01ISR Number: 3766804-3Report Type:Expedited (15-DaCompany Report #B0115085A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 3 DAY Initial or Prolonged	Amblyopia		Zyban	PS	Glaxo Wellcome	
	Burning Sensation Fatigue Myelitis Paraesthesia Peripheral Coldness Weight Decreased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/30/01ISR Number: 3767824-5Report Type:Expedited (15-DaCompany Report #B0111133A

Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Lower Respiratory Tract	Foreign	Zyban	PS	Glaxo Wellcome Inc	
150 MG / Hospitalization - SINGLE DOSE		Infection	Health				
Initial or Prolonged		Malaise	Professional	Loperamide			
		Renal Failure		Hydrochloride	C		
		Vomiting		Allopurinol	C		
				Co-Proxamol	C		
				Meloxicam	C		
				Salbutamol Sulphate	C		

Date:07/30/01ISR Number: 3767834-8Report Type:Expedited (15-DaCompany Report #B0100684A

Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Foreign	Zyban	PS	Glaxo Wellcome Inc	
150 MG				Fluoxetine			
				Hydrochloride	C		
				Ethanol	C		

Date:07/30/01ISR Number: 3767835-XReport Type:Expedited (15-DaCompany Report #B0101552A

Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bronchopneumonia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL		Cardiac Arrest					
		Coronary Artery Disease					
		Myocardial Infarction					
		Myocardial Ischaemia					

Date:07/30/01ISR Number: 3767854-3Report Type:Expedited (15-DaCompany Report #A0149272A

Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cough	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE		Haemoptysis	Health				
PER DAY/ ORAL		Respiratory Tract Neoplasm	Professional	Lansoprazole Oestradiol	C C		

Date:07/30/01ISR Number: 3767859-2Report Type:Expedited (15-DaCompany Report #B0107824A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Hypothermia	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
PER DAY/ ORAL 10 DAY		Mood Altered	Health	Aspirin (Formulation			
Initial or Prolonged		Overdose	Professional	Unknown) (Aspirin)	SS		
		Social Avoidant Behaviour					
		Stress					

Date:07/30/01ISR Number: 3767860-9Report Type:Expedited (15-DaCompany Report #B0107823A  
Age:20 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Decreased Appetite
Initial or Prolonged	Depression

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Overdose

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG/ ORAL	1 WK	Foreign Health Professional	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:07/30/01ISR Number: 3767861-0Report Type:Expedited (15-DaCompany Report #B0104495A  
Age:65 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blister	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
300 MG/ PER		Dermatitis					
DAY/ ORAL		Myocardial Infarction		Cefaclor	C		

Date:07/30/01ISR Number: 3767864-6Report Type:Expedited (15-DaCompany Report #B0115263A  
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Angina Pectoris	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/		Dyspepsia					
Initial or Prolonged		Hypocalcaemia					
VARIABLE				Bendrofluazide (Formulation Unknown)	SS		
DOSE/ ORAL				Tamoxifen	C		
2.5 MG/ PER				Aspirin	C		
DAY/				Amitriptyline	C		
				Simvastatin	C		
				Diltiazem	C		
				Nitroglycerin	C		

Atenolol C  
Enalapril C  
Bendrofluazide C

Date:07/30/01ISR Number: 3767866-XReport Type:Expedited (15-DaCompany Report #B0111862A

Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebral Artery Occlusion	Foreign	Bupropion			
Hospitalization - 150 MG/ORAL		Cerebral Artery		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Thrombosis Cheyne-Stokes Respiration Crepitations Hemiparesis Ischaemic Stroke Left Ventricular Failure Pulmonary Oedema Ventricular Hypertrophy					

Date:07/30/01ISR Number: 3767867-1Report Type:Expedited (15-DaCompany Report #B0101345A

Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coronary Artery Disease	Foreign	Bupropion			
150 MG /PER		Sudden Death		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

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Freedom Of Information (FOI) Report

DAY/ORAL

Combivent C

Date:07/30/01ISR Number: 3767868-3Report Type:Expedited (15-DaCompany Report #B0101420A  
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coronary Artery Disease Myocardial Infarction	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TWICE							
PER DAY/ORAL							

Metformin  
Hydrochloride C  
Human Insulin C  
Simvastatin C

Date:07/30/01ISR Number: 3767888-9Report Type:Expedited (15-DaCompany Report #B0101353A  
Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebrovascular Accident Diabetes Mellitus	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ TWICE							
PER DAY/ ORAL							

Aspirin C  
Salbutamol Sulphate C  
Fluticasone  
Propionate C  
Diclofenac C  
Salmeterol Xinafoate C  
Oxitropium Bromide C  
Imipramine C  
Frusemide C

Date:07/30/01ISR Number: 3767889-0Report Type:Expedited (15-DaCompany Report #B0099502A  
Age:53 YR Gender:Female I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG/ TWICE	Haemorrhagic Stroke	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
	PER DAY/ ORAL	Ruptured Cerebral Aneurysm		Estracombi	C		

Date:07/30/01ISR Number: 3767901-9Report Type:Expedited (15-DaCompany Report #B0100755A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG	Cardiac Arrest	Foreign Health	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
			Professional Company Representative				

Date:07/30/01ISR Number: 3768319-5Report Type:Expedited (15-DaCompany Report #A0125519A  
Age:49 YR Gender:Female I/FU:F

Outcome	PT
Death	Anxiety
Disability	Arrhythmia
	Cardiac Disorder
	Coronary Artery

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Hospitalization - 150MG Twice Initial or Prolonged per day	19 DAY	Arthralgia Dermatitis Hypogammaglobulinaemia Joint Swelling Liver Function Test Abnormal Red Blood Cell Sedimentation Rate Increased Serum Sickness Urticaria	Wellbutrin	PS	Glaxo Wellcome	ORAL
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Date:07/31/01ISR Number: 3767772-0Report Type:Expedited (15-DaCompany Report #B0097298A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	1 WK	Hypokalaemia Hyponatraemia Mental Disorder Polydipsia		Zyban  No Concurrent Medications	PS  C	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/31/01ISR Number: 3767777-XReport Type:Expedited (15-DaCompany Report #B0115262A  
Age:52 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300MG Per day Initial or Prolonged 1 DAY	Anaphylactic Reaction		Zyban	PS	Glaxo Wellcome	ORAL
200MG Per day			Amoxicillin	SS		ORAL
50MG Per day			Thyroxine	C	Glaxo Wellcome	ORAL
2.5MG Per day			Enalapril	C		ORAL
			Bendrofluazide	C	Glaxo Wellcome	ORAL

Date:07/31/01ISR Number: 3767779-3Report Type:Expedited (15-DaCompany Report #B0115681A  
Age:42 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG As Initial or Prolonged directed	Hypoaesthesia  Hypoaesthesia Oral  Motor Dysfunction Nervous System Disorder Pulse Abnormal		Zyban	PS	Glaxo Wellcome	

Date:07/31/01ISR Number: 3768626-6Report Type:Expedited (15-DaCompany Report #2013032  
Age:41 YR Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death	Abdominal Adhesions Hepatic Steatosis Malaise Oedema Peripheral Overdose Postoperative Adhesion Pulmonary Congestion Pulmonary Oedema Renal Cell Carcinoma	Health Professional Other	Morphine Sulfate (Similar To Andas 74-769 And 74-862) Morphine Sulfate (Similar To Nda 19-516) Oxycontin Cr Tablets, 40 Mg (Oxycodone	PS  SS		

PO	Stage Unspecified	Hydrochloride)	SS	ORAL
	Spinal Osteoarthritis	Oxycontin Cr		
	Toxicologic Test Abnormal	Tablets, 20 Mg		
		(Oxycodone		
PO		Hydrochloride)	SS	ORAL
		Bupropion	SS	
		Promethazine Hcl	SS	
		Acetaminophen	SS	
		Doxylamine	SS	
		Metoprolol	SS	
		Oxazepam	SS	
		Salicylic Acid	SS	
		Ephedrine	SS	
		Pseudophedrine Hcl	SS	
		Carbamazepine	SS	
		Trazodone Hcl	C	
		Prevacid		
		(Lansoprazole)	C	
		Diovan (Valsartan)	C	
		Baclofen (Lioresal)	C	
		Levoxyl	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/31/01ISR Number: 3777716-3Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #2001056168US

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Coordination Abnormal	Health	Celebrex	PS	Gd Searle And Co	ORAL
900 MG, QD,		Paranoia Speech Disorder	Professional	Lithobid (Lithium Carbonate)	SS		ORAL
ORAL				Seroquel (Quetiapine)	SS		ORAL
600 MG, QD,				Klonopin (Clonazepam)	SS		ORAL
ORAL				Wellbutrin-Slow Release (Amfebutamone Hydrochloride)	SS		ORAL
2 MG, QD,							
ORAL							
150 MG, QD,							
ORAL							

Date:08/01/01ISR Number: 3768424-3Report Type:Expedited (15-DaCompany Report #A0155404A  
Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:08/01/01ISR Number: 3768425-5Report Type:Expedited (15-DaCompany Report #A0155706A  
Age:58 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Systemic Lupus		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice		Erythematosis					
per day				Premarin	C		
				Tenormin	C		
				Norvasc	C		
				Cozaar	C		
				Aspirin	C		

Date:08/01/01ISR Number: 3768440-1Report Type:Expedited (15-DaCompany Report #B0115683A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blister		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day	25 DAY	Pruritus					

Date:08/01/01ISR Number: 3769202-1Report Type:Expedited (15-DaCompany Report #A0150739A  
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bipolar Disorder	Health	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
ORAL		Convulsion	Professional	Zyban Tablet - Zyban			
Required		Incest		(Bupropion			
Intervention to		Psychotic Disorder		Hydrochloride)	SS		ORAL
Prevent Permanent				Lansoprazole	C		
ORAL							
Impairment/Damage							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/01/01ISR Number: 3769593-1Report Type:Expedited (15-DaCompany Report #B0115085A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Influenza Like Illness Myelitis	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	

Date:08/01/01ISR Number: 3769603-1Report Type:Expedited (15-DaCompany Report #B0089069A

Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - ORAL Initial or Prolonged		Brain Stem Infarction Circulatory Collapse	Foreign	Bupropion Hydrochloride  Diclofenac Allopurinol	PS  C C	Glaxo Wellcome Inc	ORAL

Date:08/01/01ISR Number: 3769604-3Report Type:Expedited (15-DaCompany Report #B0090433A

Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150 MG / AS  DIRECTED /  ORAL		Coronary Artery Disease Myocardial Ischaemia  Sudden Death	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:08/01/01ISR Number: 3769605-5Report Type:Expedited (15-DaCompany Report #B0089068A

Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death ORAL		Cerebral Infarction Ruptured Cerebral	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL



Aneurysm  
Subarachnoid Haemorrhage

Date:08/01/01ISR Number: 3769606-7Report Type:Expedited (15-DaCompany Report #B0099815A  
Age:74 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Mitral Valve Disease Myocardial Infarction	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG / PER DAY / ORAL		Nausea		Salbutamol Sulphate Thyroxine Sodium Nitroglycerin Candesartan Cilexetil Fluoxetine Omeprazole Frusemide Disodium Etidronate	C C C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/02/01ISR Number: 3769152-0Report Type:Expedited (15-DaCompany Report #A0155912A  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Unknown	Completed Suicide		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Injury		Celexa	SS		ORAL
		Overdose		Alcohol	SS		ORAL
				Benadryl	SS		ORAL

Date:08/02/01ISR Number: 3769155-6Report Type:Expedited (15-DaCompany Report #B0084888A  
 Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	40MG At night	Coronary Artery Disease		Zyban	PS	Glaxo Wellcome	ORAL
Disability		Myocardial Infarction		Pravastatin	C		ORAL
Other	40MG per day	Pulmonary Infarction		Frusemide	C		ORAL
	150MG per day			Aspirin	C		ORAL
	20MG Per day			Omeprazole	C		ORAL
	5MG per day			Ramipril	C		ORAL

Date:08/02/01ISR Number: 3769161-1Report Type:Expedited (15-DaCompany Report #B0115845A  
 Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	21 DAY	Asthma		Zyban	PS	Glaxo Wellcome	
				Celecoxib	C		

Date:08/02/01ISR Number: 3769162-3Report Type:Expedited (15-DaCompany Report #B0115848A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day 6 DAY	Hyperventilation Panic Attack Tachycardia		Zyban	PS	Glaxo Wellcome	ORAL

Date:08/02/01ISR Number: 3769163-5Report Type:Expedited (15-DaCompany Report #B0115850A  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	13 DAY	Subarachnoid Haemorrhage		Zyban	PS	Glaxo Wellcome	

Date:08/02/01ISR Number: 3770225-7Report Type:Direct Company Report #  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1 BID	Drug Ineffective Fatigue Irritability		Wellbutrin 100 Mg Generic	PS		

Date:08/03/01ISR Number: 3770119-7Report Type:Expedited (15-DaCompany Report #A0155486A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	450MG Per day	Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Hypothyroidism Weight Decreased		Alcohol Unspecified	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Medications C

Date:08/03/01ISR Number: 3770127-6Report Type:Expedited (15-DaCompany Report #B0114485A  
Age:66 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day 5 DAY	Dyspnoea		Zyban	PS	Glaxo Wellcome	
Hospitalization - Initial or Prolonged		Myocardial Infarction					

Date:08/03/01ISR Number: 3770843-6Report Type:Direct Company Report #  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN		Convulsion		Benztropine	PS		
Initial or Prolonged UNKNOWN		Heat Stroke		Bupropion	SS		

Date:08/03/01ISR Number: 3770900-4Report Type:Expedited (15-DaCompany Report #WAES 01072442  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	25 MG/DAILY/ORAL	Coordination Abnormal Drug Interaction Feeling Abnormal	Health Professional Company	Vioxx 25 Mg	PS	Merck Research Laboratories Div Merck Co Inc	ORAL
		Gait Disturbance	Representative				
		Speech Disorder		Anafranil	SS		
		Vision Blurred		Lamictal	SS		
				Dipentum	SS		
				Zoloft	SS		
				Premarin	SS		
				Wellbutrin	SS		
				Klonopin	C		
				Motrin	C		

Date:08/03/01ISR Number: 3771020-5Report Type:Expedited (15-DaCompany Report #A0155404A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Study	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
ORAL		Complications Of Maternal Exposure To Therapeutic Drugs	Health Professional				

Date:08/03/01ISR Number: 3771022-9Report Type:Expedited (15-DaCompany Report #A0155706A  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Renal Impairment	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE		Systemic Lupus	Professional				
PER DAY/ORAL		Erythematosis	Company Representative	Conjugated Estrogens Atenolol Amlodipine Losartan Potassium Aspirin	C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/03/01ISR Number: 3771058-8Report Type:Expedited (15-DaCompany Report #B0115683A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 300 MG PER Prevent Permanent DAY/ORAL Impairment/Damage		Blister Pruritus	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:08/03/01ISR Number: 3771073-4Report Type:Expedited (15-DaCompany Report #S01-USA-01399-01  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 20 MG QD PO		Pancytopenia	Health Professional	Celexa	PS	Forest Laboratories Inc	ORAL
150 MG QD			Other	Wellbutrin (Amfebutamone Hydrochloride)	SS		
				Lithium	C		

Date:08/03/01ISR Number: 3771172-7Report Type:Expedited (15-DaCompany Report #B0097298A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG / TWICE PER DAY / ORAL	1 WK	Hypokalaemia Hyponatraemia Polydipsia	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL

Date:08/03/01ISR Number: 3771179-XReport Type:Expedited (15-DaCompany Report #A0155843A  
Age:26 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/ TWICE		Dermatitis	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged PER DAY/ ORAL		Hypogammaglobulinaemia	Health				
		Liver Function Test Abnormal	Professional				
		Red Blood Cell Sedimentation Rate Increased					
		Serum Sickness					

Date:08/03/01ISR Number: 3771181-8Report Type:Expedited (15-DaCompany Report #B0115262A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG/ PER		Anaphylactic Reaction	Foreign	Zyban (Bupropion Hydrochloride)	PS	Glaxo Wellcome Inc	ORAL
DAY/ ORAL				Amoxicillin (Amoxicillin)	SS		ORAL
50 MG/ ORAL				Thyroxine Sodium	C		
				Enalapril	C		
				Bendrofluazide	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/03/01ISR Number: 3771186-7Report Type:Expedited (15-DaCompany Report #B0115681A  
Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/ AS	Brain Damage Hypoaesthesia Hypoaesthesia Oral	Foreign Consumer	Zyban (Bupropion Hydrochloride)	PS	Glaxo Wellcome Inc	
DIRECTED/ UNKNOWN	Motor Dysfunction					

Date:08/03/01ISR Number: 3771344-1Report Type:Expedited (15-DaCompany Report #HQ3535223JUL2001  
Age:41 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG 1 X	Drug Interaction Grand Mal Convulsion	Foreign Health	Effexor Xr	PS	Wyeth Ayerst Laboratories	ORAL
Other PER 1 DAY	Head Injury	Professional				
ORAL		Other	Zyban (Amfebutamone Hydrochloride)	SS		
			Unspecified Antianxiety Agent (Unspecified Antianxiety Agent)	C		
			Valium (Diazepam)	C		

Date:08/03/01ISR Number: 3771553-1Report Type:Expedited (15-DaCompany Report #A0153280A  
Age:20 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY	Amnesia Grand Mal Convulsion	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL



/ ORAL	Medication Error	Company				
	Overdose	Representative	Ethanol (Formulation Unknown) (Alcohol)	SS		

Date:08/03/01ISR Number: 3771554-3Report Type:Expedited (15-DaCompany Report #A0144419A  
 Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Convulsion					
150 MG / PER		Neuropathy Peripheral	Professional				
DAY ORAL			Company Representative				

Date:08/06/01ISR Number: 3770867-9Report Type:Expedited (15-DaCompany Report #A0139670A  
 Age:74 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Anorexia		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Per day	10 MON						
Initial or Prolonged		Cardiac Failure		Rocephin	C		
INTRAVENOUS	1G Per day	8 DAY					
20MG Per day		Congestive		Pepcid	C		
		Condition Aggravated		Kcl	C		
12.5MG Twice		Convulsion		Captopril	C	Glaxo Wellcome	
per day		Depressed Level Of					
		Consciousness		Depakote	C		
500MG Per day	2 YR						
		Dyskinesia		Buspar	C		
10MG Twice		Loss Of Consciousness					
per day	2 YR						
7MG At night	1 DAY			Nicoderm Patch	C		

Freedom Of Information (FOI) Report

75MG Per day	5	YR	Seroquel	C	
			Lasix	C	
			Lanoxin	C	Glaxo Wellcome
			Albuterol	C	Glaxo Wellcome
			Spirololactone	C	
12.5MG Per day			Aspirin	C	
325MG Per day	5	YR	Antivert	C	
12.5MG As required			Artificial Tears	C	
			Proventil	C	Glaxo Wellcome
RESPIRATORY (INHALATION)		YR	Atrovent	C	Glaxo Wellcome
RESPIRATORY (INHALATION)		YR	Fluoroquinolone	C	
INTRAVENOUS					

Date:08/06/01ISR Number: 3770871-0Report Type:Expedited (15-DaCompany Report #A0154004A  
 Age:26 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blister		Zyban	PS	Glaxo Wellcome	ORAL
Other		Burning Sensation		No Concurrent Medication	C		
150MG See text	3 DAY	Erythema Insomnia Pain Pruritus Scab Scar Sensory Loss Skin Discolouration Swelling					

Date:08/06/01ISR Number: 3771519-1Report Type:Expedited (15-DaCompany Report #2013643

Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Atherosclerosis	Health	Oxycontin	PS	Purdue Pharma Lp	ORAL
MG PO		Bronchopneumonia	Professional	Amitriptyline	SS		
MG		Cardiac Arrest	Other	Cyclobenzaprine Hcl	SS		
MG		Cardiomegaly Cerebral Ischaemia		Wellbutrin (Bupropion)	SS		
MG		Foreign Body Aspiration		Diazepam	SS		
MG		Hepatic Steatosis		Lidocaine Hcl	SS		
MG		Hepatomegaly Obstructive Airways Disorder Splenomegaly Toxicologic Test Abnormal					

Date:08/06/01ISR Number: 3771653-6Report Type:Expedited (15-DaCompany Report #B0115845A

Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Asthma	Foreign	Zyban	PS	Glaxo Wellcome Inc	
150 MG	21 DAY			Celecoxib	C		

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Freedom Of Information (FOI) Report

Date:08/06/01ISR Number: 3771654-8Report Type:Expedited (15-DaCompany Report #B0115850A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Subarachnoid Haemorrhage	Foreign	Zyban	PS	Glaxo Wellcome Inc	
150 MG							

Date:08/06/01ISR Number: 3771657-3Report Type:Expedited (15-DaCompany Report #B0115848A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Hyperventilation	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ PER							
		Panic Attack					
DAY/ ORAL		Tachycardia					

Date:08/06/01ISR Number: 3771661-5Report Type:Expedited (15-DaCompany Report #B0084888A  
Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coronary Artery Disease	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
AS DIRECTED/ Disability		Coronary Artery Occlusion	Health				
ORAL							
Required		Myocardial Infarction	Professional	Pravastatin	C		
Intervention to		Pulmonary Infarction		Frusemide	C		
Prevent Permanent				Aspirin	C		
Impairment/Damage				Omeprazole	C		
				Ramipril	C		

Date:08/06/01ISR Number: 3772038-9Report Type:Expedited (15-DaCompany Report #B0114485A  
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Dyspnoea	Foreign	Bupropion			

Hospitalization - Myocardial Infarction Health Hydrochloride PS Glaxo Wellcome Inc  
 150 MG/PER  
 Initial or Prolonged Professional  
 DAY

Date:08/06/01ISR Number: 3772198-XReport Type:Expedited (15-DaCompany Report #A0155486A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 450 MG / PER Initial or Prolonged DAY / ORAL		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
		Hypothyroidism	Professional				
		Weight Decreased	Company Representative	Ethanol (Formulation Unknown) Alcohol	SS		

Date:08/06/01ISR Number: 3772342-4Report Type:Expedited (15-DaCompany Report #2000022913-1  
 Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 500 Prevent Permanent MILLIGRAMS Impairment/Damage 3.0 DAILY ORAL	37 DAY	Blood Human Chorionic Gonadotropin Negative	Consumer Health	Famvir	PS	Novartis Pharmaceuticals Corp	ORAL
		Laboratory Test Abnormal	Professional				
		Medication Error					
		Ovarian Cyst Vomiting		Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
	4 YR			Claritin (Loratadine)	C		

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Freedom Of Information (FOI) Report

Date:08/06/01ISR Number: 3772343-6Report Type:Expedited (15-DaCompany Report #A0155912A  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
150 MG ORAL		Injury	Professional	Citalopram			
		Intentional Misuse		Hydrobromide Tablet			
		Toxicologic Test Abnormal		(Citalopram			
ORAL				Hydrobromide)	SS		ORAL
				Ethanol Liquid			
ORAL				(Alcohol)	SS		
				Diphenhydramine Hcl			
				(Formulation			
				Unknown)			
				(Diphenhydramine			
ORAL				Hcl)	SS		ORAL

Date:08/07/01ISR Number: 3771426-4Report Type:Expedited (15-DaCompany Report #A0150080A  
 Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Feeling Cold		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day		Hypoaesthesia					

Date:08/07/01ISR Number: 3771428-8Report Type:Expedited (15-DaCompany Report #A0153069A  
 Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Amnesia		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG In the		Anxiety					
morning	36 DAY	Aphasia		No Concurrent			
		Coordination Abnormal		Medications	C		
		Depression					

Insomnia  
Nervousness  
Neuromyopathy  
Speech Disorder  
Weight Increased

Date:08/07/01ISR Number: 3771432-XReport Type:Expedited (15-DaCompany Report #A0156108A  
Age:45 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300MG per day Initial or Prolonged	Coagulopathy Condition Aggravated Ecchymosis Endometriosis Menorrhagia Stress		Wellbutrin Sr Herbs Homeopathy	PS C C	Glaxo Wellcome	ORAL

Date:08/07/01ISR Number: 3771437-9Report Type:Expedited (15-DaCompany Report #B0111329A  
Age:44 YR Gender:Male I/FU:F

Outcome	PT
Death	Hepatic Congestion Hepatomegaly Hyperplasia Myocardial Infarction

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Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
55	DAY	Pulmonary Haemorrhage Pulmonary Oedema Thyroid Neoplasm	Zyntabac	PS	Glaxo Wellcome	ORAL

Date:08/07/01ISR Number: 3771440-9Report Type:Expedited (15-DaCompany Report #B0114448A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Death	55 DAY	Hepatic Congestion	Zyntabac	PS	Glaxo Wellcome	ORAL
		Hepatomegaly Hyperplasia Myocardial Infarction Pulmonary Haemorrhage Pulmonary Oedema Stress Thyroid Neoplasm				

Date:08/07/01ISR Number: 3771448-3Report Type:Expedited (15-DaCompany Report #B0115843A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Death	1 DAY	Back Pain	Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -		Urinary Tract Infection	Dihydrocodeine	SS		ORAL
Initial or Prolonged		Vomiting	Estracombi	C		
TRANSDERMAL						

Date:08/07/01ISR Number: 3771449-5Report Type:Expedited (15-DaCompany Report #B0116004A  
Age:54 YR Gender:Male I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Depression Mental Disorder	Zyban	PS	Glaxo Wellcome	ORAL



Stress

Date:08/07/01ISR Number: 3771450-1Report Type:Expedited (15-DaCompany Report #B0116186A  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Twice	Abnormal Behaviour		Zyban	PS	Glaxo Wellcome	ORAL
	per day	Aggression					
	200MCG per	Disinhibition		Thyroxine	C	Glaxo Wellcome	ORAL
	day			Co-Codamol	C		ORAL

Date:08/08/01ISR Number: 3771889-4Report Type:Expedited (15-DaCompany Report #B0116466A  
 Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG	Dizziness		Zyban	PS	Glaxo Wellcome	
	Variable dose	Headache					
	3MG Three	Insomnia		Bromazepam	C		ORAL
	times per day	Petit Mal Epilepsy					
	100MG Twice	Syncope		Gtn	C	Glaxo Wellcome	ORAL
	per day	Tremor					
	100MG Per day			Atenolol	C		ORAL

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625MCG Three  
 times per  
 week

Conjugated  
 Oestrogens C ORAL

Date:08/08/01ISR Number: 3772814-2Report Type:Expedited (15-DaCompany Report #A0154004A  
 Age:26 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blister	Consumer	Bupropion			
Other		Burning Sensation		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/SEE		Erythema					
TEXT/ORAL		Hypoaesthesia					
		Oedema Peripheral					
		Pain					
		Pruritus					
		Scab					
		Scar					
		Skin Discolouration					
		Ulcer					

Date:08/08/01ISR Number: 3772858-0Report Type:Expedited (15-DaCompany Report #A0139670A  
 Age:74 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Hospitalization -		Cardiac Failure	Professional				
150 MG/ PER		Convulsion		Ceftriaxone Sodium	C		
Initial or Prolonged		Depressed Level Of		Famotidine	C		
DAY/ ORAL		Consciousness		Potassium Chloride	C		
				Captopril	C		
				Semisodium Valproate	C		
				Buspirone			
				Hydrochloride	C		

Nicotine C  
 Quetiapine Fumarate C  
 Frusemide C  
 Digoxin C  
 Salbutamol Sulphate C  
 Spironolactone C  
 Aspirin C  
 Meclozine  
 Hydrochloride C  
 Hypromellose C  
 Salbutamol Sulphate C  
 Ipratropium Bromide C  
 Fluoroquinolone C

Date:08/08/01ISR Number: 3773455-3Report Type:Expedited (15-DaCompany Report #PHBS2001US07798  
 Age:10 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Interaction Drug Level Above	Literature Health	Sandimmune	PS	Novartis Pharmaceuticals Corp	ORAL
420 MG/D, ORAL	22 DAY	Therapeutic Drug Level Below Therapeutic	Professional	Bupropion (Amfebutamone)	SS		
75 MG, BID	22 DAY			Methylphenidate			

Freedom Of Information (FOI) Report

5 MG, BID 22 DAY

(Methylphenidate Hydrochloride)	SS
Azathioprine (Azathioprine)	C
Prednisone	C
Dipyridamole (Dipyridamole)	C
Nifedipine	C
Sulfamethoxazole W/Trimethoprim (Sulfamethoxazole)	C
Calcium Carbonate	C
Aluminium Hyrdoxide W/Magnesium Hydroxide	C

Date:08/08/01ISR Number: 3773483-8Report Type:Expedited (15-DaCompany Report #A0155184A  
Age:10 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	75 MG / TWICE	Drug Interaction	Literature	Wellbutrin	PS	Glaxo Wellcome Inc	ORAL
PER DAY /		Drug Level Below	Health				
ORAL		Therapeutic	Professional				

420 MG/DAY/

Cyclosporin (Formulation Unknown) (Cyclosporine)	SS
Azathioprine	C
Prednisone	C
Dipyridamole	C
Nifedipine	C
Co-Trimoxazole	C
Calcium Carbonate	C
Aluminium Salt+ Mg(OH)2	C

Date:08/09/01ISR Number: 3772945-7Report Type:Expedited (15-DaCompany Report #A0152291A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG Twice		Blood Cholesterol		Bupropion	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	265 DAY	Increased					
1TAB Per day		Cholelithiasis		Orthotricyclen	C		ORAL
		Gallbladder Disorder Weight Decreased					

Date:08/09/01ISR Number: 3772948-2Report Type:Expedited (15-DaCompany Report #A0156426A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other 150MG Twice		Cardiac Failure		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
per day	3 YR	Congestive					
1 MON				Amantadine	SS		
				Selegiline	C		

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Date:08/09/01ISR Number: 3772952-4Report Type:Expedited (15-DaCompany Report #A0156690A  
 Age:36 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice Initial or Prolonged per day	Blood Pressure Decreased Dermatitis Hypersensitivity Loss Of Consciousness Pruritus Urticaria Viral Infection		Zyban	PS	Glaxo Wellcome	ORAL

Date:08/09/01ISR Number: 3773983-0Report Type:Expedited (15-DaCompany Report #B0116004A  
 Age:54 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death ORAL	Completed Suicide Depression Mental Disorder Stress	Foreign Literature	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:08/09/01ISR Number: 3773986-6Report Type:Expedited (15-DaCompany Report #B0116186A  
 Age:40 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150 MG / TWICE PER DAY / ORAL	Abnormal Behaviour Aggression Disinhibition	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
			Thyroxine Sodium Co-Codamol	C C		

Date:08/09/01ISR Number: 3774090-3Report Type:Expedited (15-DaCompany Report #B0115843A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Urinary Tract Infection	Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG / ORAL							
Hospitalization - Initial or Prolonged		Vomiting		Dihydrocodeine Tablet (Dihydrocodeine)	SS		ORAL
2 TABLET /							
ORAL				Estracombi	C		

Date:08/09/01ISR Number: 3774208-2Report Type:Expedited (15-DaCompany Report #A0150080A  
Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hypoaesthesia Peripheral Coldness	Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/PER							
DAY/ORAL							

Date:08/09/01ISR Number: 3774220-3Report Type:Expedited (15-DaCompany Report #B0111329A  
Age:44 YR Gender:Male I/FU:F

Outcome	PT
Death	Acute Stress Disorder Hepatomegaly Myocardial Infarction

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Pulmonary Haemorrhage Pulmonary Oedema Thyroid Neoplasm	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
150 MG/ ORAL			Health Professional				

Date:08/09/01ISR Number: 3774221-5Report Type:Expedited (15-DaCompany Report #B0114448A  
Age:44 YR Gender:Male I/FU:F

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration		Foreign	Zyban	PS	Glaxo Wellcome Inc	ORAL
Dose		Hepatomegaly					
Death		Hyperplasia Myocardial Infarction Pulmonary Haemorrhage Pulmonary Oedema Thyroid Neoplasm					
150 MG/ ORAL							

Date:08/09/01ISR Number: 3774242-2Report Type:Expedited (15-DaCompany Report #A0153069A  
Age:43 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration		Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Dose		Amnesia					
Other		Anxiety					
150 MG/ IN		Depression					
THE MORNING/ ORAL		Insomnia Neuromyopathy Weight Increased					

Date:08/09/01ISR Number: 3774246-XReport Type:Expedited (15-DaCompany Report #WAES 01072442  
Age:42 YR Gender:Female I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphasia Coordination Abnormal Difficulty In Walking	Health Professional Company	Vioxx	PS	Merck Research Laboratories Div Merck Co Inc	ORAL
25							
		Disorientation	Representative				
MG/DAILY/ORAL		Drug Interaction Speech Disorder Vision Blurred		Anafranil Klonopin Lamictal Wellbutrin Dipentum Premarin Zoloft Coumadin Motrin Lithium Carbonate	SS SS SS SS SS SS SS C C C		

Date:08/09/01ISR Number: 3774459-7Report Type:Expedited (15-DaCompany Report #WAES 01072531  
Age:37 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Spinal Disorder	Consumer Health Professional	Vioxx	PS	Merck Research Laboratories Div Merck Co Inc	ORAL
25							
MG/DAILY/PO				Wellbutrin	SS		
UNKNOWN							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Delestrogen C  
 Flexeril C  
 Lipitor C  
 Clonazepam C

Date:08/09/01ISR Number: 3774530-XReport Type:Expedited (15-DaCompany Report #A0156108A  
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Acute Stress Disorder	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged		Condition Aggravated Endometrial Disorder Increased Tendency To Bruise Menstrual Disorder		Herbal Medication Homeopathic Medication	C C		

Date:08/09/01ISR Number: 3774582-7Report Type:Expedited (15-DaCompany Report #WAES 01072531  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion Spinal Disorder	Consumer Health Professional	Vioxx	PS	Merck Research Laboratories Div Merck Co Inc	ORAL
25 MG DAILY							
PO				Wellbutrin	SS		
UNKNOWN				Delestrogen Flexeril Lipitor Clonazepam	C C C C		

Date:08/09/01ISR Number: 3774888-1Report Type:Expedited (15-DaCompany Report #A114068  
 Age:65 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Life-Threatening Hospitalization - Initial or Prolonged 300.00 MG	Arrhythmia Cardio-Respiratory Arrest Coma  Convulsion  Drug Interaction	Consumer Health Professional	Viagra  Bupropion	PS  SS	Pfizer Agricultural Div	ORAL
TOTAL: BID  ORAL  0.40 MG	Fall  Head Injury  Tachycardia		Tamsulosin	SS		ORAL
TOTAL: DAILY: 0  RAL			Atorvastatin Lorazepam Aspirin Dimethyl Sulfone Ubidecarenone Multivitamin Fish Oil Pentatonic Acid	C C C C C C C C		

Date: 08/09/01  
 ISR Number: 3775286-7  
 Report Type: Expedited (15-DaCompany Report #B0116466A)  
 Age: 49 YR Gender: Female I/FU: I

Outcome PT  
 Other Dizziness  
 Headache

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Insomnia Syncope Tremor	Report Source	Product	Role	Manufacturer	Route
150 MG	VARIABLE DOSE;		Foreign Consumer	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	
				Bromazepam	C		
				Nitroglycerin	C		
				Atenolol	C		
				Conjugated Estrogens	C		

Date:08/10/01ISR Number: 3773873-3Report Type:Expedited (15-DaCompany Report #A0127152A  
Age: Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day		Drug Level Above		Wellbutrin	PS	Glaxo Wellcome	ORAL
	100MG Twice per day		Therapeutic		Zoloft	SS		ORAL
			Overdose		Unspecified Medication	C		

Date:08/10/01ISR Number: 3773874-5Report Type:Expedited (15-DaCompany Report #A0154589A  
Age:29 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	14 DAY		Vision Blurred		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:08/11/01ISR Number: 3779413-7Report Type:Expedited (15-DaCompany Report #NSADSS2001012819  
Age:25 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Grand Mal Convulsion	Foreign Study Health Professional	Remicade (5 Mg/Ml Lyophilized Powder) (Infliximab, Recombinant)	PS		
INTRAVENOUS	5 MG/KG, 1 IN						
1 TIME(S), IV				Wellbutrin (Amfebutamone Hydrochloride)	SS		
				Pentasa (Mesalazine)	C		
				Losec (Omeprazole)	C		
				Buscopan (Hyoscine Butylbromide)	C		
				Demerol (Pethidine Hydrochloride)	C		
				Maxeran (Metoclopramide Hydrochloride)	C		
				Vitamin B12 (Cyanocobalamin)	C		
				Entocort (Budesonide)	C		
				Imuran (Azathioprine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/13/01ISR Number: 3774994-1Report Type:Expedited (15-DaCompany Report #A0157043A  
Age:68 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 50MG Per day Initial or Prolonged	Atrial Fibrillation		Wellbutrin Sr Flomax	PS C	Glaxo Wellcome	ORAL

Date:08/13/01ISR Number: 3774998-9Report Type:Expedited (15-DaCompany Report #B0109588A  
Age:51 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150MG Twice Hospitalization - per day Initial or Prolonged 150MG per day Other	Arrhythmia Asthenia Blood Pressure Decreased Dizziness Dyspnoea Feeling Abnormal Insomnia Pallor Palpitations Pulse Absent Supraventricular Tachycardia Syncope		Zyban Aspirin Ketoprofen Alcohol	PS C C C	Glaxo Wellcome	ORAL

Date:08/13/01ISR Number: 3775001-7Report Type:Expedited (15-DaCompany Report #B0114323A  
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 100MG Three times per day 2 YR	Balance Disorder Fall Tooth Loss		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:08/13/01ISR Number: 3775002-9Report Type:Expedited (15-DaCompany Report #B0116763A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	6	DAY					

Date:08/13/01ISR Number: 3775003-0Report Type:Expedited (15-DaCompany Report #B0116765A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Bipolar Disorder		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	34	DAY					
Initial or Prolonged		Hypomania		Amoxicillin	C		ORAL
500MG Per day	13	DAY		Alcohol	C		ORAL
		Mania					

Date:08/13/01ISR Number: 3776009-8Report Type:Expedited (15-DaCompany Report #PHBS2001US07876  
Age:4 MON Gender:Female I/FU:I

Outcome	PT
Other	Abdominal Pain Upper Complications Of Maternal Exposure To Therapeutic Drugs Oesophageal Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vomiting

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Health Professional	Famvir	PS	Novartis Pharmaceuticals Corp	
TRANSPLACENTAL	TRANSPLACENTA					
L			Wellbutrin (Amfebutamone Hydrochloride)	SS		
TRANSPLACENTAL	TRANSPLACENTA					
L			Claritin (Loratadine)	C		

Date:08/13/01ISR Number: 3776036-0Report Type:Expedited (15-DaCompany Report #A0156690A  
 Age:36 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY / ORAL	Blood Pressure Decreased Dermatitis Hypersensitivity Loss Of Consciousness Pruritus Urticaria Viral Infection	Foreign Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Date:08/13/01ISR Number: 3776258-9Report Type:Expedited (15-DaCompany Report #A0127152A  
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death 250 MG / PER DAY / ORAL	Drug Level Above Therapeutic	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL



Overdose

Sertraline  
Hydrochloride  
(Formulation  
Unknown) (Sertraline  
Hydrochloride)

SS

ORAL

100 MG /

TWICE PER DAY

/ ORAL

Date:08/13/01ISR Number: 3776316-9Report Type:Expedited (15-DaCompany Report #A0156426A

Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG /		Cardiac Failure Congestive	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL

TWICE PER DAY

/ ORAL 3 YR

Amantadine  
(Amantadine  
Hydrochloride)

SS

1 MON

Selegiline

C

Date:08/13/01ISR Number: 3776415-1Report Type:Expedited (15-DaCompany Report #A0152291A

Age:35 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Blood Cholesterol Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Cholecystectomy Cholelithiasis Gallbladder Disorder	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ORAL		Pain	Study	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
		Weight Decreased	Health				
			Professional	Ortho Tri-Cyclen	C		

Date:08/13/01ISR Number: 3776417-5Report Type:Expedited (15-DaCompany Report #A0154589A  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 150 MG /ORAL		Vision Blurred	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Intervention to Prevent Permanent Impairment/Damage			Professional Company Representative				

Date:08/14/01ISR Number: 3776187-0Report Type:Expedited (15-DaCompany Report #A0156593A  
Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day		Convulsion Convulsive Threshold Lowered		Wellbutrin Narcotics	PS SS	Glaxo Wellcome	ORAL

Date:08/14/01ISR Number: 3776188-2Report Type:Expedited (15-DaCompany Report #A0156711A  
Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day		Abortion Spontaneous		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Complications Of Maternal Exposure To Therapeutic Drugs		Prozac	C		

Date:08/14/01ISR Number: 3776189-4Report Type:Expedited (15-DaCompany Report #A0157266A  
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day 7 DAY Initial or Prolonged		Agitation Atrial Fibrillation Nervousness		Zyban	PS	Glaxo Wellcome	ORAL

Date:08/14/01ISR Number: 3776196-1Report Type:Expedited (15-DaCompany Report #B0116926A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Constipation Neoplasm Malignant		Zyban	PS	Glaxo Wellcome	

Date:08/15/01ISR Number: 3777433-XReport Type:Expedited (15-DaCompany Report #A0157251A  
Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day	2 WK	Laboratory Test Abnormal Multiple Sclerosis		Zyban  Premarin	PS  C	Glaxo Wellcome	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vitamin E C  
 Vitamin C C  
 Vitamin A C  
 Multivitamin C

Date:08/15/01ISR Number: 3777434-1Report Type:Expedited (15-DaCompany Report #B0113169A  
 Age:65 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice		Aortic Valve Stenosis	Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - per day	Initial or Prolonged		Cardiac Failure				
			Coronary Artery Disease				
			Decreased Appetite				
			Dry Throat				
			Dyspnoea				
			Oedema Peripheral				
			Retching				

Date:08/15/01ISR Number: 3779307-7Report Type:Periodic Company Report #2001011456-1  
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	60 MILLIGRAMS		Palpitations	Paxil	PS	Smithkline Beecham	ORAL
	1.0 DAILY		Health				
	ORAL		Professional				
	112 DAY			Wellbutrin Sr (Bupropion Hcl)	SS		ORAL
	200						
	MILLIGRAMS						
	2.0 DAILAY						
	ORAL						
	87 DAY						

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged PO	Cardiac Disorder Cardiomyopathy	Health Professional	Librium	PS	Inc Pharmaceuticals Inc	ORAL
100 MG SEE TEXT PO	Chest Pain Dyspnoea Electrocardiogram Qt Prolonged		Wellbutrin Tablet-Controlled Release	SS		ORAL
20 MG SEE TEXT PO	Intentional Misuse Laboratory Test Abnormal Peripheral Ischaemia		Mirtazapin Tablet (Mirtazapine)	SS		ORAL
	Syncope Tricuspid Valve Incompetence					

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150MG Twice Hospitalization - per day 18 DAY Initial or Prolonged	Hypersensitivity Hypotension Loss Of Consciousness Viral Infection		Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/16/01ISR Number: 3778453-1Report Type:Direct  
Age:38 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Arthritis		Wellburtin 150 Sr	PS		ORAL
150 PO BID		Prostration		Atenolol 25	C		
		Serum Sickness					
		Urticaria					

Date:08/16/01ISR Number: 3778665-7Report Type:Expedited (15-DaCompany Report #WAES 01072442  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Acute Stress Disorder	Health	Vioxx	PS	Merck Research	
		Aphasia	Professional			Laboratories Div	
		Coordination Abnormal	Company			Merck Co Inc	ORAL
25		Difficulty In Walking	Representative				
MG/DAILY/ORAL		Disorientation		Anafranil	SS		
		Drug Interaction		Klonopin	SS		
		Transient Ischaemic		Lamictal	SS		
		Attack		Wellbutrin	SS		
		Vision Blurred		Dipentum	SS		
				Premarin	SS		
				Zoloft	SS		
				Coumadin	C		
				Motrin	C		
				Lithium Carbonate	C		

Date:08/16/01ISR Number: 3779412-5Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US CDC 0426000002

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Angiopathy	Health	Librium	PS	Incn Pharmaceuticals	
Initial or Prolonged		Cardiac Disorder	Professional			Inc	ORAL
PO		Cardiomyopathy		Wellbutrin			
		Chest Pain		Tablet-Controlled			

100 MG SEE

Dyspnoea

Release

SS

ORAL

Electrocardiogram Qt

TEXT PO

Prolonged

Intentional Misuse

Laboratory Test Abnormal

Syncope

Tricuspid Valve

Incompetence

Date:08/16/01ISR Number: 3781852-5Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US CDC 0426000003\*

Outcome  
Hospitalization -  
Initial or Prolonged

PT  
Cardiac Disorder  
Cardiomyopathy  
Chest Pain  
Dyspnoea  
Electrocardiogram Qt  
Prolonged  
Intentional Misuse  
Laboratory Test Abnormal  
Syncope  
Tricuspid Valve

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Incompetence

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
PO		Health Professional	Librium (Chlorodiazepoxide Hcl)	PS		ORAL
100 MG SEE			Wellbutrin Tablet-Controlled Release	SS		ORAL
TEXT PO			Mirtazapin Tablet (Mirtazepine)	SS		ORAL
20 MG SEE						
TEXT PO						

Date:08/17/01ISR Number: 3778759-6Report Type:Expedited (15-DaCompany Report #A0156108A  
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG per day		Condition Aggravated		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Endometriosis		Herbs	C		
		Haemorrhage		Homeopathy	C		
		Haemorrhagic Disorder					
		Increased Tendency To Bruise					
		Menstrual Disorder					
		Stress					

Date:08/17/01ISR Number: 3778767-5Report Type:Expedited (15-DaCompany Report #B0115303A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN		Confusional State		Zyban	PS	Glaxo Wellcome	
		Convulsion					



Fatigue

Date:08/17/01ISR Number: 3778773-0Report Type:Expedited (15-DaCompany Report #B0117311A  
 Age:55 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day	Herpes Zoster		Zyban	PS	Glaxo Wellcome	ORAL
Other	100MCG Per day			Thyroxine	C	Glaxo Wellcome	ORAL

Date:08/17/01ISR Number: 3778774-2Report Type:Expedited (15-DaCompany Report #B0117315A  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Chronic Lymphocytic Leukaemia Skin Lesion		Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/17/01ISR Number: 3779653-7Report Type:Expedited (15-DaCompany Report #A0156711A  
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abortion Spontaneous	Study	Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Complications Of Maternal	Health				
150 MG/PER		Exposure To Therapeutic	Professional	Fluoxetine			
DAY/ORAL		Drugs		Hydrochloride	C		
		Pregnancy					

Date:08/17/01ISR Number: 3779654-9Report Type:Expedited (15-DaCompany Report #A0156593A  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Health	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
Other		Convulsive Threshold	Professional	Narcotic (Narcotic)	SS		
ORAL		Lowered	Company				
UNKNOWN			Representative				

Date:08/17/01ISR Number: 3779656-2Report Type:Expedited (15-DaCompany Report #A0157251A  
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL
Other		Joint Stiffness					
150 MG /		Multiple Sclerosis					
TWICE PER DAY		Neck Pain		Conjugated Estrogens	C		
/ ORAL		Pyrexia		Vitamin E	C		
		Tinnitus		Ascorbic Acid	C		
				Retinol	C		
				Multivitamin	C		

Date:08/17/01ISR Number: 3779710-5Report Type:Expedited (15-DaCompany Report #A0157266A  
Age:67 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Agitation	Consumer	Bupropion			
Initial or Prolonged	Atrial Fibrillation		Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
300 MG PER	Nervousness					
DAY/ORAL	Palpitations					

Date:08/17/01ISR Number: 3779818-4Report Type:Expedited (15-DaCompany Report #A114068  
Age:65 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Amnesia	Consumer	Viagra	PS	Pfizer Agricultural	
Hospitalization -	Cardio-Respiratory Arrest	Health			Div	ORAL
50.00 MG	Coma	Professional				
Initial or Prolonged	Convulsion					
TOTAL:PRN:ORA	Drug Interaction		Bupropion	SS		ORAL
Required	Fall					
L	Head Injury					
Intervention to	Tachycardia		Tamsulosin	SS		ORAL
300.00 MG						
Prevent Permanent						
TOTAL:PID:ORA						
Impairment/Damage						
L						
40 MG						
TOTAL:DAILY:O						
RAL			Atorvastatin	C		
			Lorazepam	C		
			Aspirin	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dimethyl Sulfone C  
 Ubidecarenone C  
 Multivitamin C  
 Fish Oil C  
 Pentatonic Acid C

Date:08/17/01ISR Number: 3780297-1Report Type:Expedited (15-DaCompany Report #B0113169A  
 Age:65 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aortic Valve Stenosis	Foreign	Bupropion			
Hospitalization -		Cardiac Failure	Health	Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG TWICE							
Initial or Prolonged		Coronary Artery Disease	Professional				
PER DAY ORAL							
		Decreased Appetite					
		Dry Throat					
		Retching					

Date:08/17/01ISR Number: 3785650-8Report Type:Expedited (15-DaCompany Report #B0116926A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Constipation	Foreign	Zyban Tablet - Zyban			
		Neoplasm Malignant	Consumer	(Bupropion			
				Hydrochloride)	PS		
150 MG							

Date:08/20/01ISR Number: 3779611-2Report Type:Expedited (15-DaCompany Report #A0155325A  
 Age:26 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Angioneurotic Oedema		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
150MG Twice							
Initial or Prolonged		Arthralgia					
per day	18 DAY						
		Dermatitis					
		Eyelid Oedema					

Face Oedema  
Herpes Zoster  
Joint Swelling  
Oedema Peripheral  
Serum Sickness  
Urticaria

Date:08/20/01ISR Number: 3779613-6Report Type:Expedited (15-DaCompany Report #A0155843A  
Age:26 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice Initial or Prolonged per day 19 DAY	Arthralgia Dermatitis Joint Swelling Serum Sickness Urticaria		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:08/20/01ISR Number: 3779780-4Report Type:Direct Company Report #  
Age:38 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Heart Rate Increased Hyperhidrosis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Palpitations Panic Attack Tremor	Report Source	Product	Role	Manufacturer	Route
1 DAILY ORAL				Zyban	PS		ORAL
				Coumadin Factor V	C C		

Date:08/20/01ISR Number: 3780396-4Report Type:Expedited (15-DaCompany Report #B0114323A  
Age: Gender:Female I/FU:I

Outcome Dose Other	Duration	PT Balance Disorder	Report Source Foreign	Product Wellbutrin	Role PS	Manufacturer	Route ORAL
100 MG / THREE TIMES PER DAY / ORAL	2 YR	Fall Tooth Loss	Consumer				

Date:08/20/01ISR Number: 3780397-6Report Type:Expedited (15-DaCompany Report #B0116765A  
Age:42 YR Gender:Male I/FU:I

Outcome Dose Hospitalization - Initial or Prolonged	Duration	PT Mania	Report Source Foreign	Product Zyban Tablet - Zyban (Bupropion Hydrochloride)	Role PS	Manufacturer	Route ORAL
150 MG / PER DAY/ ORAL				Amoxicillin Ethanol	C C		

Date:08/20/01ISR Number: 3780402-7Report Type:Expedited (15-DaCompany Report #B0109588A  
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged	150 MG / Required	Arrhythmia Asthenia Blood Pressure Decreased	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
Intervention to / ORAL		Dizziness					
Prevent Permanent Impairment/Damage		Dyspnoea					
		Feeling Abnormal		Aspirin	C		
		Pallor		Ketoprofen	C		
		Palpitations		Ethanol	C		
		Supraventricular Tachycardia					
		Syncope					

Date:08/20/01ISR Number: 3780521-5Report Type:Expedited (15-DaCompany Report #A0156690A  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged	150 MG / TWICE PER DAY	Hypersensitivity Viral Infection	Foreign Health Professional	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
Intervention to / ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/20/01ISR Number: 3780808-6Report Type:Expedited (15-DaCompany Report #B0116763A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Foreign	Bupropion Hydrochloride	PS	Glaxo Wellcome Inc	ORAL
150 MG/TTWICE PER DAY, ORAL							

Date:08/20/01ISR Number: 3780809-8Report Type:Expedited (15-DaCompany Report #A0157043A  
Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 50 MG/PER Initial or Prolonged DAY/ORAL		Atrial Fibrillation	Foreign	Wellbutrin Sr	PS	Glaxo Wellcome Inc	ORAL
			Consumer	Tamsulosin Hcl	C		

Date:08/21/01ISR Number: 3780539-2Report Type:Expedited (15-DaCompany Report #A0157983A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - Initial or Prolonged		Anaphylactic Reaction		Zyban	PS	Glaxo Wellcome	ORAL

Date:08/21/01ISR Number: 3780540-9Report Type:Expedited (15-DaCompany Report #B0107247A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypoaesthesia Paraesthesia		Zyban Seretide Ceclor	PS C C	Glaxo Wellcome Glaxo Wellcome	ORAL
375MG Twice							



per day

Solone

C

Date:08/21/01ISR Number: 3780541-0Report Type:Expedited (15-DaCompany Report #B0113284A  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Coronary Artery Disease Sudden Death	Zyban	PS	Glaxo Wellcome	

Date:08/21/01ISR Number: 3780545-8Report Type:Expedited (15-DaCompany Report #B0116842A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Myocardial Infarction	Zyban	PS	Glaxo Wellcome	

Date:08/21/01ISR Number: 3780547-1Report Type:Expedited (15-DaCompany Report #B0117368A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG As			Death	Zyban	PS	Glaxo Wellcome	

directed

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/21/01ISR Number: 3780548-3Report Type:Expedited (15-DaCompany Report #B0117371A

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice Initial or Prolonged per day	Angina Pectoris Chest Pain		Zyban	PS	Glaxo Wellcome	ORAL

Date:08/21/01ISR Number: 3780549-5Report Type:Expedited (15-DaCompany Report #B0117516A

Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Thermal Burn		Zyban	PS	Glaxo Wellcome	

Date:08/21/01ISR Number: 3780550-1Report Type:Expedited (15-DaCompany Report #D0019085A

Age:63 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150MG Per day 7 DAY 5MG Per day 20MG Per day	Chest Pain Constipation Dysuria Erythema Feeling Hot Genital Pruritus Female Gingivitis Headache Hyperaemia Mastitis Pruritus		Zyban Carbimazol Bisomerck Omeprazol Other Herbal Medications	PS C C C C	Glaxo Wellcome	ORAL ORAL ORAL ORAL ORAL

Date:08/22/01ISR Number: 3781489-8Report Type:Expedited (15-DaCompany Report #A0155912A

Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Unknown	Completed Suicide		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Head Injury		Celexa	SS		ORAL
		Overdose		Alcohol	SS		ORAL
		Toxicologic Test Abnormal		Benadryl	SS		ORAL

Date:08/22/01ISR Number: 3781490-4Report Type:Expedited (15-DaCompany Report #A0157654A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day 4 MON	Blood Potassium Decreased		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Blood Sodium Decreased		Zyprexa	C		
		Convulsion		Prozac	C		

Date:08/22/01ISR Number: 3781500-4Report Type:Expedited (15-DaCompany Report #B0117515A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day	Blood Pressure Decreased		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Chest Discomfort		Nicotine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/22/01ISR Number: 3781618-6Report Type:Direct  
 Age:33 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN		Chest Pain		Bupropion	PS		
Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage		Dyspnoea Electrocardiogram Abnormal Electrocardiogram St Segment Abnormal Stevens-Johnson Syndrome		Excedrin	C		

Date:08/22/01ISR Number: 3782160-9Report Type:Expedited (15-DaCompany Report #WAES 01072442  
 Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anxiety	Health	Tab Vioxx 25 Mg	PS		ORAL
25 MG/DAILY/ORAL		Aphasia	Professional				
25 MG/BID		Asthenia	Company	Anafranil	SS		
0.5 MG/BID		Balance Disorder	Representative	Klonopin	SS		
100 MG/BID		Blood Pressure Increased		Lamictal	SS		
150 MG/BID		Confusional State		Wellbutrin	SS		
250 MG		Coordination Abnormal		Dipentum	SS		
1.25 MG/DAILY		Difficulty In Walking		Premarin	SS		
100 MG/DAILY		Disorientation		Zoloft	SS		
		Dizziness		Coumadin	C		
		Drug Interaction		Motrin	C		
		Dry Skin		Lithium Carbonate	C		
		Dysarthria					
		Dysphagia					
		Dyspnoea					
		Emotional Distress					
		Feeling Abnormal					

Gait Disturbance  
 Headache  
 Mental Disorder  
 Mental Impairment  
 Motor Dysfunction  
 Respiratory Rate  
 Increased  
 Restlessness  
 Skin Warm  
 Speech Disorder  
 Transient Ischaemic  
 Attack  
 Vision Blurred

Date:08/23/01ISR Number: 3781909-9Report Type:Expedited (15-DaCompany Report #A0145968A  
 Age:54 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Per day 3 DAY Initial or Prolonged	Duration Hypersensitivity		Wellbutrin	PS	Glaxo Wellcome	ORAL
			Unknown			
			Anti-Depressant	C		
			Oxycontin	C		
			Percocet	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/23/01ISR Number: 3781918-XReport Type:Expedited (15-DaCompany Report #B0100202A

Age:59 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice	Hypertensive Crisis		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged per day 8 DAY	Transient Ischaemic Attack		Grapefruit Juice	C		
UNKNOWN			Aspirin	C		
150MG In the morning			Hrt	C		

Date:08/23/01ISR Number: 3781922-1Report Type:Expedited (15-DaCompany Report #B0117830A

Age:41 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150MG Per day	Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL

Date:08/23/01ISR Number: 3782583-8Report Type:Expedited (15-DaCompany Report #A0156108A

Age:45 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Condition Aggravated Ecchymosis Haemorrhagic Disorder Menstrual Disorder	Consumer	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL	Stress		Herbal Medication Homeopathic Medication	C C		

Date:08/23/01ISR Number: 3782735-7Report Type:Expedited (15-DaCompany Report #B0117516A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Thermal Burn	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		
150 MG			Company Representative				

Date:08/23/01ISR Number: 3782737-0Report Type:Expedited (15-DaCompany Report #B0117368A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Foreign Health Professional	Zyban Tablet- Zyban (Bupropion Hydrochloride)	PS		
150 MG / AS							
DIRECTED							

Date:08/23/01ISR Number: 3782738-2Report Type:Expedited (15-DaCompany Report #B0117371A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Angina Pectoris Chest Pain	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /			Company Representative				
TWICE PER DAY							
/ ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/23/01ISR Number: 3782741-2Report Type:Expedited (15-DaCompany Report #B0116842A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		
150 MG							

Date:08/23/01ISR Number: 3782742-4Report Type:Expedited (15-DaCompany Report #A0157983A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anaphylactic Reaction	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
Hospitalization - Initial or Prolonged ORAL							

Date:08/23/01ISR Number: 3782756-4Report Type:Expedited (15-DaCompany Report #B0117315A

Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Chronic Lymphocytic Leukaemia Lymphocytosis	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / ORAL							
Skin Disorder							

Date:08/23/01ISR Number: 3782758-8Report Type:Expedited (15-DaCompany Report #B0115303A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Confusional State Convulsion Fatigue	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
150 MG							



Date:08/23/01ISR Number: 3782759-XReport Type:Expedited (15-DaCompany Report #B0117311A  
Age:55 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required Intervention to Prevent Permanent Impairment/Damage	150 MG / PER DAY / ORAL	Herpes Zoster	Foreign	Zyban Tablet - Zybam (Bupropion Hydrochloride)	PS		ORAL
				Thyroxine Sodium	C		

Date:08/23/01ISR Number: 3782767-9Report Type:Expedited (15-DaCompany Report #B0107247A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150 MG	Hypoaesthesia Paraesthesia	Foreign Health Professional	Zyban Tablet- Zyban (Bupropion Hydrochloride)	PS		
				Fluticasone+Salmeter ol	C		
				Cefaclor	C		
				Solone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/23/01ISR Number: 3782770-9Report Type:Expedited (15-DaCompany Report #A0155325A

Age:26 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Angioneurotic Oedema Arthralgia Dermatitis Difficulty In Walking	Foreign Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL	Hepatic Function Abnormal  Herpes Zoster Hypogammaglobulinaemia Joint Swelling Red Blood Cell Sedimentation Rate Increased Serum Sickness Urticaria					

Date:08/23/01ISR Number: 3782790-4Report Type:Expedited (15-DaCompany Report #D0019085A

Age:63 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to Prevent Permanent 150 MG/ PER Impairment/Damage DAY/ ORAL	Chest Pain Constipation Dysuria Erythema  Feeling Hot Gingivitis Headache Hyperaemia Mastitis Pruritus	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)  Carbimazole Bisoprolol Omeprazole Herbal Medication	PS  C C C C		ORAL

Date:08/23/01ISR Number: 3783287-8Report Type:Expedited (15-DaCompany Report #B0113284A

Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coronary Artery Disease Overdose Sudden Death	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		

Date:08/23/01ISR Number: 3783293-3Report Type:Expedited (15-DaCompany Report #A0155843A  
Age:26 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Serum Sickness	Foreign Health Professional	Wellbutrin Tablet-Controlled Release (Buproprion Hydrochloride)	PS		ORAL

150 MG /TWICE  
PER DAY/ORAL

Date:08/24/01ISR Number: 3782396-7Report Type:Expedited (15-DaCompany Report #A0158445A  
Age:42 YR Gender:Female I/FU:F

Outcome	PT
Other	Attention Deficit/Hyperactivity Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1	MON	Bronchitis Upper Respiratory Tract Infection					
		Urticaria		Wellbutrin	PS	Glaxo Wellcome	ORAL
				Ceftin	SS	Glaxo Wellcome	ORAL
				Penicillin	SS	Glaxo Wellcome	ORAL

Date:08/24/01ISR Number: 3782945-9Report Type:Expedited (15-DaCompany Report #B0117515A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Pressure Decreased Chest Discomfort	Foreign Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/PER							
DAY/ORAL				Nicotine	C		

Date:08/24/01ISR Number: 3783160-5Report Type:Expedited (15-DaCompany Report #A0157654A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Potassium Decreased Blood Sodium Decreased Convulsion	Consumer	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER							
DAY / ORAL				Olanzapine	C		
				Fluoxetine			
				Hydrochloride	C		

Date:08/24/01ISR Number: 3783166-6Report Type:Expedited (15-DaCompany Report #A0155912A  
Age:37 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide Injury Intentional Misuse	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
	150 MG/	ORAL			Citalopram Hydrobromide Tablet (Citalopram Hydrobromide)	SS		ORAL
	ORAL				Ethanol Liquid (Alcohol)	SS		ORAL
	ORAL				Diphenhydramine Hcl (Formulation Unknown) (Diphenhydramine Hcl)	SS		ORAL

Date:08/27/01ISR Number: 3782699-6Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #

Outcome PT  
Other Arthralgia  
Joint Stiffness  
Myalgia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pyrexia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MGM			Zyban 150 Mgm/Day X 3 D (Began 17 July 01) Than Bid	PS		
WEEKLY X3 D - THEN BID						
1 DAILY			Nicoderm Patch	SS		

Date:08/27/01ISR Number: 3782987-3Report Type:Expedited (15-DaCompany Report #A0155443A  
Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day		Condition Aggravated		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Convulsion		Vioxx	SS		
		Fall		Celebrex	SS		
		Head Injury		Neurontin	C		
		Headache					
		Inflammation					
		Migraine					
		Mucous Membrane Disorder					
		Pruritus					
		Sinus Headache					
		Swelling					

Date:08/27/01ISR Number: 3783020-XReport Type:Expedited (15-DaCompany Report #B0116466A  
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG As directed	8 DAY	Dizziness		Zyban	PS	Glaxo Wellcome	
		Feeling Abnormal					

3MG Three	Headache	Bromazepam	C		ORAL
times per day	Insomnia				
100MG Twice	Syncope	Gtn	C	Glaxo Wellcome	ORAL
per day	Tremor				
100MG Per day		Atenolol	C		ORAL
625MCG Three		Conjugated Oestrogens	C		ORAL
times per					
week					

Date:08/27/01ISR Number: 3783022-3Report Type:Expedited (15-DaCompany Report #B0118184A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Gastritis		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	17 DAY						
28 DAY				Aspirin	C		ORAL
2.5MG Per day				Ramipril	C		ORAL
2MG Per day				Oestradiol	C		ORAL
UNKNOWN				Levonorgestrel	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/27/01ISR Number: 3783314-8Report Type:Expedited (15-DaCompany Report #GBR004231

Age:52 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Lethargy	Health	Meridia	PS		
Initial or Prolonged	Overdose	Professional	Wellbutrin	SS		
	Vomiting	Other	Prozac	SS		
			Serzone	SS		
			Lorazepam	SS		
			Duract	SS		

Date:08/27/01ISR Number: 3783435-XReport Type:Expedited (15-DaCompany Report #A0145968A

Age:54 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Asthenia	Health	Wellbutrin Tablet -			
Initial or Prolonged	Decreased Appetite	Professional	Controlled Release			
	Dysphagia		(Bupropion			
	Hypersensitivity		Hydrochloride)	PS		ORAL
150 MG, PER						
DAY, ORAL						
			Antidepressant	C		
			Oxycodone			
			Hydrochloride	C		
			Percocet	C		

Date:08/27/01ISR Number: 3783445-2Report Type:Expedited (15-DaCompany Report #A0158445A

Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Attention	Health	Wellbutrin Tablet			
	Deficit/Hyperactivity	Professional	(Bupropion			
	Disorder	Company	Hydrochloride)	PS		ORAL
ORAL						
	Bronchitis	Representative	Ceftin Tablet			
	Upper Respiratory Tract		(Cefuroxime Axetil)	SS		ORAL
ORAL						
	Infection		Phenoxyethylpenicil			
	Urticaria		lin K (Formulation			



Date:08/27/01ISR Number: 3783523-8Report Type:Expedited (15-DaCompany Report #A103444  
Age:79 YR Gender:Female I/FU:F

Outcome	PT
Required	Aggression
Intervention to	Blood Pressure Increased
Prevent Permanent	Blood Thyroid Stimulating
Impairment/Damage	Hormone Decreased
	Drug Ineffective
	Ecchymosis
	Epistaxis
	Fall
	Haemorrhagic Stroke
	Head Injury
	Headache
	Heart Rate Decreased
	Heart Rate Increased
	Hostility
	Hyperhidrosis
	Overdose

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Palpitations Parkinson'S Disease Syncope					
50.00 MG		Thinking Abnormal	Consumer	Zoloft Tablets	PS		
TOTAL: DAILY		Transient Ischaemic					
300.00 MG		Attack		Wellbutrin	SS		ORAL
TOTAL: BID:							
ORAL							
				Coumadin	SS		
300.00 MG				Effexor Sr	SS		ORAL
TOTAL: BID:							
ORAL							
				Toprol	C		
				Zebeta	C		
				Lanoxin	C		
				Lasix	C		
				Synthroid	C		
				Vitamin E	C		
				Estrogen	C		

Date:08/27/01ISR Number: 3783809-7Report Type:Expedited (15-DaCompany Report #B0117830A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent 150 MG / PER Impairment/Damage DAY / ORAL		Chest Pain Hypoaesthesia	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS	Glaxosmithkline	ORAL

Date:08/27/01ISR Number: 3783811-5Report Type:Expedited (15-DaCompany Report #B0100202A  
Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hypertensive Crisis Transient Ischaemic Attack	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS	Glaxosmithkline	
150 MG /							
TWICE PER DAY							

Grapefruit Juice	C
Aspirin	C
Hrt	C

Date:08/28/01ISR Number: 3783328-8Report Type:Direct Company Report #  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Alopecia		Wellbutrin Sr/ 150 Mg	PS		ORAL
1 PER/ ORAL							

Date:08/28/01ISR Number: 3783481-6Report Type:Expedited (15-DaCompany Report #B0117307A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Feeling Abnormal		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day	14 DAY	Mental Impairment Pupillary Disorder		Femodene	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/28/01ISR Number: 3783484-1Report Type:Expedited (15-DaCompany Report #B0118326A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cold Sweat		Zyban	PS	Glaxo Wellcome	ORAL
6 DAY		Pain In Extremity Pallor Palpitations					

Date:08/28/01ISR Number: 3783904-2Report Type:Direct Company Report #  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Urticaria		Zyban	PS		
150 MG SR BID							

Date:08/29/01ISR Number: 3784006-1Report Type:Expedited (15-DaCompany Report #B0117827A  
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Back Pain		Zyban	PS	Glaxo Wellcome	
UNKNOWN	33 DAY	Depression Difficulty In Walking Dizziness Dystonia Headache Malaise Pain In Extremity					

Date:08/29/01ISR Number: 3785299-7Report Type:Expedited (15-DaCompany Report #A0155443A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Convulsion Fall	Health Professional	Wellbutrin Tablet-Controlled			

150 MG/ PER	Head Injury	Release (Bupropion			
	Headache	Hydrochloride)	PS	Glaxosmithkline	ORAL
DAY / ORAL	Increased Viscosity Of				
	Bronchial Secretion	Rofecoxib			
	Migraine	(Formulation			
	Pruritus	Unknown) (Rofecoxib)	SS		
	Sinus Headache	Celecoxib			
	Swelling	(Formulation			
		Unknown) (Celecoxib)	C		
		Gabapentin	C		

Date:08/29/01ISR Number: 3785604-1Report Type:Expedited (15-DaCompany Report #B0118184A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Dyspepsia	Foreign	Zyban Tablet - Zyban			
Intervention to		Gastritis		(Bupropion			
Prevent Permanent				Hydrochloride)	PS		ORAL
150 MG /							
Impairment/Damage							
TWICE PER DAY							
/ ORAL							
				Aspirin	C		
				Ramipril	C		
				Oestradiol	C		
				Levonorgestrel	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/29/01ISR Number: 3785620-XReport Type:Expedited (15-DaCompany Report #B0116466A

Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Foreign	Zyban Tablet - Zyban			
		Headache	Consumer	(Bupropion			
150 MG/AS		Insomnia		Hydrochloride)	PS		
		Petit Mal Epilepsy					
DIRECTED		Syncope		Bromazepam	C		
		Tremor		Nitroglycerin	C		
				Atenolol	C		
				Conjugated Estrogens	C		

Date:08/30/01ISR Number: 3785121-9Report Type:Expedited (15-DaCompany Report #A0155912A

Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice		Head Injury	Professional				
per day		Overdose		Celexa	SS		ORAL
		Toxicologic Test Abnormal		Alcohol	SS		ORAL
				Benadryl	SS		ORAL

Date:08/30/01ISR Number: 3785124-4Report Type:Expedited (15-DaCompany Report #A0159165A

Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
200MG Twice		Bone Graft					
per day	6	MON		Toprol Xl	SS		ORAL
50MG Per day		Drug Interaction					
		Liver Function Test		Nitrodur	C	Glaxo Wellcome	
		Abnormal		Ranitidine	C	Glaxo Wellcome	
		Scleral Discolouration		Hydroxyzine	C		

Tinnitus  
Tooth Disorder

Date:08/30/01ISR Number: 3785131-1Report Type:Expedited (15-DaCompany Report #B0118186A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	9 DAY	Hemiparesis Hypoaesthesia Hyporeflexia Nervousness Sensory Disturbance					

Date:08/30/01ISR Number: 3785132-3Report Type:Expedited (15-DaCompany Report #B0118455A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Thrombosis		Zyban	PS	Glaxo Wellcome	
150MG As							

directed

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/01ISR Number: 3785916-1Report Type:Direct  
 Age:64 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG SR Q AM		Agitation Coordination Abnormal		Wellbutrin / Bupropion Sr 100mg	PS		
		Hyperhidrosis		Clonazepam	C		
		Mydriasis		Zoloft	C		
		Tachycardia		Mertazapine	C		
		Tremor					

Date:08/31/01ISR Number: 3785967-7Report Type:Expedited (15-DaCompany Report #A0141089A  
 Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day		Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Fall		Alcohol	SS		
		Intentional Misuse		Flovent	C	Glaxo Wellcome	
		Multiple Fractures		Serevent	C	Glaxo Wellcome	
		Stupor		Atrovent	C	Glaxo Wellcome	
		Suicide Attempt		Proventil	C	Glaxo Wellcome	

Date:08/31/01ISR Number: 3785982-3Report Type:Expedited (15-DaCompany Report #A0159004A  
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Coronary Artery Surgery		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day		Heart Rate Decreased					
81MG Per day		Syncope		Aspirin	C		ORAL
				Glucophage	C		
				Nitroglycerin	C	Glaxo Wellcome	
				Altace	C		
				Zocor	C		
				Niaspan	C		



Date:08/31/01ISR Number: 3785985-9Report Type:Expedited (15-DaCompany Report #B0089512A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG Twice		Erythema Multiforme Psoriasis		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
per day	21 DAY	Rash Pustular Skin Disorder		Co-Proxamol Dothiepin	C C		ORAL ORAL

Date:08/31/01ISR Number: 3785986-0Report Type:Expedited (15-DaCompany Report #B0089979A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2 WK		Erythema Multiforme		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Psoriasis Skin Disorder		Coproxamol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/01ISR Number: 3785994-XReport Type:Expedited (15-DaCompany Report #B0118584A

Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Single	Myocardial Infarction		Zyban	PS	Glaxo Wellcome	
dose							

Date:08/31/01ISR Number: 3787093-XReport Type:Periodic Company Report #A115200

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	25.00 MG	Dermatitis	Health	Ziprasidone Po	PS		
TOTAL:DAILY		Glossitis	Professional				
			Company Representative	Wellbutrin (Ibuproprion)	SS		

Date:08/31/01ISR Number: 3787140-5Report Type:Periodic Company Report #A117193

Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	160.00 MG	Akathisia	Health	Ziprasidone Po	PS		ORAL
TOTAL:PID:ORA		Mania	Professional				

L

Wellbutrin	SS
Klonopin	C
Luvox	C

Date:08/31/01ISR Number: 3787319-2Report Type:Expedited (15-DaCompany Report #B0117827A

Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anhedonia Depression Difficulty In Walking Dizziness Dystonia Malaise Pain	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		

Date:08/31/01ISR Number: 3787767-0Report Type:Periodic Company Report #A118566  
 Age:27 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - NOT SPECIFIED		Convulsion	Health	Ziprasidone Po	PS		ORAL
Initial or Prolonged Required		Electrocardiogram Qt Prolonged	Professional Company	Wellbutrin Prozac	SS SS		
Intervention to Prevent Permanent Impairment/Damage		Intentional Misuse Tachycardia	Representative	Depakote	SS		

Date:08/31/01ISR Number: 3788846-4Report Type:Periodic Company Report #A108581  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Health	Ziprasidone Po	PS		ORAL
150.00 MG		Drug Interaction	Professional				

TOTAL: BID: ORA  
 L  
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

250.00 MG Wellbutrin SS ORAL

TOTAL: BID: ORA

L

Date: 09/04/01 ISR Number: 3786883-7 Report Type: Expedited (15-DaCompany Report #B0115303A  
 Age: 27 YR Gender: Female I/FU: F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Anxiety		Zyban	PS	Glaxo Wellcome	
UNKNOWN	20 DAY						
Hospitalization -		Bite		Dianette	C		ORAL
1TAB Per day							
Initial or Prolonged		Confusional State					
Other		Grand Mal Convulsion					
		Loss Of Consciousness					
		Tongue Disorder					

Date: 09/04/01 ISR Number: 3786885-0 Report Type: Expedited (15-DaCompany Report #B0116842A  
 Age: Gender: Male I/FU: F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Hospitalization -							
per day							
Initial or Prolonged							

Date: 09/04/01 ISR Number: 3788024-9 Report Type: Direct Company Report #  
 Age: 42 YR Gender: Female I/FU: I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Aggression		Zoloft 100 Mg Pfizer	PS	Pfizer	ORAL
3 TABS/ EVERY							
Other		Agitation					
MORNIN/ORAL							

Anger  
Bipolar I Disorder

Wellbutrin Sr 100 Mg  
Glaxo

SS Glaxo

ORAL

2 TABS / 2

Crying

TIMES/ ORAL

Dizziness  
Euphoric Mood  
Fear  
Feeling Abnormal  
Irritability  
Nausea  
Premenstrual Syndrome  
Suicidal Ideation  
Tinnitus

Date:09/05/01ISR Number: 3787337-4Report Type:Expedited (15-DaCompany Report #A0158740A  
Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Parkinson'S Disease		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:09/05/01ISR Number: 3787350-7Report Type:Expedited (15-DaCompany Report #B0118929A  
Age:28 YR Gender:Male I/FU:I

Outcome  
Hospitalization -  
Initial or Prolonged

PT  
Blood Pressure Abnormal  
Dysphagia  
Erythema  
Eyelid Oedema

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Face Oedema Heart Rate Increased Jaundice					
		Skin Discolouration		Zyban	PS	Glaxo Wellcome	ORAL

Date:09/05/01ISR Number: 3788288-1Report Type:Expedited (15-DaCompany Report #A0141089A  
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthralgia Blood Pressure Increased Condition Aggravated	Health Professional	Zyban Tablets - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ PER DAY/ ORAL		Fracture					
		Liver Function Test Abnormal Renal Impairment Suicide Attempt		Ethanol (Formulation Unknown) (Alcohol) Fluticasone Propionate Salmeterol Xinafoate Ipratropium Bromide Salbutamol Sulphate	SS C C C C		

Date:09/05/01ISR Number: 3788369-2Report Type:Expedited (15-DaCompany Report #B0118186A  
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aggression Fluid Overload Hemiparesis	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER DAY/ ORAL		Hypoaesthesia  Hyporeflexia Nervousness					

Date:09/05/01ISR Number: 3788370-9Report Type:Expedited (15-DaCompany Report #B0118455A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Thrombosis	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
Intervention to Prevent Permanent 150 MG / AS Impairment/Damage DIRECTED /							

Date:09/05/01ISR Number: 3788381-3Report Type:Expedited (15-DaCompany Report #A0155912A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Injury Intentional Misuse	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE PER DAY/ORAL							
				Citalopram Hydrobromide Tablet (Citalopram Hydrobromide)	SS		ORAL
				Ethanol Liquid (Alcohol)	SS		ORAL
				Diphenhydramine Hcl			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL (Formulation Unknown) (Diphenhydramine Hcl) SS ORAL

Date:09/05/01ISR Number: 3788382-5Report Type:Expedited (15-DaCompany Report #A0159165A  
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Amnesia Drug Interaction Loss Of Consciousness Scleral Discolouration	Consumer	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
200 MG TWICE PER DAY ORAL		Tinnitus					
50 MG PER DAY ORAL		Tooth Disorder		Metoprolol Succinate (Formulation Unknown) (Metoprolol Succinate)	SS		ORAL
				Nitroglycerin	C		
				Ranitidine Hydrochloride	C		
				Hydroxyzine	C		

Date:09/05/01ISR Number: 3788391-6Report Type:Expedited (15-DaCompany Report #A0159004A  
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Heart Rate Decreased Quadruple Vessel Bypass Graft	Consumer	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG TWICE PER DAY ORAL				Aspirin Metformin	C		



Hydrochloride	C
Nitroglycerin	C
Ramipril	C
Simvastatin	C
Nicotinic Acid	C

Date:09/05/01ISR Number: 3788401-6Report Type:Expedited (15-DaCompany Report #B0089979A  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	2 WK	Erythema Multiforme Psoriasis	Foreign Health Professional Company Representative	Zyban Tablet - Zyban (Bupropion Hydrochloride) Co-Proxamol	PS C		ORAL

Date:09/05/01ISR Number: 3788412-0Report Type:Expedited (15-DaCompany Report #B0089512A  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Erythema Multiforme Psoriasis	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

150 MG /

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Freedom Of Information (FOI) Report

TWICE PER DAY

/ ORAL

Co-Proxamol C  
 Dothiepin C

Date:09/05/01ISR Number: 3788677-5Report Type:Expedited (15-DaCompany Report #B0118584A  
 Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
150 MG/							

SINGLE DOSE

Date:09/06/01ISR Number: 3787894-8Report Type:Expedited (15-DaCompany Report #A0158740A  
 Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Parkinson'S Disease		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:09/06/01ISR Number: 3787907-3Report Type:Expedited (15-DaCompany Report #B0118929A  
 Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Pressure Abnormal Dysphagia Erythema Eyelid Oedema Face Oedema Heart Rate Increased Jaundice Skin Discolouration		Zyban	PS	Glaxo Wellcome	ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	6 WK	Asthenia Back Pain Difficulty In Walking Fatigue Feeling Jittery Muscle Spasms Myalgia Pain		Zyban Vitamins	PS C	Glaxo Wellcome	ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death		Arthralgia		Zyban	PS	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/06/01ISR Number: 3788948-2Report Type:Expedited (15-DaCompany Report #2010427

Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Diverticulum Drug Level Above Therapeutic Overdose	Study Health Professional Other	Oxycontin Cr Tablets, 40 Mg (Oxycodone Hydrochloride)	PS		ORAL
40 MG TID PO							
		Pulmonary Congestion Pulmonary Oedema		Oxycontin Cr Tablets, 20 Mg (Oxycodone Hydrochloride)	SS		ORAL
20 MG TID PO							
				Bupropion Hydrochloride	SS		
				Promethazine Hydrochloride	SS		
				Diazepam	SS		
				Acetaminophen	SS		
				Tetrahydrocannabinol Premarin (Conjugated Estrogens)	SS C		
				Wellbutrin (Bupropion)	C		
				Lortab (Hydrocodone/Acetami nophen)	C		
				Diazepam	C		

Date:09/07/01ISR Number: 3789066-XReport Type:Expedited (15-DaCompany Report #A0159874A

Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	26 DAY	Difficulty In Walking					
		Drug Hypersensitivity Dry Mouth Insomnia Muscle Spasms Oedema Peripheral					

Pyrexia  
Urticaria

Date:09/07/01ISR Number: 3789080-4Report Type:Expedited (15-DaCompany Report #B0118244A  
Age:63 YR Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG Twice per day	11 DAY	Abdominal Tenderness Chest Pain Circulatory Collapse Dyspepsia Dyspnoea Heart Rate Increased Pain Pharyngolaryngeal Pain Vertigo		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/07/01ISR Number: 3789085-3Report Type:Expedited (15-DaCompany Report #B0119109A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Zyban	PS	Glaxo Wellcome	

Date:09/07/01ISR Number: 3789087-7Report Type:Expedited (15-DaCompany Report #B0119241A

Age:57 YR Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebrovascular Accident		Zyban	PS	Glaxo Wellcome	

Date:09/07/01ISR Number: 3789088-9Report Type:Expedited (15-DaCompany Report #B0119242A

Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Haemorrhagic Stroke		Zyban	PS	Glaxo Wellcome	

Date:09/07/01ISR Number: 3789584-4Report Type:Expedited (15-DaCompany Report #HQ3100411JUL2001

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Complications Of Maternal	Health	Effexor	PS		ORAL
Initial or Prolonged ORAL		Exposure To Therapeutic Drugs	Professional	Remeron (Mirtazapine)	SS		ORAL
ORAL		Convulsion Neonatal		Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL

Date:09/07/01ISR Number: 3789585-6Report Type:Expedited (15-DaCompany Report #B0115303A

Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged	150 MG	Confusional State Grand Mal Convulsion Loss Of Consciousness	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
Other Required Intervention to Prevent Permanent Impairment/Damage				Dianette	C		

Date:09/07/01ISR Number: 3789587-XReport Type:Expedited (15-DaCompany Report #B0116842A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - Initial or Prolonged	150 MG /	Cardiac Arrest	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

TWICE PER DAY

/ ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/10/01ISR Number: 3789554-6Report Type:Expedited (15-DaCompany Report #A0159171A  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	100MG Per day 2	MON	Asthenia	Wellbutrin	PS	Glaxo Wellcome	ORAL
			Chest Pain	Phenergan	SS		
INTRAVENOUS							
			Circulatory Collapse	Atenolol	C		
			Convulsion	Lipitor	C		
			Dizziness	Methotrexate	C		
			Drug Interaction	Folic Acid	C		
			Ecchymosis	Ambien	C		
			Face Oedema	Ativan	C		
			Fall	Zantac	C	Glaxo Wellcome	
			Headache				
			Muscle Spasms				
			Nausea				
			Paralysis				
			Sedation				
			Tremor				

Date:09/10/01ISR Number: 3789556-XReport Type:Expedited (15-DaCompany Report #B0105578A  
 Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice		Arthralgia	Zyban	PS	Glaxo Wellcome	ORAL
	per day		Cerebrovascular Accident				
	100UG Twice		Liver Function Test	Eltroxin	C	Glaxo Wellcome	ORAL
	per day		Abnormal				
	5MG Per day			Vasace	C		ORAL
				Mirtazapine	C		
				Pravastatin	C		
				Aspirin	C		
				Omeprazole	C		
				Atenolol	C		
				Isosorbide	C		
				Ramipril	C		



Gemfibrozil

C

Date:09/10/01ISR Number: 3790054-8Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG PO QD X 3 DAYS THEN 150MG BID		Convulsion		Wellbutrin Sr 150 Mg (Glaxowellcome)	PS	Glaxowellcome	ORAL
				Ultram	C		
				Paxil	C		
				Tylenol #4	C		

Date:09/11/01ISR Number: 3790241-9Report Type:Expedited (15-DaCompany Report #B0119467A  
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 8 DAY Other RESPIRATORY (INHALATION) RESPIRATORY (INHALATION)		Purpura		Zyban	PS	Glaxo Wellcome	ORAL
				Combivent	C		
				Salmeterol	C	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/11/01ISR Number: 3790242-0Report Type:Expedited (15-DaCompany Report #B0119683A

Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice per day	Alcohol Poisoning		Zyban	PS	Glaxo Wellcome	ORAL
	275MG At night	Drug Toxicity		Prothiaden	SS		ORAL
	5MG Twice per day			Alcohol Olanzapine	SS C		ORAL
	5MG At night			Lithium Nitrazepam	C C		ORAL ORAL
				Oxybutynin	C		ORAL

Date:09/11/01ISR Number: 3791424-4Report Type:Expedited (15-DaCompany Report #B0117307A

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage DAY/ORAL	300 MG/PER DAY/ORAL	Emotional Disorder Eye Disorder Mental Disorder	Foreign	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
				Femodene	C		

Date:09/11/01ISR Number: 3791444-XReport Type:Expedited (15-DaCompany Report #A0158740A

Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Parkinson'S Disease	Health Professional	Wellbutrin Unspecified Tablet			

ORAL (Bupropion Hydrochloride) PS ORAL

Date:09/11/01ISR Number: 3791505-5Report Type:Expedited (15-DaCompany Report #B0118326A  
Age:33 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required Intervention to Prevent Permanent 150 MG/ ORAL Impairment/Damage	Cold Sweat Pain In Extremity Pallor Palpitations	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:09/11/01ISR Number: 3791518-3Report Type:Expedited (15-DaCompany Report #B0118929A  
Age:28 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG/ ORAL	Blood Pressure Abnormal Dysphagia Eyelid Oedema Face Oedema Heart Rate Increased Jaundice Skin Discolouration	Foreign Consumer	Zyban Tablet- Zyban (Bupropion Hydrochloride)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/12/01ISR Number: 3792211-3Report Type:Expedited (15-DaCompany Report #B0105578A  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arthralgia Cerebrovascular Accident Liver Function Test	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY / ORAL		Abnormal		Thyroxine Sodium Cilazapril Mirtazapine Pravastatin Aspirin Omeprazole Atenolol Isosorbide Ramipril Gemfibrozil	C C C C C C C C C C		

Date:09/12/01ISR Number: 3792243-5Report Type:Expedited (15-DaCompany Report #A0159874A  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG / Initial or Prolonged TWICE PER DAY / ORAL		Arthropathy Difficulty In Walking Drug Hypersensitivity	Foreign Consumer	Zyban	PS		ORAL

Date:09/12/01ISR Number: 3792267-8Report Type:Expedited (15-DaCompany Report #B0118244A  
 Age:63 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Tenderness Angina Unstable	Foreign Study	Zyban Tablet - Zyban (Bupropion			

150 MG /	Chest Pain	Health	Hydrochloride)	PS	ORAL
TWICE PER DAY	Dyspepsia	Professional			
/ ORAL	Dyspnoea				
	Heart Rate Increased				
	Pharyngolaryngeal Pain				

Date:09/12/01ISR Number: 3792269-1Report Type:Expedited (15-DaCompany Report #B0119242A  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Haemorrhagic Stroke	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		

150 MG

Date:09/12/01ISR Number: 3792413-6Report Type:Expedited (15-DaCompany Report #A0159171A  
Age:48 YR Gender:Female I/FU:I

Outcome	PT
Other	Chest Pain
	Circulatory Collapse
	Convulsion
	Dizziness
	Drug Interaction
	Ecchymosis

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG/ PER DAY/ORAL		Face Oedema Fall Gait Disturbance	Consumer	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
		Head Injury Muscle Spasms Nausea Paralysis					
		Refusal Of Treatment By Patient Sedation		Promethazine Hcl Injection (Promethazine Hcl)	SS		
INTRAVENOUS DOSE/INTRAVENOUS	SINGLE			Atenolol Atorvasatin Calcium Methotrexate Folic Acid Zolpidem Tartrate Lorazepam Ranitidine Hydrochloride	C C C C C C C		

Date:09/12/01ISR Number: 3792885-7Report Type:Expedited (15-DaCompany Report #A0159343A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL	6 WK	Asthenia Difficulty In Walking Fatigue Feeling Jittery Muscle Spasms Pain	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)  Multivitamin	PS  C		ORAL

Date:09/12/01ISR Number: 3792926-7Report Type:Expedited (15-DaCompany Report #B0119241A  
Age:57 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebrovascular Accident	Foreign	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		
UNKNOWN							

Date:09/12/01ISR Number: 3792929-2Report Type:Expedited (15-DaCompany Report #B0119109A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Foreign	Zyban Tablet - Zyban(Bupropion Hydrochloride)	PS		
UNKNOWN	150 MG						

Date:09/13/01ISR Number: 3792103-XReport Type:Expedited (15-DaCompany Report #B0119682A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 10TABS Per day		Anger Depression Overdose		Zyban Paracetamol  Panadeine	PS SS  SS	Glaxo Wellcome	
10TABS Per							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

day

Mersyndol Forte SS

10TABS Per

day

Date:09/14/01ISR Number: 3792818-3Report Type:Expedited (15-DaCompany Report #A0155912A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice	Completed Suicide		Wellbutrin	PS	Glaxo Wellcome	ORAL
	per day	Head Injury					
		Overdose		Celexa	SS		ORAL
		Toxicologic Test Abnormal		Alcohol	SS		ORAL
				Benadryl	SS		ORAL

Date:09/14/01ISR Number: 3792819-5Report Type:Expedited (15-DaCompany Report #A0159903A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Transient Ischaemic Attack		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL

Date:09/14/01ISR Number: 3792829-8Report Type:Expedited (15-DaCompany Report #B0120029A  
Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Zyban	PS	Glaxo Wellcome	
		Overdose		Slow K	SS		
				Temazepam	C		
				Hydrochlorothiazide	C		
				Salbutamol	C	Glaxo Wellcome	
				Fluticasone	C	Glaxo Wellcome	



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Zyban	PS	Glaxo Wellcome	
8 DAY		Circulatory Collapse					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dermatitis		Zyban 150mg Catalytica	PS	Catalytica	ORAL
150MG BID							
ORAL				Nicoderm Patch	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Chest Pain Vomiting		Zyban 150mg Catalytica	PS	Catalytica	ORAL
150MG BID							
ORAL				Ortho-Novum 777	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nicoderm C

Date:09/17/01ISR Number: 3793868-3Report Type:Direct  
Age:31 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash Generalised		Zyban	PS	Catalytica	ORAL
150 MG/ BID/		Rash Pruritic					
ORAL							

Nicoderm C

Date:09/17/01ISR Number: 3793874-9Report Type:Direct  
Age:27 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis		Zyban	PS	Catalytica	ORAL
150 MG/ QD TO		Pruritus					
BID/ ORAL							

Nicoderm C

Verapamil C

Date:09/17/01ISR Number: 3794365-1Report Type:Expedited (15-DaCompany Report #WAES 01072442  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Health	Tab Vioxx 25 Mg	PS		ORAL
25		Aphasia	Professional				
MG/DAILY/ORAL		Asthenia	Company	Anafranil Unk	SS		
25 MG/BID		Balance Disorder	Representative	Klonopin Unk	SS		
0.5 MG/BID		Blood Pressure Increased		Lamictal Unk	SS		
100 MG/BID							

150 MG/BID	Confusional State	Wellbutrin Sr Unk	SS
	Coordination Abnormal	Dipentum Unk	SS
250 MG	Difficulty In Walking	Premarin Unk	SS
1.25 MG/DAILY	Disorientation	Zoloft Unk	SS
100 MG/DAILY	Dizziness	Coumadin	C
	Drug Interaction	Motrin	C
	Dry Skin	Lithium Carbonate	C
	Dysarthria		
	Dysphagia		
	Dyspnoea		
	Emotional Distress		
	Headache		
	Mental Disorder		
	Motor Dysfunction		
	Respiratory Rate Increased		
	Restlessness		
	Skin Warm		
	Speech Disorder		
	Transient Ischaemic Attack		
	Vision Blurred		

Date:09/17/01ISR Number: 3794470-XReport Type:Expedited (15-DaCompany Report #B0119682A

Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Anger
Initial or Prolonged	Depression

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Overdose

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG		Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
10 TABLETS			Paracetamol (Formulation Unknown) (Acetaminophen)	SS		
PER DAY						
10 TABLETS			Panadeine (Formulation Unknown) (Panadeine)	SS		
PER DAY						
10 TABLETS			Mersyndol (Formulation Unknown) (Mersyndol)	SS		
PER DAY						

Date:09/17/01ISR Number: 3794485-1Report Type:Expedited (15-DaCompany Report #B0119683A  
 Age:46 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG /	TWICE PER DAY	Alcohol Poisoning Drug Toxicity	Foreign Health Professional	Zyban Tablet -Zyban (Bupropion Hydrochloride)	PS		ORAL
		/ ORAL			Dothiepin Hydrochloride (Dothiepin)			

275 MG / AT

Hydrochloride)

SS

ORAL

NIGHT / ORAL

Ethanol (Alcohol)

SS

Olanzapine

C

Lithium Salt

C

Nitrazepam

C

Oxybutynin

C

Date:09/17/01ISR Number: 3794506-6Report Type:Expedited (15-DaCompany Report #B0119467A

Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required		Purpura	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage				Combivent Salmeterol Xinafoate	C		
					C		

Date:09/19/01ISR Number: 3795450-0Report Type:Direct

Company Report #

Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	2 AM 1 PM	Drug Effect Decreased		Wellbutrin Sr 100mg	PS		

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Freedom Of Information (FOI) Report

Date:09/20/01ISR Number: 3795895-9Report Type:Expedited (15-DaCompany Report #B0114598A

Age:56 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice Initial or Prolonged per day	Abdominal Pain Accident		Zyban	PS	Glaxo Wellcome	ORAL
Duration 62 DAY	Aggression Amnesia Angioneurotic Oedema Balance Disorder Blood Bilirubin Increased Blood Pressure Increased Chest Pain Circulatory Collapse Complement Factor C3 Increased Confusional State Convulsion Dermatitis Diarrhoea Disorientation Disturbance In Attention Dizziness Dry Skin Dysphagia Dyspnoea Eyelid Oedema Face Oedema Feeling Hot Flatulence Flushing Hyperhidrosis Illusion Immunology Test Abnormal Irritability Irritable Bowel Syndrome Loss Of Consciousness Malaise Nausea Ocular Hyperaemia Oedema Peripheral Pharyngeal Oedema Pruritus Rhinitis					

Speech Disorder  
Tinnitus  
Tongue Oedema  
Urine Abnormality  
Urticaria

Date:09/20/01ISR Number: 3795904-7Report Type:Expedited (15-DaCompany Report #B0119910A  
Age:42 YR Gender:Unknown I/FU:I

Outcome PT  
Hospitalization - Bacterial Infection  
Initial or Prolonged C-Reactive Protein  
Increased  
Cd4 Lymphocytes Decreased  
Dermatitis  
Neutrophil Count  
Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	White Blood Cell Count Increased	Report Source	Product	Role	Manufacturer	Route
				Zyban	PS	Glaxo Wellcome	ORAL

Date:09/20/01ISR Number: 3795910-2Report Type:Expedited (15-DaCompany Report #B0120302A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN	1TAB Single	Convulsion		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged dose		Loss Of Consciousness					
		Metabolic Acidosis Tachycardia		Ibuprofen	C		

Date:09/20/01ISR Number: 3795911-4Report Type:Expedited (15-DaCompany Report #D0014305A  
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 20 DAY		Angioneurotic Oedema		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged Disability	3 YR	Arthritis Gonococcal Pain In Extremity		Plastulen N Cyclo Ostrogynal	C C		ORAL ORAL
		Red Blood Cell Sedimentation Rate Increased Serum Sickness Vasculitis					

Date:09/20/01ISR Number: 3795929-1Report Type:Expedited (15-DaCompany Report #B0108747A  
Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN	300MG per day 3 WK	Abortion Spontaneous		Zyban	PS	Glaxo Wellcome	



Complications Of Maternal  
Exposure To Therapeutic  
Drugs

Date:09/20/01ISR Number: 3795931-XReport Type:Expedited (15-DaCompany Report #B0111884A  
Age:61 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Hypoxia		Zyban	PS	Glaxo Wellcome	
Hospitalization - Initial or Prolonged		Joint Effusion Respiratory Failure					

Date:09/20/01ISR Number: 3795941-2Report Type:Expedited (15-DaCompany Report #B0120405A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Per day		Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY (INHALATION)	2PUFF Twice	Joint Dislocation Spinal Fracture		Alcohol Salbutamol	SS C	Glaxo Wellcome	
per day				Seretide	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)	2PUFF Twice						
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/20/01ISR Number: 3795942-4Report Type:Expedited (15-DaCompany Report #B0120407A  
Age:49 YR Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG As Initial or Prolonged directed	Psoriasis		Zyban	PS	Glaxo Wellcome	ORAL

Date:09/20/01ISR Number: 3795943-6Report Type:Expedited (15-DaCompany Report #B0120468A  
Age:48 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Per day 20 DAY Initial or Prolonged	Back Pain Oesophageal Disorder Urticaria	Health Professional	Zyntabac Unspecified Medication Gliclazide	PS C C	Glaxo Wellcome	

Date:09/20/01ISR Number: 3795944-8Report Type:Expedited (15-DaCompany Report #B0120523A  
Age:61 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 42 DAY UNKNOWN	Cardiac Failure Myocardial Fibrosis Sudden Death	Consumer	Zyban Sotalol Warfarin	PS C C	Glaxo Wellcome	ORAL

Date:09/21/01ISR Number: 3796366-6Report Type:Expedited (15-DaCompany Report #A0055937A  
Age:55 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Disability	Agoraphobia Alopecia Anaphylactic Shock Anxiety

Bipolar Disorder  
Blood Triglycerides  
Increased  
Bradycardia  
Chills  
Condition Aggravated  
Convulsion  
Cystocele  
Dizziness  
Drug Dependence  
Drug Hypersensitivity  
Dysarthria  
Dyspepsia  
Dyspnoea  
Dysuria  
Fatigue  
Haematuria  
Headache  
Hyperglycaemia  
Hypertension  
Hyperventilation  
Hypoaesthesia  
Labile Blood Pressure  
Mania  
Memory Impairment  
Micturition Urgency  
Mitral Valve Incompetence

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	12 DAY	Muscle Twitching Muscular Weakness Musculoskeletal Stiffness Neck Pain	Zyban	PS	Glaxo Wellcome	ORAL
1.25MG Per day		Nervousness Obsessive-Compulsive Disorder	Premarin	C		
		Oedema	Vitamin	C		
		Oesophageal Disorder	Asa	C		
		Panic Disorder Without Agoraphobia	Melatonin	C		
		Paraesthesia				
		Renal Disorder				
		Renal Impairment				
		Schizophrenia				
		Sciatica				
		Sensation Of Heaviness				
		Stupor				
		Tachycardia				
		Tinnitus				
		Tobacco Abuse				
		Transient Ischaemic Attack				
		Tremor				
		Urethral Stricture				
		Urinary Incontinence				
		Urinary Retention				
		Ventricular Extrasystoles				
		Weight Increased				

Date:09/21/01ISR Number: 3796371-XReport Type:Expedited (15-DaCompany Report #A0160261A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day	Crying	Wellbutrin	PS	Glaxo Wellcome	ORAL
6 WK	2 WK	Depression	Meridia	SS		ORAL
		Diabetes Mellitus	Toprol	C		
		Memory Impairment	Hctz	C		

Mood Swings  
Palpitations  
Serotonin Syndrome  
Suicidal Ideation

Date:09/21/01ISR Number: 3796381-2Report Type:Expedited (15-DaCompany Report #B0120525A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice	Cardiac Failure		Zyban	PS	Glaxo Wellcome	ORAL
	per day	Congestive Cardiomyopathy					
	150MG per day	Drug Interaction		Aspirin	C		ORAL
	50MG per day			Atenolol	C		ORAL
	RESPIRATORY			Beclomethasone	C	Glaxo Wellcome	
	(INHALATION)	200MG per day					
	1UNIT per day			Co-Amilofruse	C		ORAL
	60MG per day			Isosorbide	C		ORAL
	15ML Twice			Lactulose	C		ORAL
	per day						
	20MG Twice			Lisinopril	C		ORAL
	per day						

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

100MG Twice			Tramadol	C		ORAL
per day						
20MG Per day			Fluoxetine	C		ORAL
10MG Per day			Cetirizine	C	Glaxo Wellcome	ORAL
TOPICAL			Betamethasone	C	Glaxo Wellcome	

Date:09/21/01ISR Number: 3796382-4Report Type:Expedited (15-DaCompany Report #B0120692A  
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	41 DAY	Asthma	Zyban	PS	Glaxo Wellcome	ORAL
			Bronchitis Chronic				
			Lower Respiratory Tract				
			Infection				

Date:09/21/01ISR Number: 3796993-6Report Type:Direct Company Report #  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Dyskinesia	Wellbutrin Sr	PS		
			Dysphagia				
			Emotional Disorder				
			Hemiparesis				
			Hyperacusis				
			Hyperaesthesia				
			Photosensitivity Reaction				
			Visual Disturbance				

Date:09/21/01ISR Number: 3797707-6Report Type:Expedited (15-DaCompany Report #B0120031A  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Required Intervention to Prevent Permanent Dose Impairment/Damage	Duration	PT	Report Source	Product	Role	Manufacturer	Route
BID		Blood Pressure Decreased	Health	Ziprasidone Po	PS		ORAL
BID		Convulsion	Professional Company	Wellbutrin Prozac	SS		
		Electrocardiogram	Representative	Depakote	SS		
		Abnormal Electrocardiogram Qt Prolonged Myocardial Infarction Overdose Sinus Tachycardia					

Date:09/24/01ISR Number: 3798115-4Report Type:Expedited (15-DaCompany Report #B0108747A  
 Age:34 YR Gender:Female I/FU:F

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abortion Spontaneous	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		

Date:09/24/01ISR Number: 3798116-6Report Type:Expedited (15-DaCompany Report #B0120405A  
 Age:44 YR Gender:Male I/FU:I

Outcome Dose Disability	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Convulsion Joint Dislocation Spinal Fracture	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

150 MG / PER

DAY/ ORAL

Ethanol (Formulation Unknown) (Alcohol)	SS
Salbutamol Sulphate	C
Fluticasone+Salmeterol	C



Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Aggression Angioneurotic Oedema Balance Disorder	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG TWICE PER DAY ORAL	Blood Pressure Increased Circulatory Collapse Convulsion Disorientation Dizziness Dry Skin Irritable Bowel Syndrome Rhinitis Stress Urticaria					

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Freedom Of Information (FOI) Report

Date:09/24/01ISR Number: 3798563-2Report Type:Expedited (15-DaCompany Report #B0120407A  
Age:49 YR Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG AS DIRECTED / ORAL	Psoriasis	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:09/24/01ISR Number: 3798565-6Report Type:Expedited (15-DaCompany Report #B0120302A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1 TABLET SINGLE DOSE	Convulsion Loss Of Consciousness Metabolic Acidosis Tachycardia	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
			Ibuprofen	C		

Date:09/24/01ISR Number: 3798637-6Report Type:Expedited (15-DaCompany Report #B0111884A  
Age:61 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Hospitalization - Initial or Prolonged 150 MG	Hypoxia Joint Effusion Respiratory Failure	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		

Date:09/24/01ISR Number: 3798686-8Report Type:Expedited (15-DaCompany Report #D0020665A  
Age: Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Cardiac Arrest	Foreign Literature Health	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL				Professional				

Date:09/24/01ISR Number: 3798687-XReport Type:Expedited (15-DaCompany Report #B0120523A  
 Age:61 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Cardiac Failure Myocardial Fibrosis Sudden Death	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / ORAL					Sotalol Warfarin Sodium	C C		

Date:09/24/01ISR Number: 3798688-1Report Type:Expedited (15-DaCompany Report #B0120468A  
 Age:48 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Back Pain Oesophageal Disorder Urticaria	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
150 MG / PER					Gliclazide	C		

DAY 20 DAY

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/24/01ISR Number: 3802045-9Report Type:Expedited (15-DaCompany Report #D0014305A  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Disability		Angioneurotic Oedema Antinuclear Antibody Positive	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ ORAL		Pain In Extremity Serum Sickness Urticaria Vasculitis		Plastulen N Cycloostrogynal	C C		

Date:09/24/01ISR Number: 3802048-4Report Type:Expedited (15-DaCompany Report #B0119910A  
Age:42 YR Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Bacterial Infection C-Reactive Protein Increased	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL		Dermatitis Lymphocyte Count Decreased Neutrophil Count Increased White Blood Cell Count Increased					

Date:09/25/01ISR Number: 3797887-2Report Type:Expedited (15-DaCompany Report #A0161283A  
Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day	10 DAY	Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:09/25/01ISR Number: 3797888-4Report Type:Expedited (15-DaCompany Report #A0161303A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Per day			Abortion Spontaneous	Bupropion	PS	Glaxo Wellcome	ORAL
			Complications Of Maternal Exposure To Therapeutic Drugs Pregnancy				

Date:09/25/01ISR Number: 3797890-2Report Type:Expedited (15-DaCompany Report #B0097007A  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged			Arthralgia Drug Hypersensitivity Erysipelas Streptococcal Infection	Zyban	PS	Glaxo Wellcome	

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Freedom Of Information (FOI) Report

Date:09/25/01ISR Number: 3797891-4Report Type:Expedited (15-DaCompany Report #B0110588A  
Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 13 DAY Initial or Prolonged	Convulsion		Zyban	PS	Glaxo Wellcome	
	Fall Headache Insomnia Muscle Spasms Salivary Hypersecretion Sensation Of Pressure					

Date:09/25/01ISR Number: 3797896-3Report Type:Expedited (15-DaCompany Report #B0118244A  
Age:63 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150MG Twice per day	Abdominal Pain Upper Abdominal Tenderness		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
11 DAY	Angina Unstable Blood Pressure Increased Cardiac Failure Congestive Chest Pain Dyspnoea Heart Rate Increased Pharyngolaryngeal Pain					

Date:09/25/01ISR Number: 3797897-5Report Type:Expedited (15-DaCompany Report #B0119920A  
Age:30 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice per day	Discomfort		Zyban	PS	Glaxo Wellcome	ORAL
15 DAY	Face Oedema Inflammation Loss Of Consciousness		Stimulant	C		

Rash Erythematous  
Rash Papular  
Rash Pruritic  
Respiratory Disorder  
Urticaria

Date:09/25/01ISR Number: 3799243-XReport Type:Direct  
Age:37 YR Gender:Female I/FU:I

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Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG BID PO		Lacrimation Increased Pruritus Urticaria		Zyban (150 Mg) (Gsk)	PS	Gsk	ORAL

Date:09/25/01ISR Number: 3799988-1Report Type:Direct  
Age:48 YR Gender:Male I/FU:I

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Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG SA 1 PO BID		Convulsion		Bupropion (150mg Sa) (Gsk)	PS	Gsk	ORAL

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Freedom Of Information (FOI) Report

Date:09/25/01ISR Number: 3800753-7Report Type:Expedited (15-DaCompany Report #A0160261A  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Depression Diabetes Mellitus Serotonin Syndrome	Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
150 MG/PER DAY /ORAL	2	WK		Sibutramine Hydrochloride Tablet (Sibutramine Hydrochloride)	SS		ORAL
ORAL	6	WK		Metoprolol Succinate Hydrochlorothiazide	C C		

Date:09/25/01ISR Number: 3801259-1Report Type:Expedited (15-DaCompany Report #B0120525A  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Failure Congestive Cardiomyopathy	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY/ ORAL				Aspirin Atenolol Beclomethasone Dipropion. Co-Amilofruse Isosorbide Lactulose Lisinopril Tramadol Hydrochloride Fluoxetine Cetirizine	C C C C C C C C C C C		



Hydrochloride C  
Betamethasone C

Date:09/25/01ISR Number: 3801262-1Report Type:Expedited (15-DaCompany Report #B0120692A  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bronchitis Chronic	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /PER DAY/ ORAL							

Date:09/25/01ISR Number: 3806051-XReport Type:Expedited (15-DaCompany Report #A0055937A  
Age:55 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Alopecia
Disability	Anxiety
	Autonomic Nervous System
	Imbalance
	Blood Glucose Increased

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Dose	Duration	Adverse Event	Report Source	Product	Role	Manufacturer	Route
150 MG/PER	DAY/ORAL	Blood Pressure Increased Blood Triglycerides Increased Bradycardia Condition Aggravated Convulsion	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
		Drug Hypersensitivity					
		Dysarthria Dyspepsia Dyspnoea Dysthymic Disorder Enuresis Euphoric Mood Faeces Pale Feeling Hot And Cold Fluid Retention Haematuria Hyperventilation Hypoaesthesia Hypomania Insomnia Intervertebral Disc Protrusion Labile Blood Pressure Memory Impairment Mental Impairment Mitral Valve Incompetence Musculoskeletal Stiffness Oedema Palpitations Panic Disorder Paraesthesia Renal Disorder Renal Impairment Sensation Of Pressure Syncope Vasovagal Tachycardia Tinnitus Tongue Disorder Transient Ischaemic Attack Tremor Urethral Stricture Urinary Incontinence		Conjugated Estrogens Vitamin Aspirin Melatonin	C C C C		

Weight Increased

Date:09/26/01ISR Number: 3798439-0Report Type:Expedited (15-DaCompany Report #A0157251A

Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Twice			Brain Scan Abnormal	Zyban	PS	Glaxo Wellcome	ORAL
per day	2	WK	Headache				
			Multiple Sclerosis	Premarin	C		
			Musculoskeletal Stiffness	Vitamin E	C		
			Neck Pain	Vitamin C	C		
			Pyrexia	Vitamin A	C		
			Tinnitus	Multivitamin	C		

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Date:09/26/01ISR Number: 3798443-2Report Type:Expedited (15-DaCompany Report #B0118455A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG As directed		Drug Interaction Thrombosis		Zyban  Coumarin	PS  SS	Glaxo Wellcome	  ORAL

Date:09/26/01ISR Number: 3798446-8Report Type:Expedited (15-DaCompany Report #B0120825A  
Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day 25 DAY		Hypersensitivity		Zyban  Aspirin Levothyroxine	PS  C C	Glaxo Wellcome  Glaxo Wellcome	ORAL  ORAL

Date:09/26/01ISR Number: 3798447-XReport Type:Expedited (15-DaCompany Report #B0120882A  
Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG As Initial or Prolonged directed		Dyspepsia Haematemesis		Quomem	PS	Glaxo Wellcome	ORAL

Date:09/26/01ISR Number: 3798449-3Report Type:Expedited (15-DaCompany Report #D0018145A  
Age:70 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death UNKNOWN		Haemorrhagic Stroke Hypertensive Crisis		Zyban Diclofenac	PS C	Glaxo Wellcome	ORAL

Date:09/26/01ISR Number: 3798866-1Report Type:Direct  
Age:14 YR Gender:Male I/FU:I

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Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 250 MG QID PO	Dermatitis		Keflex (Cephalexin)	PS		ORAL
Initial or Prolonged 100SR 1 PO Q	Erythema Multiforme		Bupropion	SS		ORAL
Required AM X 2 THEN Intervention to BID Prevent Permanent Impairment/Damage	Pruritus					

Date:09/27/01ISR Number: 3798981-2Report Type:Expedited (15-DaCompany Report #A0141870A  
Age:68 YR Gender:Male I/FU:F

Outcome	PT
Death	Balance Disorder
Life-Threatening	Blood Creatinine
Hospitalization -	Increased
Initial or Prolonged	Blood Urea Increased
Other	Cerebral Ischaemia
	Cerebrovascular Accident
	Coma
	Depressed Level Of Consciousness
	Fall
	Grand Mal Convulsion
	Head Injury

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Dose	Duration	Insomnia Lethargy Oral Intake Reduced Rib Fracture	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	9 DAY			Wellbutrin	PS	Glaxo Wellcome	ORAL
100MG Twice per day				Celexa	C		ORAL
				Cyclosporin	C		ORAL
10MG Per day				Prednisone	C		ORAL
25MG Twice per day				Metoprolol	C		ORAL
TOPICAL	.2MG Per day			Transderm Nitro	C	Glaxo Wellcome	
10MG At night				Zocor	C		ORAL
40MG Per day				Lasix	C		ORAL
1TAB Per day				Enteric Coated Aspirin	C		ORAL
1TAB Twice per day				Tums	C		ORAL

Date:09/27/01ISR Number: 3798991-5Report Type:Expedited (15-DaCompany Report #A0161553A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Vasculitis Cerebral		Zyban	PS	Glaxo Wellcome	ORAL

Date:09/27/01ISR Number: 3798994-0Report Type:Expedited (15-DaCompany Report #B0111884A  
Age:61 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Emphysema		Zyban	PS	Glaxo Wellcome	
Hospitalization - Initial or Prolonged		Joint Effusion Respiratory Failure					

Date:09/27/01ISR Number: 3798995-2Report Type:Expedited (15-DaCompany Report #B0119050A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 18 DAY Hospitalization - Initial or Prolonged		Angioneurotic Oedema Dermatitis Dysphagia Dyspnoea Pyrexia Rash Generalised		Zyntabac	PS	Glaxo Wellcome	ORAL

Date:09/27/01ISR Number: 3798999-XReport Type:Expedited (15-DaCompany Report #B0120884A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG As directed		Blister Dizziness Sleep Disorder		Zyban	PS	Glaxo Wellcome	

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Freedom Of Information (FOI) Report

Date:09/27/01ISR Number: 3799000-4Report Type:Expedited (15-DaCompany Report #B0120885A  
Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Idiopathic		Zyban	PS	Glaxo Wellcome	
150MG Per day	4 DAY						
		Thrombocytopenic Purpura					

Date:09/27/01ISR Number: 3799001-6Report Type:Expedited (15-DaCompany Report #B0121084A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Blood Cholesterol Increased		Zyban	PS	Glaxo Wellcome	
		Myocardial Infarction					

Date:09/27/01ISR Number: 3799002-8Report Type:Expedited (15-DaCompany Report #B0121088A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsive Threshold		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	3 DAY						
Initial or Prolonged		Lowered		Trimipramine	SS		ORAL
350MG Per day							
		Drug Interaction		Fluctine	SS		ORAL
40MG Per day		Epilepsy					

Date:09/27/01ISR Number: 3799003-XReport Type:Expedited (15-DaCompany Report #B0121093A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Clumsiness		Zyban	PS	Glaxo Wellcome	ORAL
Other		Dizziness		Microgynon	C		ORAL
		Fluid Retention					
		Flushing					
		Hypertension					



Insomnia  
Weight Increased

Date:09/27/01ISR Number: 3800235-2Report Type:Direct  
Age:49 YR Gender:Female I/FU:I

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus Urticaria		Bupropion Gsk	PS		

Date:09/28/01ISR Number: 3800386-2Report Type:Expedited (15-DaCompany Report #A0161559A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion Drug Interaction		Wellbutrin Sr Ultram	PS SS	Glaxo Wellcome	ORAL

Date:09/28/01ISR Number: 3800403-XReport Type:Expedited (15-DaCompany Report #B0120883A  
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT
Hospitalization - Initial or Prolonged		Dysgeusia Nervous System Disorder

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Nervousness

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	39 DAY		Zyban	PS	Glaxo Wellcome	
750MG Twice			Amoxicillin	C		
per day	8 DAY					
100MG Per day			Hydroquinine Hydrobromide	C		

Date:09/28/01ISR Number: 3800405-3Report Type:Expedited (15-DaCompany Report #B0121198A  
 Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Sudden Death					
per day	63 DAY			Aspirin	C		ORAL
75MG per day				Lisinopril	C		ORAL
20MG per day				Simvastatin	C		ORAL
40MG per day							

Date:09/28/01ISR Number: 3801921-0Report Type:Expedited (15-DaCompany Report #A122280  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Agitation	Consumer	Zoloft Tablets	PS		
Initial or Prolonged		Cognitive Disorder		Bupropion	SS		ORAL
225.00 MG		Depression					
TOTAL:DAILY:0		Disturbance In Attention					
RAL		Hallucination					

Memory Impairment  
Mood Swings  
Personality Change

Date:09/28/01ISR Number: 3802134-9Report Type:Expedited (15-DaCompany Report #A0161283A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs	Study Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY / ORAL	10 DAY						

Date:09/28/01ISR Number: 3802135-0Report Type:Expedited (15-DaCompany Report #A0161303A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs	Study Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER DAY / ORAL		Pregnancy					

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Date:09/28/01ISR Number: 3802410-XReport Type:Expedited (15-DaCompany Report #B0110588A  
Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Convulsion Headache Insomnia Intraocular Pressure Increased	Foreign	Zyban Tablet- (Bupropion Hydochloride)	PS		

Date:09/28/01ISR Number: 3802453-6Report Type:Expedited (15-DaCompany Report #B0097007A  
Age:37 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG	Erysipelas Hypersensitivity Streptococcal Infection	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		

Date:09/28/01ISR Number: 3802456-1Report Type:Expedited (15-DaCompany Report #B0118244A  
Age:63 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG TWICE PER DAY ORAL	Abdominal Tenderness Angina Unstable Cardiac Failure Congestive Chest Pain Condition Aggravated Dyspepsia Dyspnoea Exacerbated Heart Rate Increased Pharyngolaryngeal Pain	Foreign Study Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:09/28/01ISR Number: 3802457-3Report Type:Expedited (15-DaCompany Report #B0119920A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Immunoglobulin E Increased	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG TWICE PER DAY ORAL		Inflammation					
		Loss Of Consciousness Rash Erythematous Rash Papular Rash Pruritic Respiratory Disorder Urticaria		Stimulant	C		

Date:09/28/01ISR Number: 3802660-2Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #2000UW03177

Outcome	PT
Other	Confusional State Drug Interaction Face Oedema Hyperhidrosis Hypoaesthesia Influenza Like Illness

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral Pain					
100 MG PO		Toothache	Consumer	Seroquel "Zeneca"	PS	Zeneca	ORAL
100 MG PO		Tremor		Celebrex	SS		ORAL
300 MG				Oxycodone	SS		
10 MG PRN				Wellbutrin	SS		
				Valium	SS		
				Valium	SS		
				Ambien	SS		
				Ambien	SS		

Date:10/01/01ISR Number: 3801625-4Report Type:Expedited (15-DaCompany Report #A0159343A  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Jittery		Zyban	PS	Glaxo Wellcome	ORAL
Other		Myopathy					
150MG Twice per day	6 WK			Vitamins	C		

Date:10/01/01ISR Number: 3801630-8Report Type:Expedited (15-DaCompany Report #A0161900A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pain		Wellbutrin	PS	Glaxo Wellcome	ORAL
Other		Tardive Dyskinesia					

Date:10/01/01ISR Number: 3801642-4Report Type:Expedited (15-DaCompany Report #B0121348A  
 Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Angina Pectoris Zyban PS Glaxo Wellcome  
 7 WK  
 Initial or Prolonged Dyspepsia  
 Dyspnoea

Date:10/01/01ISR Number: 3801643-6Report Type:Expedited (15-DaCompany Report #B0121383A  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	Overdose		Zyban	PS	Glaxo Wellcome	ORAL
UNKNOWN		Respiratory Failure		Desipramine	SS		
UNKNOWN				Dihydrocodeine	SS		ORAL
UNKNOWN				Imipramine	SS		
UNKNOWN				Dothiepin	C		

Date:10/01/01ISR Number: 3802965-5Report Type:Direct Company Report #  
 Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	75 MG Q AM PO	Blepharospasm		Bupropion	PS		ORAL
		Eye Movement Disorder		Olanzapine	C		
		Eyelid Ptosis		Miratazapine	C		
		Movement Disorder		Thiamine	C		
		Muscle Spasms		Lorazepam	C		
		Strabismus		Gosereline Acetate	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/01/01ISR Number: 3803192-8Report Type:Expedited (15-DaCompany Report #B0120884A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blister Dizziness Sleep Disorder	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
150 MG / AS							

DIRECTED

Date:10/01/01ISR Number: 3803324-1Report Type:Expedited (15-DaCompany Report #2001UW11959

Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Foreign Other	Lidocaine Atropine Chloral Hydrate Diazepam Methadone Sertraline Bupropion Venlafaxine	PS SS SS SS SS SS SS SS		

Date:10/01/01ISR Number: 3803801-3Report Type:Expedited (15-DaCompany Report #B0111884A

Age:61 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - Initial or Prolonged		Hypoxia Joint Effusion Respiratory Failure	Foreign Health Professional	Bupropion Hydrochloride Tablet - Zyban (Bupropion Hydrochloride)	PS		
150MG							

Date:10/01/01ISR Number: 3804109-2Report Type:Expedited (15-DaCompany Report #A0161553A

Age: Gender:Female I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hemiparesis Stress Vasculitis Cerebral	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Date:10/01/01ISR Number: 3804110-9Report Type:Expedited (15-DaCompany Report #B0119050A

Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization - Initial or Prolonged 300 MG/ORAL		Angioneurotic Oedema Dysphagia Oedema  Photosensitivity Reaction Pyrexia Rash Erythematous Rash Generalised Respiratory Disorder	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/01/01ISR Number: 3804111-0Report Type:Expedited (15-DaCompany Report #B0120885A  
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent 150 MG/PER Impairment/Damage DAY		Idiopathic Thrombocytopenic Purpura	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		

Date:10/01/01ISR Number: 3804112-2Report Type:Expedited (15-DaCompany Report #B0121084A  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death  150 MG		Blood Cholesterol Increased Myocardial Infarction	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		

Date:10/01/01ISR Number: 3804113-4Report Type:Expedited (15-DaCompany Report #B0121088A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged  150 MG/PER DAY/ORAL		Drug Interaction Epilepsy	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
350 MG/PER DAY/ORAL				Surmonatil (Formulation Unknown)	SS		ORAL
40 MG/PER				Fluctine (Formulation Unknown) (Fluctine)	SS		ORAL

DAY/ORAL

Date:10/01/01ISR Number: 3804114-6Report Type:Expedited (15-DaCompany Report #B0121093A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required		Clumsiness Dizziness Feeling Hot And Cold	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage		Fluid Retention Flushing Hypertension Insomnia Weight Increased		Microgynon	C		

Date:10/01/01ISR Number: 3804176-6Report Type:Expedited (15-DaCompany Report #A0141870A  
Age:68 YR Gender:Male I/FU:F

Outcome  
Death  
Life-Threatening  
Hospitalization -  
Initial or Prolonged  
Required  
Intervention to  
Prevent Permanent

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Freedom Of Information (FOI) Report

Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG /		Balance Disorder	Health	Wellbutrin			
		Bite	Professional	Tablet-Controlled			
		Blood Creatine	Company	Release (Bupropion			
		Phosphokinase Increased	Representative	Hydrochloride)	PS		ORAL
TWICE PER DAY		Blood Urea Increased					
/ ORAL		Coma					
		Ecchymosis		Citalompram			
		Fall		Hydrobromide	C		
		Grand Mal Convulsion		Cyclosporin	C		
		Head Injury		Prednisone	C		
		Insomnia		Metoprolol	C		
		Ischaemic Stroke		Nitroglycerin	C		
		Nervous System Disorder		Simvasatin	C		
		Oral Intake Reduced		Frusemide	C		
		Pain		Aspirin	C		
		Platelet Count Decreased		Calcium Carbonate	C		
		Rib Fracture					
		Tongue Disorder					

Date:10/02/01ISR Number: 3803547-1Report Type:Expedited (15-DaCompany Report #B0120882A  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Dyspepsia	Foreign	Zyban Tablet -Zyban			
Initial or Prolonged		Haematemesis	Health	(Bupropion			
150 MG / AS			Professional	Hydrochlorde)	PS		ORAL
DIRECTED /							
ORAL							

Date:10/02/01ISR Number: 3803641-5Report Type:Direct  
 Age:32 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsive Threshold Lowered		Bupropion 150mg Wellcome	PS	Wellcome	ORAL
150MG QAM							
ORAL				Bupropion	SS	Wellcome	ORAL
75 MG / 1PM /							
ORAL				Olanzapine	C		
				Nortriptyline	C		
				Divalproex	C		
				Mercaptopurine	C		

Date:10/02/01ISR Number: 3803643-9Report Type:Direct Company Report #  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Arthralgia Pain In Extremity Tendon Injury		Wellbutrin Sr 150 Mgs. Twice Daily Glaxo	PS	Glaxo	ORAL
150 MGS. /							
TWICE DAILY /		Urticaria					
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/02/01ISR Number: 3803909-2Report Type:Expedited (15-DaCompany Report #A0161559A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Health	Wellbutrin			
Other		Drug Interaction	Professional	Tablet-Controlled			
			Company	Release (Bupropion			
			Representative	Hydrochloride)	PS		ORAL
UNK/UNK/ORAL				Tramadol			
				Hydrochloride			
				(Formulation			
				Unknown) (Tramadol			
				Hydrochloride)	SS		

Date:10/02/01ISR Number: 3804076-1Report Type:Expedited (15-DaCompany Report #D0018145A

Age:70 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Haemorrhagic Stroke	Foreign	Zyban Tablet - Zyban			
Death		Hypertensive Crisis	Health	(Bupropion			
			Professional	Hydrochloride)	PS		ORAL
150 MG ORAL				Diclofenac	C		

Date:10/02/01ISR Number: 3804077-3Report Type:Expedited (15-DaCompany Report #B0118455A

Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Foreign	Zyban Tablet - Zyban			
Required		Thrombosis	Health	(Bupropion			
Intervention to			Professional	Hydrochloride)	PS		
Prevent Permanent							
150 MG AS				Coumarin			
Impairment/Damage				(Formulation			
DIRECTED				Unknown) (Coumarin)	SS		ORAL
ORAL							

Date:10/02/01ISR Number: 3804217-6Report Type:Expedited (15-DaCompany Report #A0157251A  
Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache	Health	Zyban Tablet - Zyban			
		Musculoskeletal Stiffness	Professional	(Bupropion			
		Neck Pain		Hydrochloride)	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL		Nervous System Disorder					
		Nuclear Magnetic		Conjugated Estrogens	C		
		Resonance Imaging		Vitamin E	C		
		Abnormal		Ascorbic Acid	C		
		Pyrexia		Retinol	C		
		Tinnitus		Multivitamin	C		

Date:10/02/01ISR Number: 3805552-8Report Type:Expedited (15-DaCompany Report #B0120883A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dysgeusia	Foreign	Zyban Tablet - Zyban			
Initial or Prolonged		Nervous System Disorder		(Bupropion			
		Nervousness		Hydrochloride)	PS		
150 MG / PER							
DAY /							
				Amoxicillin	C		

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Freedom Of Information (FOI) Report

Hydroquinine  
Hydrobromide C

Date:10/02/01ISR Number: 3805553-XReport Type:Expedited (15-DaCompany Report #B0121198A  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia Sudden Death Syncope	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							

Aspirin C  
Lisinopril C  
Simvastatin C

Date:10/02/01ISR Number: 3805564-4Report Type:Expedited (15-DaCompany Report #B0120825A  
Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage DAY / ORAL		Drug Hypersensitivity	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Aspirin C  
Thyroxine Sodium C

Date:10/03/01ISR Number: 3803375-7Report Type:Expedited (15-DaCompany Report #B0121689A  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Twice		Chronic Obstructive		Zyban	PS	Glaxo Wellcome	ORAL



per day	Pulmonary Disease		
120MG per day	Respiratory Failure	Isosorbide	C
40MG per day	Sudden Death	Nifedipine	C
		Alverine	C
		Aspirin	C

Date:10/03/01ISR Number: 3803377-0Report Type:Expedited (15-DaCompany Report #D0021164A

Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Unknown Initial or Prolonged	Anaemia		Zyban	PS	Glaxo Wellcome	ORAL
	Myeloproliferative Disorder Thrombocythaemia					

Date:10/04/01ISR Number: 3803916-XReport Type:Expedited (15-DaCompany Report #A0161749A

Age:47 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150MG Per day 4 YR	Cardiac Murmur		Wellbutrin	PS	Glaxo Wellcome	ORAL
	Mitral Valve Incompetence		Pepcid	C		
	Mitral Valve Prolapse		Prevacid	C		
			Multivitamin	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/04/01ISR Number: 3803918-3Report Type:Expedited (15-DaCompany Report #A0162004A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death				Wellbutrin	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged				Zyban	SS	Glaxo Wellcome	ORAL

Date:10/05/01ISR Number: 3804372-8Report Type:Expedited (15-DaCompany Report #A0161755A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day				Wellbutrin	PS	Glaxo Wellcome	ORAL
				Fluphenazine	C		
				Unspecified Medication	C		
				Amantadine	C		
				Depakote	C		

Date:10/05/01ISR Number: 3804376-5Report Type:Expedited (15-DaCompany Report #A0162391A

Age:22 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged				Zyban	PS	Glaxo Wellcome	ORAL
				Vitamin B	C		
				Vitamin C	C		
				Mvi	C		

Date:10/05/01ISR Number: 3804385-6Report Type:Expedited (15-DaCompany Report #B0121094A

Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization -	Confusional State	Zyban	PS	Glaxo Wellcome	ORAL
15TAB Per day 1 DAY					
Initial or Prolonged	Convulsion	Alcohol	C		ORAL
1 DAY	Hallucination				
	Hypokalaemia				
	Overdose				

Date:10/05/01ISR Number: 3806715-8Report Type:Expedited (15-DaCompany Report #A0161900A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pain	Health	Wellbutrin Tablet-			
		Tardive Dyskinesia	Professional	Controlled Release			
			Company	(Bupropion			
			Representative	Hydrochloride)	PS		ORAL
ORAL							

Date:10/05/01ISR Number: 3806720-1Report Type:Expedited (15-DaCompany Report #A0159343A  
Age:37 YR Gender:Female I/FU:F

Outcome	PT
Other	Asthenia
	Back Pain
	Difficulty In Walking

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Feeling Jittery Muscle Spasms Myopathy	Report Source	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL	6 WK		Health Professional	Zyban Tablet	PS		ORAL
				Multivitamin	C		

Date:10/05/01ISR Number: 3808950-1Report Type:Expedited (15-DaCompany Report #B0121383A  
Age:39 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG PER DAY ORAL		Overdose Renal Disorder Respiratory Failure Toxicologic Test Abnormal	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
					Desipramine (Formulation Unknown) (Desipramine)	SS		
					Dihydrocodeine (Formulation Unknown) (Dihydrocodeine)	SS		ORAL
					Imipramine (Formulation Unknown) (Imipramine)	SS		
					Dothiepin	C		

Date:10/05/01ISR Number: 3808956-2Report Type:Expedited (15-DaCompany Report #B0121348A  
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Angina Pectoris Dyspepsia Dyspnoea	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		

Date:10/08/01ISR Number: 3805036-7Report Type:Expedited (15-DaCompany Report #B0118584A  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Single dose		Circulatory Collapse Myocardial Infarction		Zyban	PS	Glaxo Wellcome	

Date:10/08/01ISR Number: 3805041-0Report Type:Expedited (15-DaCompany Report #B0122029A  
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1 DAY		Blood Gases Abnormal		Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/09/01ISR Number: 3805419-5Report Type:Expedited (15-DaCompany Report #A0159874A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	27 DAY	Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
		Blister					
		Dry Mouth					
		Insomnia					
		Muscle Spasms					
		Oedema Peripheral					
		Pyrexia					
		Serum Sickness					
		Urticaria					

Date:10/09/01ISR Number: 3805422-5Report Type:Expedited (15-DaCompany Report #A0162388A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Twice per day		Completed Suicide		Wellbutrin	PS	Glaxo Wellcome	ORAL
9 DAY				Depakote	C		
				Dexamethasone	C		

Date:10/09/01ISR Number: 3805423-7Report Type:Expedited (15-DaCompany Report #A0162626A  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900MG Per day	5 YR	Balance Disorder		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Blood Potassium Abnormal		Eskalith Cr	SS	Glaxo Wellcome	ORAL
UNKNOWN		Blood Sodium Abnormal		Zyprexa	SS		
UNKNOWN		Condition Aggravated		Prilosec	SS		
		Diplopia					

Disorientation  
Electrolyte Imbalance  
Fall  
Tremor

Date:10/09/01ISR Number: 3805427-4Report Type:Expedited (15-DaCompany Report #A0162932A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Decreased Cardiac Disorder Loss Of Consciousness		Zyban	PS	Glaxo Wellcome	ORAL

Date:10/09/01ISR Number: 3805432-8Report Type:Expedited (15-DaCompany Report #B0120704A  
Age:46 YR Gender:Male I/FU:F

Outcome	PT
Other	Agitation Breast Pain Condition Aggravated Confusional State Depression Disturbance In Attention Dizziness Feeling Abnormal Hallucinations, Mixed





Ranitidine C Glaxo Wellcome  
Aspirin C

75MG per day

Date:10/10/01ISR Number: 3806165-4Report Type:Expedited (15-DaCompany Report #A0143570A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Bundle Branch Block	Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice Hospitalization - per day			Completed Suicide				
Initial or Prolonged 40MG Per day			Intentional Misuse	Celexa	C		ORAL
15MG Twice per day			Loss Of Employment	Buspar	C		ORAL
			Suicide Attempt				
250MG Per day				Lamisil	C		ORAL
10MG Per day				Zyrtec	C	Glaxo Wellcome	ORAL
30MG Twice per day				Prevacid	C		ORAL

Date:10/10/01ISR Number: 3806173-3Report Type:Expedited (15-DaCompany Report #B0122442A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 12 DAY			Dermatitis Bullous	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged 1 DAY			Erythema Multiforme	Propofan	SS		ORAL
			Oral Soft Tissue Disorder	Lysanxia	SS		
			Rash Pustular	Stilnox	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/10/01ISR Number: 3808086-XReport Type:Expedited (15-DaCompany Report #D0021164A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anaemia Myeloproliferative Disorder	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / ORAL		Thrombocythaemia					

Date:10/10/01ISR Number: 3808153-0Report Type:Expedited (15-DaCompany Report #A0161749A

Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cardiac Murmur Mitral Valve Incompetence Mitral Valve Prolapse	Health Professional	Wellbutrin Tablet- Controlled Release (Bupropion Hydrochloried)	PS		ORAL
150 MG/ PER				Famotidine	C		
				Lansoprazole	C		
				Multivitamin	C		
DAY/ORAL							

Date:10/10/01ISR Number: 3808154-2Report Type:Expedited (15-DaCompany Report #A0162004A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - Initial or Prolonged		Cardiac Failure	Consumer	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Zyban Tablet- Zyban (Bupropion Hydrochloride)	SS		ORAL
ORAL							

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Corneal Oedema	Consumer	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY / ORAL				Fluphenazine Amantadine Semisodium Valproate	C C C		

Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Fall	Health Professional	Detrol (Tolterodine) Tablet	PS		ORAL
ORAL 3285 DAY	2 DAY	Haematoma		Wellbutrin (Amfebutam one Hydrochloride)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/10/01ISR Number: 3809150-1Report Type:Expedited (15-DaCompany Report #2001023091-1  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - SEE IMAGE	5 YR	Balance Disorder	Consumer	Eskalith	PS	Smith Beecham	ORAL
Initial or Prolonged		Condition Aggravated		Wellbutrin			
		Diplopia		(Bupropion)	SS	Glaxowellcome	
		Disorientation		Zyprexa	SS	Lilly	
		Electrolyte Imbalance		Prilosec			
		Fall		(Omeprazole)	SS	Astrazeneca	
		Tremor					

Date:10/10/01ISR Number: 3809257-9Report Type:Expedited (15-DaCompany Report #WAES 01072442  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	25	Abdominal Distension	Health	Tab Vioxx 25 Mg	PS		ORAL
Hospitalization - MG/DAILY/ORAL		Anxiety	Professional				
Initial or Prolonged		Aphasia	Company	Anafranil	SS		
25 MG/BID							
Other		Asthenia	Representative	Klonopin	SS		
0.5 MG/BID							
100 MG/BID		Balance Disorder		Lamictal	SS		
150 MG/BID		Blood Pressure Systolic		Wellbutrin Sr	SS		
250 MG		Increased		Dipentum	SS		
1.25 MG/DAILY		Confusional State		Premarin	SS		
100 MG/DAILY		Coordination Abnormal		Zoloft	SS		
		Difficulty In Walking		Coumadin	C		
		Disorientation		Motrin	C		
		Dizziness		Lithium Carbonate	C		
		Drug Interaction					
		Dysphagia					
		Dyspnoea					
		Emotional Distress					
		Fear					

Gait Disturbance  
 Headache  
 Nausea  
 Palpitations  
 Restlessness  
 Speech Disorder  
 Transient Ischaemic  
 Attack  
 Vision Blurred

Date:10/10/01ISR Number: 3809500-6Report Type:Expedited (15-DaCompany Report #2001023368-1  
 Age:25 YR Gender:Male I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
30 MILLIGRAMS		Conjunctival Haemorrhage Conjunctival Hyperaemia	Health Professional	Paxil Glaxosmithkline	PS	Glaxosmithkline	ORAL
1.0 DAILY		Constipation					
ORAL	99 DAY	Ecchymosis Orgasm Abnormal		Wellbutrin Sr Glaxosmithkline	SS	Glaxosmithkline	ORAL
150 MILLIGRAMS							
2.0 DAILY							
ORAL	44 DAY			Wellbutrin Sr Glaxosmithkline	SS	Glaxosmithkline	ORAL
150 MILLIGRAMS							
1.0 DAILY							

Freedom Of Information (FOI) Report

ORAL 3 DAY

Date:10/10/01ISR Number: 3809549-3Report Type:Expedited (15-DaCompany Report #B0121094A  
Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 15 TABLET/PER DAY/ORAL	Confusional State Convulsion Hallucination Hypokalaemia	Foreign Health Professional	Zyban (Bupropion Hydrochloride) Ethanol	PS C		ORAL

Date:10/10/01ISR Number: 3809569-9Report Type:Expedited (15-DaCompany Report #A0162391A  
Age:22 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Bronchospasm Dermatitis Eyelid Oedema Peripheral Circulatory Failure Rash Generalised Respiratory Distress	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride) Vitamin B Ascorbic Acid M.V.I.	PS C C C		ORAL

Date:10/10/01ISR Number: 3809666-8Report Type:Expedited (15-DaCompany Report #B0121689A  
Age:62 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death 150 MG / TWICE PER DAY / ORAL	Angina Pectoris Chronic Obstructive Pulmonary Disease Mesenteric Artery Stenosis	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Respiratory Failure  
Sudden Death

Isosorbide C  
Nifedipine C  
Alverine Citrate C  
Aspirin C

Date:10/12/01ISR Number: 3807770-1Report Type:Expedited (15-DaCompany Report #A0067430A  
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	3 YR	Anaemia Asthenia		Wellbutrin	PS	Glaxo Wellcome	ORAL
10MG Per day		Blood Iron Increased Confusional State		Lamictal Paxil	SS SS	Glaxo Wellcome Glaxo Wellcome	ORAL ORAL
UNKNOWN		Dizziness		Celexa	SS		
INTRAVENOUS		Dyspnoea		Blood Transfusion	SS		
		Fatigue Haematocrit Decreased Headache Memory Impairment Myelofibrosis Nausea Nervousness Palpitations Pruritus Rash Papular		Zyprexa Glucophage Allegra Ativan	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/12/01ISR Number: 3807777-4Report Type:Expedited (15-DaCompany Report #A0143021A  
Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300MG per day	Condition Aggravated		Wellbutrin	PS	Glaxo Wellcome	ORAL
Hospitalization -	100MG Three	Confusional State		Amantadine	SS		ORAL
Initial or Prolonged	times per day	Delirium					
	YR	Drug Toxicity		Prozac	C		
		Extrapyramidal Disorder		Loxapine	C		
		Fall		Seroquel	C		
		Feeling Abnormal		Risperdal	C		
		Hallucination, Auditory					
		Injury					
		Insomnia					
		Memory Impairment					
		Paranoia					
		Schizophrenia					
		Tremor					

Date:10/12/01ISR Number: 3807782-8Report Type:Expedited (15-DaCompany Report #A0157251A  
Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Twice	Brain Stem Auditory		Zyban	PS	Glaxo Wellcome	ORAL
per day	2 WK	Evoked Response Abnormal					
		Deafness Neurosensory		Premarin	C		
		Headache		Vitamin E	C		
		Multiple Sclerosis		Vitamin C	C		
		Musculoskeletal Stiffness		Vitamin A	C		
		Neck Pain		Multivitamin	C		
		Nervous System Disorder					
		Pyrexia					
		Tinnitus					

Date:10/12/01ISR Number: 3807786-5Report Type:Expedited (15-DaCompany Report #A0162941A  
Age: Gender:Female I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Brain Neoplasm		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL

Date:10/12/01ISR Number: 3807787-7Report Type:Expedited (15-DaCompany Report #A0162942A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Brain Neoplasm		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL

Date:10/12/01ISR Number: 3807793-2Report Type:Expedited (15-DaCompany Report #B0111329A  
 Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Hepatic Congestion		Zyntabac	PS	Glaxo Wellcome	ORAL
55 DAY		Hepatomegaly Myocardial Infarction Pulmonary Haemorrhage Pulmonary Oedema Thyroid Disorder Thyroid Neoplasm					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/12/01ISR Number: 3807794-4Report Type:Expedited (15-DaCompany Report #B0112406A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Zyntabac	PS	Glaxo Wellcome	

Date:10/12/01ISR Number: 3807797-XReport Type:Expedited (15-DaCompany Report #B0114445A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day	61 DAY	Arthralgia Difficulty In Walking Dyspnoea Headache Malaise Nausea Pain In Extremity Polymyalgia Rheumatica Red Blood Cell Sedimentation Rate Increased Sedation Tremor		Zyban	PS	Glaxo Wellcome	

Date:10/12/01ISR Number: 3807810-XReport Type:Expedited (15-DaCompany Report #B0121704A  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG Per day	9 DAY	Asthma		Zyban	PS	Glaxo Wellcome	ORAL
				Aspirin	C		ORAL

Date:10/12/01ISR Number: 3808484-4Report Type:Expedited (15-DaCompany Report #B0122029A  
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Blood Gases Abnormal	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:10/12/01ISR Number: 3808886-6Report Type:Expedited (15-DaCompany Report #B0118584A  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Circulatory Collapse Myocardial Infarction Pulse Absent	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
150 MG							
SINGLE DOSE							

Date:10/15/01ISR Number: 3808163-3Report Type:Expedited (15-DaCompany Report #D0018145A  
Age:70 YR Gender:Male I/FU:F

Outcome	PT
Death	Bradycardia
Hospitalization - Initial or Prolonged	Brain Oedema Cerebral Ischaemia Depressed Level Of

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Consciousness Dyspnoea Haemorrhagic Stroke Hemiplegia	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Hypertension		Zyban	PS	Glaxo Wellcome	ORAL
UNKNOWN	100MG As	Hypotension		Diclofenac	C		
required		Loss Of Consciousness		Anti-Hyperlipidaemic (Unspecified)	C		
UNKNOWN				Trental	C		
UNKNOWN							

Date:10/16/01ISR Number: 3808985-9Report Type:Expedited (15-DaCompany Report #B0122683A  
Age:71 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	4 DAY	Chronic Obstructive		Zyban	PS	Glaxo Wellcome	ORAL
	UNKNOWN	150MG	Airways Disease Twice Exacerbated		Ranitidine	C	Glaxo Wellcome	
per day			Respiratory Failure		Alendronate Sodium	C		
UNKNOWN	10MG Per day				Gaviscon Advance	C		
UNKNOWN	8ML	Three			Alfuzosin	C		
times per day					Combivent	C		
UNKNOWN	5MG	Twice per						
day								
RESPIRATORY								
(INHALATION)	2.5ML	Four						
times per day					Aminophylline	C		
UNKNOWN	225MG	As						

required

Salmeterol C Glaxo Wellcome

RESPIRATORY

(INHALATION)

Fluticasone C Glaxo Wellcome

UNKNOWN 250MCG Per

day

Date:10/16/01ISR Number: 3809424-4Report Type:Direct  
Age:27 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Wellbutrin Sr (150)	PS		ORAL
150; 1 PO TID		Grand Mal Convulsion Medication Error					

Date:10/16/01ISR Number: 3809425-6Report Type:Direct  
Age:20 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin Sr (150 1 Bid)	PS		ORAL
150 BID PO							

Date:10/16/01ISR Number: 3810151-8Report Type:Expedited (15-DaCompany Report #B0122089A  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Coronary Artery Disease Coronary Artery Occlusion	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /							

TWICE PER DAY

/ ORAL

Bisoprolol C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Diclofenac C  
 Ranitidine C  
 Hydrochloride C  
 Aspirin C

Date:10/16/01ISR Number: 3810152-XReport Type:Expedited (15-DaCompany Report #B0121882A  
 Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Toxicity Headache	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG ORAL				Olanzapine (Formulation Unknown) (Olanzapine)	SS		ORAL
ORAL				Methadone Hydrochloride (Formulation Unknown) (Methadone Hydrochloride)	SS		
				Lithium Carbonate	C		
				Carbimazole	C		

Date:10/16/01ISR Number: 3810153-1Report Type:Expedited (15-DaCompany Report #A0162932A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Pressure Decreased Cardiac Disorder Loss Of Consciousness	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / ORAL							

Date:10/16/01ISR Number: 3810154-3Report Type:Expedited (15-DaCompany Report #B0120704A  
 Age:46 YR Gender:Male I/FU:I

Outcome PT

Other

Agitation  
Anxiety  
Breast Pain  
Confusional State  
Depression  
Dermatitis  
Disturbance In Attention  
Dizziness  
Fatigue  
Feeling Abnormal  
Hallucinations, Mixed  
Insomnia  
Irritability  
Mental Disorder  
Mood Altered  
Nausea  
Pain  
Pollakiuria  
Pruritus  
Psoriasis  
Seborrhoea  
Sleep Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /		Tremor Visual Disturbance	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
TWICE PER DAY				Zopiclone	C		
				Oxazepam	C		
				Naproxen	C		

Date:10/16/01ISR Number: 3810156-7Report Type:Expedited (15-DaCompany Report #A0159874A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthralgia Blister Drug Hypersensitivity	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG TWICE		Dry Mouth					
PER DAY ORAL		Insomnia Oedema Peripheral Serum Sickness					

Date:10/16/01ISR Number: 3810209-3Report Type:Expedited (15-DaCompany Report #B0122442A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dermatitis Bullous Erythema Multiforme Mouth Ulceration	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ORAL				Propofan (Formulation Unknown) (Propofan)	SS		ORAL
ORAL				Prazepam			



UNKNOWN

(Formulation  
Unknown) (Prasepam) SS

Zolpidem  
(Formulation  
Unknown)( Zolpidem) SS

UNKNOWN

Date:10/16/01ISR Number: 3810447-XReport Type:Expedited (15-DaCompany Report #A118566

Age:27 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Blood Pressure Systolic
Initial or Prolonged	Decreased
Required	Bundle Branch Block Left
Intervention to	Convulsion
Prevent Permanent	Electrocardiogram
Impairment/Damage	Abnormal
	Electrocardiogram Qrs
	Complex Prolonged
	Electrocardiogram Qt
	Corrected Interval
	Prolonged
	Electrocardiogram St
	Segment Abnormal
	Infarction

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Overdose Sinus Tachycardia				
Dose	Duration		Report Source	Product	Role	Manufacturer
			Health	Ziprasidone Po	PS	
BID			Professional Company Representative	Wellbutrin (Bupropion) Prozac (Fluoxetine)	SS SS	
BID				Depakote (Divalproex Sodium)	SS	

Date:10/16/01ISR Number: 3810504-8Report Type:Expedited (15-DaCompany Report #A0143570A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bundle Branch Block	Health	Wellbutrin			
Hospitalization - Initial or Prolonged		Completed Suicide Intentional Misuse Suicide Attempt	Professional	Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY / ORAL		Toxicologic Test Abnormal		Citalopram Hydrobromide Buspirone Hydrochloride Terbinafine Cetirizine Hydrochloride Lansoprazole	C C C C C		

Date:10/16/01ISR Number: 3810515-2Report Type:Expedited (15-DaCompany Report #A0162626A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Balance Disorder Condition Aggravated	Consumer	Wellbutrin Unspecified Tablet			

ORAL	Diplopia	(Bupropion		
	Disorientation	Hydrochloride)	PS	ORAL
	Electrolyte Imbalance	Lithium Carbonate		
	Fall	(Formulation		
	Tremor	Unknown) (Lithium		
900 MG / PER		Carbonate)	SS	ORAL
DAY / ORAL				
		Olanzapine		
		(Formulation		
		Unknown)		
		(Olanzapine)	SS	
		Omeprazole		
		(Formulation		
		Unknown)		
		(Omeprazole)	SS	

Date:10/16/01ISR Number: 3810517-6Report Type:Expedited (15-DaCompany Report #A0162388A  
Age:49 YR Gender:Male I/FU:I

Outcome	PT	Report Source
Death	Completed Suicide	Health
	Injury Asphyxiation	Professional
		Company

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Representative

Dose	Duration	Product	Role	Manufacturer	Route
150 MG / TWICE PER DAY / ORAL		Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
		Semisodium Valproate	C		
		Dexamethasone	C		

Date:10/17/01ISR Number: 3809799-6Report Type:Expedited (15-DaCompany Report #A0153715A  
Age:17 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200MG Twice per day	1 YR	Depression Malabsorption Medication Residue		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:10/17/01ISR Number: 3809810-2Report Type:Expedited (15-DaCompany Report #A0163486A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly 300MG Per day		Atrial Septal Defect Complications Of Maternal Exposure To Therapeutic Drugs Pregnancy Pulmonary Valve Stenosis Congenital		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:10/17/01ISR Number: 3809812-6Report Type:Expedited (15-DaCompany Report #A0163666A  
Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Three		Drug Level Above					
times per day		Therapeutic		Lamictal	C	Glaxo Wellcome	ORAL
225MG Twice							
per day				Zyprexa	C		ORAL
35MG per day				Luvox	C		ORAL

Date:10/18/01ISR Number: 3810573-5Report Type:Expedited (15-DaCompany Report #B0106367A  
Age:16 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Dizziness					
Hospitalization -							
per day 21 DAY		Dyspnoea		Microgynon	C		ORAL
Initial or Prolonged							
1TAB Per day		Rash Generalised					
Disability		Visual Disturbance					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/18/01ISR Number: 3810574-7Report Type:Expedited (15-DaCompany Report #B0110958A  
Age:16 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150MG Twice	Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - per day 21 DAY	Dizziness					
Initial or Prolonged 1TAB Per day	Dyspnoea		Microgynon	C		ORAL
Disability	Rash Generalised Visual Disturbance					

Date:10/18/01ISR Number: 3810577-2Report Type:Expedited (15-DaCompany Report #B0123297A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Arrhythmia Circulatory Collapse		Zyban	PS	Glaxo Wellcome	

Date:10/18/01ISR Number: 3810578-4Report Type:Expedited (15-DaCompany Report #D0021744A  
Age:35 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150MG Unknown 4 DAY	Erythema		Zyban	PS	Glaxo Wellcome	ORAL
	Fatigue		Nitrendipin	C		ORAL
	Oedema Peripheral		Bisoprolol	C		ORAL

Date:10/18/01ISR Number: 3811102-2Report Type:Direct Company Report #  
Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG PO QD	Angioneurotic Oedema		Wellbutrin Sr	PS		ORAL
			Zoloft	C		

Date:10/18/01ISR Number: 3811478-6Report Type:Expedited (15-DaCompany Report #B0112406A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
150 MG							

Date:10/18/01ISR Number: 3811504-4Report Type:Expedited (15-DaCompany Report #A0143021A  
Age:53 YR Gender:Female I/FU:F

Outcome	PT
Death	Confusional State
Hospitalization - Initial or Prolonged	Delirium Extrapyramidal Disorder Fall Feeling Abnormal Hallucination, Auditory Insomnia Limb Injury Medication Error Memory Impairment Paranoia Schizophrenia Toxicologic Test Abnormal

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
SEE TEXT /		Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL			Amantadine Tablet (Amantadine Hydrochloride)	SS		ORAL
100 MG/ THREE						
TIMES PER						
DAY/ ORAL			Fluoxetine Hydrochloride	C		
			Loxapine	C		
			Quetiapine Fumarate	C		
			Risperidone	C		

Date:10/18/01ISR Number: 3811510-XReport Type:Expedited (15-DaCompany Report #A0157251A  
 Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness Neurosensory Headache Multiple Sclerosis	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE		Musculoskeletal Stiffness					
PER DAY/ ORAL		Neck Pain		Conjugated Estrogens	C		
		Pyrexia		Vitamin E	C		
		Tinnitus		Ascorbic Acid	C		
				Retinol	C		
				Multivitamin	C		



Date:10/18/01ISR Number: 3811519-6Report Type:Expedited (15-DaCompany Report #B0111329A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Hepatic Congestion Hepatomegaly Hyperplasia	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ORAL		Myocardial Infarction Pulmonary Haemorrhage Pulmonary Oedema Thyroid Neoplasm					

Date:10/18/01ISR Number: 3811578-0Report Type:Expedited (15-DaCompany Report #B0121704A  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent 300 MG/PER Impairment/Damage DAY/ORAL		Asthma	Foreign	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
				Aspirin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/18/01ISR Number: 3811774-2Report Type:Expedited (15-DaCompany Report #A0162942A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Brain Neoplasm	Health Professional	Wellbutrin Tablet - Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Date:10/18/01ISR Number: 3811776-6Report Type:Expedited (15-DaCompany Report #A0162941A

Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Brain Neoplasm	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Date:10/18/01ISR Number: 3811828-0Report Type:Expedited (15-DaCompany Report #A0067430A

Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaemia Anxiety Asthenia Biopsy Bone Marrow	Consumer	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG /							
TWICE PER DAY							
/ ORAL							
ORAL							
		Abnormal Blood Iron Increased Confusional State Depression		Lamictal Tablet (Lamotrigine)	SS		ORAL
		Dizziness Dyspnoea Fatigue Headache		Paroxetine Hydrochloride (Formulation Unknown) (Paroxetine			

10 MG / PER	Memory Impairment	Hydrochloride)	SS	ORAL
DAY / ORAL	Myelofibrosis			
	Nausea	Citalopram		
	Nervousness	Hydrobromide		
	Palpitations	(Formulation		
	Pruritus	Unknown) (Citalopram		
	Rash Papular	Hydrobromide)	SS	
		Blood (Formulation		
		Uknown) (Blood)	SS	
INTRAVENOUS	EVERY TWO			
WEEKS /				
INTRAVENOUS		Olanzapine	C	
		Metformin		
		Hydrochloride	C	
		Fexofenadine		
		Hydrochlorid	C	
		Lorazepam	C	

Date:10/18/01ISR Number: 3812060-7Report Type:Expedited (15-DaCompany Report #B0114445A  
Age: Gender:Female I/FU:F

Outcome PT  
Other Arthralgia  
Asthenia  
Decreased Activity

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG /	TWICE PER DAY	Difficulty In Walking Dyspnoea Fatigue Feeling Abnormal Headache Ill-Defined Disorder Insomnia Malaise Pain In Extremity Polymyalgia Rheumatica Red Blood Cell Sedimentation Rate Increased Sedation Tremor	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		

Date:10/19/01ISR Number: 3811174-5Report Type:Expedited (15-DaCompany Report #A0163484A  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	3 WK	Abnormal Behaviour Hostility Social Avoidant Behaviour Social Phobia		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:10/19/01ISR Number: 3811181-2Report Type:Expedited (15-DaCompany Report #B0121084A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Twice per day	20 DAY	Coronary Artery Disease Myocardial Infarction Myocardial Ischaemia		Zyban	PS	Glaxo Wellcome	

Date:10/19/01ISR Number: 3811188-5Report Type:Expedited (15-DaCompany Report #B0123296A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Decreased Appetite		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	10 DAY	Malaise					
225MG Per day		Psoriasis		Cyclosporin	C		ORAL
		Skin Infection Vomiting					

Date:10/19/01ISR Number: 3812553-2Report Type:Expedited (15-DaCompany Report #B0122683A  
Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Chronic Obstructive Airways Disease Exacerbated	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER DAY / ORAL		Respiratory Failure		Ranitidine Hydrochloride	C		
				Alendronate Sodium	C		
				Gaviscon Advance	C		
				Alfuzosin	C		
				Combivent	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Aminophylline C  
 Salmeterol Xinafoate C  
 Fluticasone  
 Propionate C

Date:10/22/01ISR Number: 3813327-9Report Type:Direct  
 Age:47 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG ONCE		Disturbance In Attention		Wellbutrin 150 Mg Sr	PS	Sr	ORAL
PO QD							

Date:10/22/01ISR Number: 3813341-3Report Type:Direct  
 Age:31 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG QD X 3		Tachycardia		Zyban	PS		
THEN BID				Nicorette	C		

Date:10/22/01ISR Number: 3813800-3Report Type:Expedited (15-DaCompany Report #A0055937A  
 Age:55 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Agitation
Initial or Prolonged	Agoraphobia
Disability	Anaphylactic Shock
	Anxiety
	Arrhythmia
	Supraventricular
	Autonomic Nervous System
	Imbalance
	Blood Glucose Increased
	Blood Triglycerides
	Increased

Bradycardia  
Cognitive Disorder  
Condition Aggravated  
Convulsion  
Delusion  
Depersonalisation  
Depression  
Drug Hypersensitivity  
Dyspnoea  
Dysthymic Disorder  
Haematuria  
Hallucination  
Hyperhidrosis  
Hypertension  
Hypomania  
Intervertebral Disc  
Degeneration  
Labile Blood Pressure  
Memory Impairment  
Mental Impairment  
Mitral Valve Incompetence  
Mood Swings  
Muscle Spasms

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	12 DAY	Neurosis Obsessive-Compulsive Disorder	Zyban	PS	Glaxo Wellcome	ORAL
1.25MG Per day		Palpitations Panic Attack Paranoia Psychomotor Hyperactivity Renal Disorder Renal Impairment Schizophrenia Sciatica Speech Disorder Stress Stupor Tachycardia Tinnitus Tobacco Abuse Transient Ischaemic Attack Urethral Stricture Urinary Incontinence Ventricular Extrasystoles Weight Increased	Premarin  Vitamin Asa Melatonin	C  C C		

Date:10/22/01ISR Number: 3813810-6Report Type:Expedited (15-DaCompany Report #B0121094A  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day	3 WK	Agitation		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged 2250MG Single Other dose	1 DAY	Confusional State Delirium		Zyban	SS	Glaxo Wellcome	ORAL
1 DAY		Grand Mal Convulsion		Alcohol	SS		ORAL
		Hallucination Hypokalaemia Leukocytosis Overdose					



Date:10/22/01ISR Number: 3813817-9Report Type:Expedited (15-DaCompany Report #B0123589A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day 6 DAY	Angioneurotic Oedema		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged		Depressed Level Of Consciousness Ecchymosis Panic Reaction					

Date:10/22/01ISR Number: 3813818-0Report Type:Expedited (15-DaCompany Report #B0123591A  
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Twice	Bronchospasm		Zyban	PS	Glaxo Wellcome	ORAL
per day	15 DAY			Salbutamol	C	Glaxo Wellcome	
RESPIRATORY ( INHALATION)							
RESPIRATORY ( INHALATION)				Dihydrocodeine Eformoterol	C C		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

60MG per day			Indoramin	C	Glaxo Wellcome	ORAL
RESPIRATORY			Ipratropium	C	Glaxo Wellcome	
(INHALATION)						
RESPIRATORY			Ipratropium	C	Glaxo Wellcome	
(INHALATION)	8PUFF per day					

Date:10/22/01ISR Number: 3813820-9Report Type:Expedited (15-DaCompany Report #B0123592A  
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxo Wellcome	ORAL
Other		Hyperaesthesia					
150MG Twice		Micturition Urgency					
per day	31 DAY	Penis Disorder					
		Pollakiuria					
		Premature Ejaculation					

Date:10/22/01ISR Number: 3813821-0Report Type:Expedited (15-DaCompany Report #B0123594A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxo Wellcome	
Hospitalization -		Hepatitis					
1UNIT Per day	15 DAY			Microval	SS		ORAL
Initial or Prolonged		Hypokalaemia		Cholstat	SS		ORAL

Date:10/23/01ISR Number: 3811928-5Report Type:Expedited (15-DaCompany Report #A0155443A  
Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxo Wellcome	ORAL
Hospitalization -		Condition Aggravated					
150MG Per day							

Initial or Prolonged	Convulsion	Vioxx	SS	
Other	Fall	Celebrex	SS	
14 DAY				
	Head Injury	Neurontin	C	ORAL
300MG Four				
	Headache			
times per day	YR			
	Inflammation			
	Migraine			
	Mucous Membrane Disorder			
	Paralysis			
	Pruritus			
	Sinus Headache			
	Sinus Polyp			
	Swelling			

Date:10/23/01ISR Number: 3811944-3Report Type:Expedited (15-DaCompany Report #D0021964A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Arrhythmia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	33 DAY	Loss Of Consciousness					
		Palpitations					
		Pulse Absent					
		Tachycardia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/23/01ISR Number: 3812699-9Report Type:Expedited (15-DaCompany Report #B0106367A  
 Age:16 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening Hospitalization - Initial or Prolonged	Arthralgia Dizziness Dyspnoea	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE Disability PER DAY/ORAL	Rash Generalised Visual Disturbance		Microgynon	C		

Date:10/23/01ISR Number: 3812884-6Report Type:Expedited (15-DaCompany Report #A0163666A  
 Age:33 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Convulsion Drug Level Above Therapeutic Drug Level Below Therapeutic	Health Professional Company Representative	Wellbutrin-Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / THREE TIMES PER DAY / ORAL			Lamotrigine Olanzapine Fluvoxamine Maleate	C C C		

Date:10/23/01ISR Number: 3812886-XReport Type:Expedited (15-DaCompany Report #A0163486A  
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Congenital Anomaly	Atrial Septal Defect Complications Of Maternal Exposure To Therapeutic Drugs	Study Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
300 MG / PER						

DAY / ORAL

Pregnancy

Pulmonary Valve Stenosis  
Congenital

Date:10/23/01ISR Number: 3813572-2Report Type:Expedited (15-DaCompany Report #D0018145A

Age:70 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bradycardia	Foreign	Zyban Tablet - Zyban			
Hospitalization - Initial or Prolonged		Brain Oedema	Health	(Bupropion			
150 MG PER		Cerebral Ischaemia	Professional	Hydrochloride)	PS		ORAL
DAY ORAL		Depressed Level Of					
		Consciousness		Diclofenac	C		
		Dyspnoea		Anti-Hyperlipidemic	C		
		Haemorrhagic Stroke		Oxpentifylline	C		
		Hypertension					
		Loss Of Consciousness					
		Muscle Spasms					
		Pupils Unequal					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/23/01ISR Number: 3813664-8Report Type:Expedited (15-DaCompany Report #B0110958A  
Age:16 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - Initial or Prolonged 150 MG TWICE Disability PER DAY/ ORAL						
	Arthropathy	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
	Dizziness					
	Dyspnoea					
	Rash Generalised					
	Tenderness		Mycrogynon	C		
	Visual Disturbance					

Date:10/23/01ISR Number: 3813724-1Report Type:Expedited (15-DaCompany Report #B0123297A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG						
	Arrhythmia	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
	Circulatory Collapse	Health Professional				

Date:10/23/01ISR Number: 3813732-0Report Type:Expedited (15-DaCompany Report #D0021744A  
Age:35 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150 MG / ORAL						
	Erythema	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
	Fatigue	Health				
	Oedema Peripheral	Professional				
			Nitrendipine	C		
			Bisoprolol	C		

Date:10/23/01ISR Number: 3814284-1Report Type:Expedited (15-DaCompany Report #A0153715A  
Age:17 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Required Intervention to Prevent Permanent Impairment/Damage 200/MG/TWICE	Depression Malabsorption Medication Residue	Health Professional	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS	ORAL
PER DAY/ORAL					

Date:10/24/01ISR Number: 3812620-3Report Type:Expedited (15-DaCompany Report #A0121035A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG Twice Initial or Prolonged per day		Pancreatitis		Wellbutrin	PS	Glaxo Wellcome	ORAL
Other 10MG Per day	104 DAY			Adderall	C		ORAL
37.5MG Per day				Effexor	C		ORAL

Date:10/24/01ISR Number: 3812629-XReport Type:Expedited (15-DaCompany Report #B0122925A  
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 4 DAY Initial or Prolonged 8UNIT per day		Anxiety		Zyban	PS	Glaxo Wellcome	ORAL
2MG As required		Dyskinesia		Co-Proxamol	C		ORAL
				Diazepam	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Diclofenac	C		ORAL
		Paraffin	C		
250MG Four		Erythromycin	C	Glaxo Wellcome	ORAL
times per day					
RESPIRATORY		Fluticasone	C	Glaxo Wellcome	
(INHALATION)	125MCG Twice				
per day					
TOPICAL		Gtn	C	Glaxo Wellcome	
		Hydrocortisone	C	Glaxo Wellcome	
120MG per day		Isosorbide	C		ORAL
RESPIRATORY		Ipratropium	C	Glaxo Wellcome	
(INHALATION)					
TOPICAL		Mucogel	C		ORAL
AURICULAR		Oilatum Plus	C		
(OTIC)		Otomize	C		
300MG per day		Quinine Sulphate	C		ORAL
150MG per day		Ranitidine	C	Glaxo Wellcome	ORAL
RESPIRATORY		Salbutamol	C	Glaxo Wellcome	
(INHALATION)					
RESPIRATORY		Salmeterol	C	Glaxo Wellcome	
(INHALATION)	50MCG Twice				
per day					
OPHTHALMIC	1DROP Four	Sodium Cromoglycate	C		
times per day					



100MG Three		Tramadol	C		ORAL
times per day					
300MG Per day		Diltiazem	C	Glaxo Wellcome	ORAL
TOPICAL		Aqueous Cream	C		
75MG per day		Aspirin	C		ORAL
TRANSDERMAL		Gtn	C	Glaxo Wellcome	
RESPIRATORY		Ipratropium	C	Glaxo Wellcome	
(INHALATION)	250MCG Four				
times per day					
RESPIRATORY		Salbutamol	C	Glaxo Wellcome	
(INHALATION)	2.5MG As				
required					
AURICULAR		Locorten Vioform	C		
(OTIC)	2DROP Twice				
per day					

Date:10/24/01ISR Number: 3814906-5Report Type:Expedited (15-DaCompany Report #B0123296A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Decreased Appetite Malaise Pain	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE		Psoriasis					
PER DAY/ORAL		Skin Infection Vomiting		Cyclosporin	C		

Date:10/24/01ISR Number: 3815064-3Report Type:Expedited (15-DaCompany Report #A0163484A  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hostility Phobia Social Avoidant Behaviour	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion			

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Freedom Of Information (FOI) Report

150 MG/ TWICE  
 PER DAY/ ORAL 3 WK  
 Hydrochloride) PS ORAL

Date:10/24/01ISR Number: 3815090-4Report Type:Expedited (15-DaCompany Report #A0143021A  
 Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Condition Aggravated	Health	Bupropion			
Hospitalization - Initial or Prolonged		Confusional State Delirium Drug Toxicity Fall	Professional	Hydrochloride Tablet - Controlled Release (Bupropion Hydrochloride)	PS		ORAL
SEE TEXT/ORAL		Memory Impairment Paranoia Schizophrenia		Amantadine Tablet (Amantadine Hydrochloride)	SS		ORAL
SEE TEXT/ORAL		Tremor		Fluoxetine Hydrochloride Loxapine Quetiapine Fumarate Risperidone Lorazepam	C C C C C		

Date:10/24/01ISR Number: 3815293-9Report Type:Expedited (15-DaCompany Report #B0121084A  
 Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coronary Artery Disease Myocardial Infarction Myocardial Ischaemia	Foreign Health Professional	Zyban Tablet- (Bupropion Hydrochloride)	PS		
150 MG							
TWICE PER DAY							

Date:10/25/01ISR Number: 3814631-0Report Type:Direct Company Report #  
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG PO	Acute Respiratory Distress Syndrome Laryngeal Oedema Medication Error Obstructive Airways Disorder		Zyban  Serevent Azmacort	PS  C C		ORAL

Date:10/25/01ISR Number: 3819398-8Report Type:Periodic Company Report #HQ4496309AUG2001  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged OVERDOSE,  ORAL		Overdose	Health Professional	Ativan (Lorazepam, Tablet)	PS		ORAL
OVERDOSE,  ORAL				Duract (Bromfenac, Capsule)	SS		ORAL
OVERDOSE,  ORAL				Meridia (Sibutramine Hydrochloride, )	SS		ORAL
OVERDOSE,  ORAL				Prozac (Fluoxetine Hydrochloride)	SS		ORAL

Freedom Of Information (FOI) Report

OVERDOSE,  ORAL	Serzone (Mefazodone Hydrochloride, )	SS	ORAL
 OVERDOSE,  ORAL	 Wellbutrin (Amfebutamone Hydrochloride, )	 SS	 ORAL

Date:10/26/01ISR Number: 3814734-0Report Type:Expedited (15-DaCompany Report #A0093320A  
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Agitation		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	307 DAY	Asthenia					
		Crying		Azithromycin	C		
		Eye Disorder		Zantac	C	Glaxo Wellcome	
		Facial Palsy		Nasonex	C		
		Hypoaesthesia					
		Irritability					
		Mental Disorder					
		Transient Ischaemic Attack					

Date:10/26/01ISR Number: 3814738-8Report Type:Expedited (15-DaCompany Report #A0164646A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Complications Of Maternal Exposure To Therapeutic Drugs		Zyban No Concurrent Medications	PS  C	Glaxo Wellcome	ORAL
		Intra-Uterine Death					
		Umbilical Cord Abnormality					

Date:10/26/01ISR Number: 3814744-3Report Type:Expedited (15-DaCompany Report #B0105568A  
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebrovascular Accident		Zyban	PS	Glaxo Wellcome	ORAL
		Dizziness		Levothyroxine	C	Glaxo Wellcome	ORAL
10MCG Twice							
per day							
5MG Per day				Cilazapril	C		

Date:10/26/01ISR Number: 3814745-5Report Type:Expedited (15-DaCompany Report #B0105578A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Liver Function Test					
per day	32 DAY	Abnormal		Mirtazapine	C		
30MG Per day		Nausea		Pravastatin	C		
20MG Per day				Aspirin	C		
75MG Per day				Omeprazole	C		
20MG Per day				Atenolol	C		ORAL
25MG Per day				Isosorbide Mononitrate	C		
60MG Per day				Ramipril	C		ORAL
2MG Per day							

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Freedom Of Information (FOI) Report

20MG Per day

Date:10/26/01ISR Number: 3814746-7Report Type:Expedited (15-DaCompany Report #B0114323A  
 Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	100MG Three	Balance Disorder		Wellbutrin	PS	Glaxo Wellcome	ORAL
Hospitalization -	times per day	Deafness					
Initial or Prolonged		Depression		Citalopram	C		
	300MG Twice	Fall		Sodium Valproate	C		
	per day	Feeling Jittery					
	40MG per day	Loss Of Consciousness		Fluoxetine	C		
1 DAY		Multiple Sclerosis		Zopiclone	C		ORAL
		Orthostatic Hypotension					
		Photopsia					
		Road Traffic Accident					
		Syncope					
		Tooth Injury					
		Visual Acuity Reduced					

Date:10/26/01ISR Number: 3814747-9Report Type:Expedited (15-DaCompany Report #B0114925A  
 Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	24 DAY	Amnesia		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Diplopia					
Disability		Ecchymosis					
Other		Eye Pain					
		Facial Bones Fracture					
		Sedation					
		Sleep Disorder					
		Wrist Fracture					

Date:10/26/01ISR Number: 3814754-6Report Type:Expedited (15-DaCompany Report #B0123953A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypersensitivity		Zyban	PS	Glaxo Wellcome	ORAL
150MG	Per day	5 DAY					

Date:10/26/01ISR Number: 3814755-8Report Type:Expedited (15-DaCompany Report #B0123954A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Eruption		Zyban	PS	Glaxo Wellcome	ORAL
11	DAY	Psoriasis		Salbutamol	C	Glaxo Wellcome	

RESPIRATORY

(INHALATION) 2PUFF Four

times per day

RESPIRATORY

(INHALATION) 2PUFF Three

times per day

RESPIRATORY

(INHALATION)

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/26/01ISR Number: 3816065-1Report Type:Expedited (15-DaCompany Report #A123115

Age:38 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 160.00 MG	Anxiety	Health	Ziprasidone Po	PS		ORAL
Initial or Prolonged TOTAL;BID;ORA	Lethargy	Professional				
L	Overdose		Bupropriion	SS		ORAL
100.00 MG						
TOTAL;DAILY;O						
RAL			Trazodone	SS		ORAL
ORAL			Alprazolam	SS		

Date:10/30/01ISR Number: 3816532-0Report Type:Expedited (15-DaCompany Report #A0156108A

Age:45 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300MG per day	Condition Aggravated		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Ecchymosis Haemorrhage Hysterectomy Menorrhagia Stress		Herbs Homeopathy	C C		

Date:10/30/01ISR Number: 3816543-5Report Type:Expedited (15-DaCompany Report #B0123957A

Age:42 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice	Laryngeal Oedema		Zyntabac	PS	Glaxo Wellcome	ORAL
per day	Mouth Ulceration					
22 DAY						

Date:10/30/01ISR Number: 3816545-9Report Type:Expedited (15-DaCompany Report #B0124107A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Twice	Angina Pectoris		Zyban	PS	Glaxo Wellcome	
Hospitalization - per day		Feeling Hot					
Initial or Prolonged							
Other							

Date:10/31/01ISR Number: 3817276-1Report Type:Expedited (15-DaCompany Report #A0155864A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Infectious Mononucleosis Neutropenia		Wellbutrin No Concurrent Medications	PS C	Glaxo Wellcome	ORAL

Date:10/31/01ISR Number: 3817278-5Report Type:Expedited (15-DaCompany Report #A0162401A  
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Drug Effect Decreased Mood Swings		Zyban Topamax	PS SS	Glaxo Wellcome	ORAL
UNKNOWN		Suicidal Ideation		Zyprexa	SS		
UNKNOWN				Klonopin	SS		
UNKNOWN				Ziprasidone	SS		

Freedom Of Information (FOI) Report

UNKNOWN

Depakote SS

Date:10/31/01ISR Number: 3817284-0Report Type:Expedited (15-DaCompany Report #B0107035A  
Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2 WK	Anxiety		Zyban	PS	Glaxo Wellcome	ORAL
		Decreased Appetite Depression Disturbance In Attention Irritability Mood Swings Vision Blurred		Chloroquine	C		

Date:10/31/01ISR Number: 3817291-8Report Type:Expedited (15-DaCompany Report #B0124387A  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	13 DAY	Abdominal Pain		Zyban	PS	Glaxo Wellcome	ORAL
Other SUBLINGUAL		Albuminuria Condition Aggravated Haematuria Malignant Hypertension Pancreatitis Retinopathy Hypertensive		Glyceryl Trinitrate	C	Glaxo Wellcome	

Date:10/31/01ISR Number: 3817292-XReport Type:Expedited (15-DaCompany Report #B0124405A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anaphylactic Shock Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 50.00 MG		Balance Disorder	Consumer	Zoloft Tablets	PS		
Intervention to TOTAL:DAILY		Cerebrovascular Accident					
Prevent Permanent 300.00 MG		Convulsion		Wellbutrin	SS		ORAL
Impairment/Damage TOTAL:PID:ORA		Drug Ineffective					
L		Fall					
		Heart Rate Increased		Coumadin	SS		
		Hostility		Effexor Sr	SS		ORAL
300.00 MG		Hyperhidrosis					
TOTAL:PID:ORA		Laceration					
L		Mental Impairment		Toprol	C		
		Prothrombin Time		Zebeta	C		
		Prolonged		Lanoxin	C		
		Vaginal Haemorrhage		Lasix	C		
				Synthroid	C		
				Vitamin E	C		
				Estrogen	C		
				Amantadine	C		
				Zocor	C		
				Estradiol Vaginal			
				Ring	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/01/01ISR Number: 3817868-XReport Type:Expedited (15-DaCompany Report #A0159004A

Age:57 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice	Bradycardia		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	Syncope		Aspirin	C		ORAL
81MG Per day			Glucophage	C		
			Nitroglycerin	C	Glaxo Wellcome	
			Altace	C		
			Zocor	C		
			Niaspan	C		

Date:11/01/01ISR Number: 3817869-1Report Type:Expedited (15-DaCompany Report #A0163666A

Age:33 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Three	Condition Aggravated		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged times per day	Convulsion					
Other 300MG Twice	Depression		Lamictal	C	Glaxo Wellcome	ORAL
per day	Drug Level Above					
35MG per day	Therapeutic		Zyprexa	C		ORAL
	Grand Mal Convulsion		Luvox	C		ORAL
			Remeron	C		
			Lithium	C		

Date:11/01/01ISR Number: 3817875-7Report Type:Expedited (15-DaCompany Report #A0165315A

Age:40 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice	Deafness		Wellbutrin	PS	Glaxo Wellcome	ORAL

Tinnitus  
 per day 10 DAY  
 Paxil C  
 Klonopin C  
 Glaxo Wellcome

Date:11/01/01ISR Number: 3817884-8Report Type:Expedited (15-DaCompany Report #B0120884A  
 Age:73 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aortic Aneurysm		Zyban	PS	Glaxo Wellcome	
150MG As		Blister					
directed		Dizziness					
		Insomnia					
		Tremor					

Date:11/02/01ISR Number: 3818796-6Report Type:Expedited (15-DaCompany Report #A0165224A  
 Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Complications Of Maternal		Wellbutrin	PS	Glaxo Wellcome	ORAL
200MG Twice		Exposure To Therapeutic					
per day		Drugs		Buspar	C		
		Oesophageal Atresia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/02/01ISR Number: 3818801-7Report Type:Expedited (15-DaCompany Report #B0101026A  
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coronary Artery Disease		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice per day	18 DAY	Sudden Death					
				Bisoprolol	C		
10MG At night				Voltarol Sr	C	Glaxo Wellcome	
UNKNOWN	75MG Twice						
				Ranitidine	C	Glaxo Wellcome	
				Aspirin Ec	C		
UNKNOWN							

Date:11/02/01ISR Number: 3818803-0Report Type:Expedited (15-DaCompany Report #B0114323A  
Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Balance Disorder		Wellbutrin	PS	Glaxo Wellcome	ORAL
100MG Three Hospitalization - times per day		Deafness					
Initial or Prolonged		Depression		Citalopram	C		
300MG Twice		Fall		Sodium Valproate	C		
		Feeling Jittery					
per day		Loss Of Consciousness		Fluoxetine	C		
40MG per day		Nervous System Disorder		Zopiclone	C		ORAL
1 DAY		Orthostatic Hypotension					
		Overdose					
		Photopsia					
		Road Traffic Accident					
		Syncope					
		Tooth Loss					
		Tremor					
		Visual Disturbance					

Date:11/02/01ISR Number: 3818804-2Report Type:Expedited (15-DaCompany Report #B0115008A  
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Toxicity		Zyban	PS	Glaxo Wellcome	
		Physical Assault					

Date:11/02/01ISR Number: 3818805-4Report Type:Expedited (15-DaCompany Report #B0118455A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deep Vein Thrombosis		Zyban	PS	Glaxo Wellcome	
150MG As							
directed		Drug Interaction					
				Coumarin	SS		ORAL

Date:11/02/01ISR Number: 3818807-8Report Type:Expedited (15-DaCompany Report #B0122089A  
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Coronary Artery Disease		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	18 DAY	Coronary Artery Occlusion					
				Bisoprolol	C		ORAL
10MG Per day							
				Diclofenac	C		
75MG Twice							
per day							
				Ranitidine	C	Glaxo Wellcome	
				Aspirin	C		
75MG per day							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/02/01ISR Number: 3818808-XReport Type:Expedited (15-DaCompany Report #B0123963A  
 Age:65 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Disorder		Zyntabac	PS	Glaxo Wellcome	ORAL
150MG Per day	23 DAY	Hypertension		Enalapril	C		
20MG Per day		Monoparesis Transient Ischaemic Attack					

Date:11/02/01ISR Number: 3819944-4Report Type:Expedited (15-DaCompany Report #200110-1486 (0)  
 Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Bite Grand Mal Convulsion	Health Professional	Sudafed - Dosage Form Unspecified (Pseudoephedrine)	PS		ORAL
PER ORAL		Tongue Disorder		Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
150 MG BID,							
PER ORAL				Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		
				Oral Contraceptive Quetiapine Fumarate (Quetiapine Fumarate)	C		

Date:11/05/01ISR Number: 3820684-6Report Type:Expedited (15-DaCompany Report #02720  
 Age:46 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Literature	Naproxen	PS		

Bupropine Sr

SS

Date:11/05/01ISR Number: 3820703-7Report Type:Expedited (15-DaCompany Report #02597

Age:42 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Carbamazepine	PS		
		Intentional Misuse		Bupropion	SS		

Date:11/05/01ISR Number: 3820704-9Report Type:Expedited (15-DaCompany Report #02611

Age:37 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Carbamazepine	PS		
		Intentional Misuse		Bupropion	SS		
				Olanzapine	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/05/01ISR Number: 3820725-6Report Type:Expedited (15-DaCompany Report #02612

Age:36 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Clonazepam	PS		
		Intentional Misuse		Trazodone	SS		
				Bupropion	SS		

Date:11/05/01ISR Number: 3820727-XReport Type:Expedited (15-DaCompany Report #02608

Age:45 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Clonidine	PS		
		Intentional Misuse	Health	Amtriptyline	SS		
			Professional	Bupropion	SS		

Date:11/05/01ISR Number: 3820746-3Report Type:Expedited (15-DaCompany Report #02613

Age:21 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Clonazepam	PS		
		Intentional Misuse		Bupropion	SS		
				Olanzapine	SS		

Date:11/05/01ISR Number: 3820763-3Report Type:Expedited (15-DaCompany Report #02663

Age:46 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Alprazolam	PS		
		Intentional Misuse		Bupropion	SS		
				Olanzapine	SS		

Date:11/06/01ISR Number: 3820310-6Report Type:Expedited (15-DaCompany Report #A0165053A

Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 12 YR Initial or Prolonged		Bacterial Infection Crying Derealisation Drug Withdrawal Syndrome Pregnancy		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:11/06/01ISR Number: 3820314-3Report Type:Expedited (15-DaCompany Report #B0123589A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Per day 6 DAY Hospitalization - Initial or Prolonged		Angioneurotic Oedema Convulsion Depressed Level Of Consciousness Ecchymosis Panic Reaction		Zyban	PS	Glaxo Wellcome	ORAL

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Freedom Of Information (FOI) Report

Date:11/06/01ISR Number: 3820317-9Report Type:Expedited (15-DaCompany Report #B0124304A  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG		Irritability		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged Variable dose		Logorrhoea					
2700MG Single dose	1 DAY	Nausea		Zyban	SS	Glaxo Wellcome	ORAL
		Overdose					
		Suicide Attempt		Alcohol	C		
		Tearfulness					

Date:11/07/01ISR Number: 3820835-3Report Type:Expedited (15-DaCompany Report #A0164667A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Pain In Extremity		Zyban	PS	Glaxo Wellcome	ORAL
4 MON		Tremor					

Date:11/07/01ISR Number: 3820836-5Report Type:Expedited (15-DaCompany Report #A0164944A  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG See		Amnesia		Wellbutrin	PS	Glaxo Wellcome	ORAL
dosage text		Blood Potassium Decreased					
60MG Per day		Dizziness		Prozac	C		ORAL
		Grand Mal Convulsion					
		Stress					

Date:11/07/01ISR Number: 3820839-0Report Type:Expedited (15-DaCompany Report #B0108772A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Bupropion Hcl Aspirin	PS C	Glaxo Wellcome	ORAL ORAL
150MG Per day				Atenolol	C		ORAL
1UNIT Per day				Beclomethasone	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)	200MCG Twice						
per day				Co-Amilofruse	C		ORAL
1UNIT Per day				Imdur	C		ORAL
60MG Per day				Lactulose	C		ORAL
15ML Twice							
per day				Lisinopril	C		ORAL
20MG Twice							
per day				Zydol	C		ORAL
100MG Twice							
per day				Fluoxetine	C		ORAL
20MG Per day				Cetirizine	C	Glaxo Wellcome	ORAL
10MG Per day				Betnovate	C	Glaxo Wellcome	
TOPICAL	.1PCT Twice						
per day							

Date:11/07/01ISR Number: 3820842-0Report Type:Expedited (15-DaCompany Report #B0120525A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Failure		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Congestive Cardiomyopathy					
per day		Drug Interaction		Aspirin	C		ORAL
150MG per day							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

50MG per day		Atenolol	C		ORAL
RESPIRATORY		Beclomethasone	C	Glaxo Wellcome	
(INHALATION)	200MG per day				
1UNIT per day		Co-Amilofruse	C		ORAL
60MG per day		Isosorbide	C		ORAL
15ML Twice		Lactulose	C		ORAL
per day					
20MG Twice		Lisinopril	C		ORAL
per day					
100MG Twice		Tramadol	C		ORAL
per day					
20MG Per day		Fluoxetine	C		ORAL
10MG Per day		Cetirizine	C	Glaxo Wellcome	ORAL
TOPICAL		Betamethasone	C	Glaxo Wellcome	

Date:11/07/01ISR Number: 3823287-2Report Type:Expedited (15-DaCompany Report #M0697-2001  
 Age:39 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 22.5 MGS PO	Asthenia	Literature	Remeron	PS		ORAL
Initial or Prolonged L.I.D.	Body Temperature Increased		Valproic Acid	SS		ORAL
500 MGS PO	Electrocardiogram					
QHS	Abnormal		Bupropion	SS		ORAL
150 MG PO BID						



7.5 MG PO AT      Respiratory Rate      Bupropion      SS      ORAL  
 Increased  
 NOON  
 Supraventricular  
 Tachycardia

Date:11/08/01ISR Number: 3821680-5Report Type:Expedited (15-DaCompany Report #B0124639A  
 Age:68 YR    Gender:Male      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Erythema Psoriasis		Zyban	PS	Glaxo Wellcome	

Date:11/08/01ISR Number: 3821682-9Report Type:Expedited (15-DaCompany Report #B0125311A  
 Age:47 YR    Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 4    WK		Dyspepsia  Glomerulonephritis Nephrotic Syndrome Weight Increased		Zyban	PS	Glaxo Wellcome	ORAL

Date:11/08/01ISR Number: 3821683-0Report Type:Expedited (15-DaCompany Report #D0020665A  
 Age:      Gender:Unknown      I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction		Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/08/01ISR Number: 3822367-5Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Tongue Oedema		Wellbutrin 150mg Tid	PS		ORAL
150MG PO TID						
Required			Zocor	C		
Intervention to			Diltiazem	C		
Prevent Permanent			Aspirin	C		
Impairment/Damage			Atenol	C		
			Glucotrol	C		

Date:11/08/01ISR Number: 3822847-2Report Type:Expedited (15-DaCompany Report #A0159165A  
 Age:65 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Abnormal Behaviour	Consumer	Wellbutrin Sr			
	Amnesia		Tablet-Controlled			
	Conjunctival		Release (Bupropion			
	Discolouration		Hydrochloride)	PS		ORAL
SEE DOSAGE						
	Deafness					
TEXT / ORAL						
	Depression		Metoprolol Succinate			
	Drug Interaction		(Formulation			
	Liver Function Test		Unknown) (Metoprolol			
	Abnormal		Succinate)	SS		ORAL
50 MG PER DAY						
	Tinnitus					
ORAL						
	Tooth Disorder		Nitroglycerin	C		
			Ranitidine			
			Hydrochloride	C		
			Hydroxyzine	C		

Date:11/08/01ISR Number: 3823862-5Report Type:Expedited (15-DaCompany Report #B0125004A  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Aggression	Foreign	Zyban Tablet - Zyban			

Initial or Prolonged	Depression	Consumer	(Bupropion	
150 MG	Fatigue		Hydrochloride)	PS
	Loss Of Consciousness			
	Weight Decreased			

Date:11/08/01ISR Number: 3823893-5Report Type:Expedited (15-DaCompany Report #B0123286A  
 Age:25 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening Hospitalization - Initial or Prolonged 150 MG/ PER Required DAY	Angioneurotic Oedema Convulsion Depressed Level Of Consciousness	Foreign Health Professional Company	Zyban Tablet- Zyban (Bupropion Hydrochloride)	PS		
Intervention to Prevent Permanent Impairment/Damage	Head Injury Hyperventilation Obstructive Airways Disorder Panic Reaction	Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/08/01ISR Number: 3823907-2Report Type:Expedited (15-DaCompany Report #B0124852A  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Aggression	Foreign	Zyban Tablet- Zyban			
Intervention to		Dyskinesia		(Bupropion			
Prevent Permanent		Fall		Hydrochloride)	PS		ORAL
150 MG/TWICE							
Impairment/Damage		Head Injury					
PER DAY/ORAL		Loss Of Consciousness		Bendrofluazide	C		

Date:11/08/01ISR Number: 3823918-7Report Type:Expedited (15-DaCompany Report #B0115303A  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Bite	Foreign	Zyban Tablet - Zyban			
Hospitalization -		Confusional State	Health	(Bupropion			
Initial or Prolonged		Decreased Activity	Professional	Hydrochloride)	PS		
150 MG							
Other		Dyspnoea		Dianette	C		
Required		Fatigue					
Intervention to		Grand Mal Convulsion					
Prevent Permanent		Loss Of Consciousness					
Impairment/Damage		Tongue Disorder					

Date:11/08/01ISR Number: 3823920-5Report Type:Expedited (15-DaCompany Report #B0119682A  
Age:18 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Acute Psychosis	Foreign	Zyban Tablet -Zyban			
		Anger	Health	(Bupropion			
		Depression	Professional	Hydrochloride)	PS		ORAL
150 MG/ ORAL							
		Overdose		Paracetamol			
		Suicidal Ideation		(Formulation			
				Unknown)			
				(Acetaminophen)	SS		

10 TABLETS

PER DAY

Paracetamol +  
Codeine Po4  
(Formulation  
Unknown)  
(Acetaminophen + SS

10 TABLETS

PER DAY

Mersyndol  
(Formulation  
Unknown) (Mersyndol) SS

Date:11/08/01ISR Number: 3823945-XReport Type:Expedited (15-DaCompany Report #B0122882A

Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability		Ageusia Rash Erythematous	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

150 MG/ AS

DIRECTED/

ORAL 2 WK

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Date:11/13/01ISR Number: 3823065-4Report Type:Expedited (15-DaCompany Report #A0157251A

Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	2 WK		Cerebrovascular Disorder	Zyban	PS	Glaxo Wellcome	ORAL
			Deafness Neurosensory				
			Headache	Premarin	C		
			Joint Stiffness	Vitamin E	C		
			Neck Pain	Vitamin C	C		
			Pyrexia	Vitamin A	C		
			Tinnitus	Multivitamin	C		

Date:11/13/01ISR Number: 3823066-6Report Type:Expedited (15-DaCompany Report #A0160261A

Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150MG Per day	2 WK		Amnesia	Wellbutrin	PS	Glaxo Wellcome	ORAL
6 WK			Crying	Meridia	SS		ORAL
			Diabetes Mellitus	Toprol	C		
			Feeling Abnormal	Hctz	C		
			Mood Swings				
			Palpitations				
			Psychomotor Hyperactivity				
			Serotonin Syndrome				
			Suicidal Ideation				

Date:11/13/01ISR Number: 3823067-8Report Type:Expedited (15-DaCompany Report #A0163660A

Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice per day	14 DAY		Peyronie'S Disease	Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day			Priapism				
Disability 22 DAY			Scar	Catapres Tts	C		

Other

Desyrel

C

Date:11/13/01ISR Number: 3823068-XReport Type:Expedited (15-DaCompany Report #A0165555A  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Vomiting		Wellbutrin Alcohol Hydrocodone	PS SS C	Glaxo Wellcome	

Date:11/13/01ISR Number: 3823069-1Report Type:Expedited (15-DaCompany Report #A0166622A  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 3 WK		Agitation  Cholelithiasis Convulsion Muscle Twitching Nightmare		Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/13/01ISR Number: 3823071-XReport Type:Expedited (15-DaCompany Report #A0166978A  
Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Per day		Complications Of Maternal Exposure To Therapeutic Drugs					

Date:11/13/01ISR Number: 3823073-3Report Type:Expedited (15-DaCompany Report #B0111855A  
Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG Twice		Coronary Artery Disease		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	7 WK	Diplopia					
Other		Echocardiogram Abnormal		Citalopram	C		
20MG Per day		Hypercholesterolaemia		Thiamine	C		
300MG Per day		Myocardial Infarction		Alprazolam	C		
.25MG Twice		Nystagmus					
per day		Respiratory Failure		Fluticasone	C	Glaxo Wellcome	
2U Per day		Supraventricular		Salmeterol	C	Glaxo Wellcome	
2U Per day		Extrasystoles Tachycardia Ventricular Extrasystoles Vith Nerve Paralysis					

Date:11/13/01ISR Number: 3823079-4Report Type:Expedited (15-DaCompany Report #B0124304A  
Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Life-Threatening 150MG As Hospitalization - directed 9 DAY	Abnormal Behaviour Anger Dry Mouth Dry Skin	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged 2700MG Single dose 1 DAY	Dysuria Excitability Heart Rate Increased Impulsive Behaviour Insomnia Irritability Logorrhoea Mydriasis Nervousness Overdose Sedation Suicide Attempt Tearfulness Vomiting	Zyban  Alcohol	SS  C	Glaxo Wellcome	ORAL

Date:11/13/01ISR Number: 3823086-1Report Type:Expedited (15-DaCompany Report #B0125839A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2 DAY		Circulatory Collapse Face Oedema Shock Urticaria		Zyban	PS	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/14/01ISR Number: 3823315-4Report Type:Expedited (15-DaCompany Report #B0120884A  
 Age:73 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death				Zyban	PS	Glaxo Wellcome	
150MG As							
directed	9	DAY					

Date:11/14/01ISR Number: 3823325-7Report Type:Expedited (15-DaCompany Report #B0125993A  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening				Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Hospitalization -							
per day							
Initial or Prolonged				Alcohol	C		
Other							

Date:11/14/01ISR Number: 3823326-9Report Type:Expedited (15-DaCompany Report #B0125995A  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death				Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day							
INTRAMUSCULAR	1	DAY		Hydrocortisone	C	Glaxo Wellcome	

Date:11/14/01ISR Number: 3823327-0Report Type:Expedited (15-DaCompany Report #B0125999A  
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Asthma		Zyban	PS	Glaxo Wellcome	ORAL
9 DAY				Fluticasone	C	Glaxo Wellcome	

RESPIRATORY  
(INHALATION) 1G Twice per day

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Montelukast	C		ORAL
				Combivent	C		

RESPIRATORY  
(INHALATION)

Date:11/14/01ISR Number: 3823328-2Report Type:Expedited (15-DaCompany Report #B0126006A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Sudden Death		Zyban	PS	Glaxo Wellcome	

Date:11/14/01ISR Number: 3825082-7Report Type:Expedited (15-DaCompany Report #2001UW14424  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Balance Disorder	Other	Prilosec	PS		
Initial or Prolonged		Condition Aggravated		Zyprexa	SS		
		Disorientation		Eskalith	SS		
		Fall		Wellbutrin	SS		

Freedom Of Information (FOI) Report

Date:11/15/01ISR Number: 3824478-7Report Type:Expedited (15-DaCompany Report #A0160255A  
 Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Arrhythmia		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - per day	14 DAY	Asthenia					
Initial or Prolonged 10MG Twice		Atrioventricular Block		Thioridazine	C		ORAL
per day	5 YR	Blood Creatine Phosphokinase Increased Bradycardia Face Oedema Hypotension Loss Of Consciousness Pruritus Red Blood Cell Sedimentation Rate Increased Scrotal Oedema Serum Sickness Sinoatrial Block Urticaria White Blood Cell Count Increased					

Date:11/15/01ISR Number: 3824490-8Report Type:Expedited (15-DaCompany Report #B0126090A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 15 DAY		Hypoaesthesia		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Monoparesis					

Date:11/19/01ISR Number: 3825687-3Report Type:Expedited (15-DaCompany Report #A0132245A  
 Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 150MG Twice Initial or Prolonged per day 1 MON	Blindness Unilateral	Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice per day	Deafness				
	Dizziness	Zyban	SS	Glaxo Wellcome	ORAL
	Hyperglycaemia				
	Hypoglycaemia	Multivitamin	C		
	Nervous System Disorder	Vitamin E	C		
	Tinnitus	Vitamin C	C		
		Aspirin	C		

Date:11/19/01ISR Number: 3825689-7Report Type:Expedited (15-DaCompany Report #A0162932A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Angiopathy Blood Pressure Decreased Cardiac Disorder Loss Of Consciousness		Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/19/01ISR Number: 3825693-9Report Type:Expedited (15-DaCompany Report #A0166999A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice		Complications Of Maternal					
per day		Exposure To Therapeutic Drugs		Synthroid	C	Glaxo Wellcome	

Date:11/19/01ISR Number: 3825694-0Report Type:Expedited (15-DaCompany Report #A0167029A  
Age:72 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aneurysm		Wellbutrin	PS	Glaxo Wellcome	ORAL
200MG Twice		Dizziness					
per day	4 MON	Drug Withdrawal Syndrome					

Date:11/19/01ISR Number: 3825708-8Report Type:Expedited (15-DaCompany Report #B0126404A  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day 27 DAY							
Initial or Prolonged				Lansoprazole	C		ORAL

Date:11/19/01ISR Number: 3825709-XReport Type:Expedited (15-DaCompany Report #B0126406A  
Age:43 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Gingival Bleeding		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Neutropenia		Lansoprazole	C		ORAL
		Thrombocytopenia					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression	Health	Xenical (Orlistat)			
		Drug Interaction	Professional	120 Mg	PS		ORAL

120 MG 3 PER

DAY ORAL

Wellbutrin Sr (Bupropion Hydrochloride)	SS
Synthroid (Levothyroxine Sodium)	C
Dexedrine (Dextroamphetamine Sulfate)	C
Effexor Xr (Venlafaxine Hydrochloride)	C
Pindolol (Pindolol)	C
Norvasc (Amlodipine Besylate)	C
Ambien (Zolpidem Tartrate)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/19/01  
Age:57 YR  
Gender:Male  
I/FU:I

Report Type:Direct  
Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -							
3 TABS BID	1 MON	Mental Impairment		Ms Contin (60)	PS		
Initial or Prolonged		Respiratory Failure		Wellbutrin Sr	SS		
1 TAB BID	1 MON						
Other				Serzone	C		
				Neurontin	C		
				Synthroid	C		

Date:11/20/01  
Age:26 YR  
Gender:Male  
I/FU:F

Report Type:Expedited (15-Da  
Company Report #A0154004A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blister		Zyban	PS	Glaxo Wellcome	ORAL
150MG See		Burning Sensation					
dosage text	3 DAY	Burns Third Degree		No Concurrent Medication	C		
		Feeling Abnormal					
		Flat Affect					
		Insomnia					
		Oedema Peripheral					
		Pain					
		Pain In Extremity					
		Penis Disorder					
		Pruritus					
		Rash Erythematous					
		Scab					
		Scar					
		Skin Discolouration					
		Vascular Injury					

Date:11/20/01  
Age:48 YR  
Gender:Female  
I/FU:F

Report Type:Expedited (15-Da  
Company Report #A0159171A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Other		Chest Pain	Wellbutrin	PS	Glaxo Wellcome	ORAL
100MG Twice						
		Circulatory Collapse				
per day	2	MON				
		Convulsion	Phenothiazine	SS		
INTRAVENOUS						
		Dizziness	Atenolol	C		
		Drug Interaction	Lipitor	C		
		Ecchymosis	Methotrexate	C		
		Face Oedema	Folic Acid	C		
		Fall	Ambien	C		
		Head Injury	Ativan	C		
		Muscle Spasms	Zantac	C	Glaxo Wellcome	
		Nausea				
		Paralysis				
		Sedation				
		Tremor				

Date:11/20/01ISR Number: 3826288-3Report Type:Expedited (15-DaCompany Report #B0114598A  
Age:56 YR Gender:Male I/FU:F

Outcome PT  
Hospitalization - Abdominal Pain  
Initial or Prolonged Aggression  
Amnesia  
Anaphylactic Shock  
Angioneurotic Oedema

Freedom Of Information (FOI) Report

Dose	Duration	Balance Disorder Blister Blood Pressure Increased	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	62 DAY	Chest Pain Circulatory Collapse Confusional State Convulsion Diarrhoea Disorientation Dizziness Dry Skin Dysphagia Dyspnoea Feeling Drunk Feeling Hot Flatulence Gastroenteritis Hyperhidrosis Irritability Irritable Bowel Syndrome Loss Of Consciousness Mental Impairment Nausea Oedema Peripheral Pharyngeal Oedema Pruritus Rash Generalised Rhinitis Speech Disorder Tinnitus Tongue Oedema Urticaria		Zyban	PS	Glaxo Wellcome	ORAL

Date:11/20/01  
Age: Gender:Male

ISR Number: 3827397-5  
I/FU:I

Report Type:Direct  
Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other 40MG BID		Dyspnoea		Ziprasidone(Geodon)	PS		
		Throat Tightness		Bupropion Sr			

100MG BID

(Wellbutrin Sr)	SS
Clozapine	C
Ddvp	C
Lamotrigine	C
Oxcarbezeprine	C

Date:11/20/01ISR Number: 3827400-2Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Dysphagia		Ziprasidone(Geodon)	PS		ORAL
40MG PO BID							
Other		Dyspnoea		Bupropion			
150MG PO BID				Sr(Wellbutrin Sr)	SS		ORAL
				Lithium Carbonate	C		
				Clozapine	C		
				Guanfacine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/21/01ISR Number: 3827086-7Report Type:Expedited (15-DaCompany Report #A0157983A  
 Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Anaphylactic Reaction	Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice			Cardiac Arrest				
Hospitalization -	27 DAY		Cardiac Failure	Hctz	C		ORAL
per day			Cardiomegaly	Vasotec	C		ORAL
Initial or Prolonged			Coronary Artery Disease				
5MG Per day			Hypersensitivity				
10MG Per day			Loss Of Consciousness				
			Ventricular Fibrillation				
			Vomiting				

Date:11/21/01ISR Number: 3827110-1Report Type:Expedited (15-DaCompany Report #B0126786A  
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Fall	Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day			Head Injury	Nevirapine	C		ORAL
200MG per day			Headache	Stavudine	C		ORAL
40MG per day			Insomnia	Famciclovir	C	Glaxo Wellcome	
500MG per day				Lamivudine	C	Glaxo Wellcome	ORAL
150MG per day				Testosterone	C	Glaxo Wellcome	
INTRAMUSCULAR							

Date:11/21/01ISR Number: 3827111-3Report Type:Expedited (15-DaCompany Report #B0126795A  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization - Atrial Flutter  
150MG Per day 15 DAY  
Initial or Prolonged

Zyban

PS

Glaxo Wellcome

Date:11/21/01ISR Number: 3827119-8Report Type:Expedited (15-DaCompany Report #D0022804A  
Age:43 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Bowel Sounds Abnormal		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Coma		Amphetamine Sulphate	SS		ORAL
800MG per day						
	Drug Toxicity		Ritalin	SS		ORAL
60MG per day						
	Fatigue		Wine	SS		ORAL
3BT per day						
	Suicide Attempt					
	Tachycardia					

Date:11/21/01ISR Number: 3829553-9Report Type:Direct Company Report #  
Age:46 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
150 MG TAB PO	Insomnia		Zyban	PS		ORAL
BID						
			Habitrol 21mg	C		

Date:11/26/01ISR Number: 3828457-5Report Type:Expedited (15-DaCompany Report #A0167897A  
Age:53 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Completed Suicide		Wellbutrin	PS	Glaxo Wellcome	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/26/01ISR Number: 3828463-0Report Type:Expedited (15-DaCompany Report #B0126564A

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Asthma		Zyban	PS	Glaxo Wellcome	
UNKNOWN		Complications Of Maternal Exposure To Therapeutic Drugs Eczema Neonatal Disorder Pregnancy Ventricular Septal Defect					

Date:11/26/01ISR Number: 3828465-4Report Type:Expedited (15-DaCompany Report #B0126796A

Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Confusional State		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged per day	5 DAY	Feeling Of Despair Heart Rate Increased Oxygen Saturation Decreased					

Date:11/26/01ISR Number: 3828466-6Report Type:Expedited (15-DaCompany Report #B0126849A

Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day		Multiple Sclerosis		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged				Interferon Beta	SS		
UNKNOWN				Lioresal	SS		
UNKNOWN							

Date:11/26/01ISR Number: 3828471-XReport Type:Expedited (15-DaCompany Report #A0167897A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:11/26/01ISR Number: 3828477-0Report Type:Expedited (15-DaCompany Report #B0126564A  
Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly UNKNOWN		Asthma		Zyban	PS	Glaxo Wellcome	
		Complications Of Maternal Exposure To Therapeutic Drugs Eczema Neonatal Disorder Pregnancy Ventricular Septal Defect					

Date:11/26/01ISR Number: 3828479-4Report Type:Expedited (15-DaCompany Report #B0126796A  
Age:47 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Confusional State Feeling Of Despair Heart Rate Increased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Oxygen Saturation Decreased					
Dose	Duration		Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	5 DAY			Zyban	PS	Glaxo Wellcome	

Date:11/26/01ISR Number: 3828480-0Report Type:Expedited (15-DaCompany Report #B0126849A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG Per day Initial or Prolonged UNKNOWN		Multiple Sclerosis		Zyban	PS	Glaxo Wellcome	ORAL
				Interferon Beta	SS		
				Lioresal	SS		

Date:11/26/01ISR Number: 3828485-XReport Type:Expedited (15-DaCompany Report #A0167897A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:11/26/01ISR Number: 3828491-5Report Type:Expedited (15-DaCompany Report #B0126564A  
Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Congenital Anomaly UNKNOWN		Asthma		Zyban	PS	Glaxo Wellcome	
		Complications Of Maternal Exposure To Therapeutic Drugs Eczema Neonatal Disorder Pregnancy					



Ventricular Septal Defect

Date:11/26/01ISR Number: 3828493-9Report Type:Expedited (15-DaCompany Report #B0126796A  
 Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice				Zyban	PS	Glaxo Wellcome	
Initial or Prolonged per day	5 DAY	Dyspnoea					
		Feeling Of Despair					
		Heart Rate Increased					

Date:11/26/01ISR Number: 3828494-0Report Type:Expedited (15-DaCompany Report #B0126849A  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day				Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged UNKNOWN		Multiple Sclerosis		Interferon Beta	SS		
UNKNOWN				Lioresal	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/26/01ISR Number: 3828499-XReport Type:Expedited (15-DaCompany Report #A0167897A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:11/26/01ISR Number: 3828505-2Report Type:Expedited (15-DaCompany Report #B0126564A  
Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly UNKNOWN		Asthma		Zyban	PS	Glaxo Wellcome	
		Complications Of Maternal Exposure To Therapeutic Drugs Congenital Anomaly Eczema Neonatal Disorder Pregnancy Ventricular Septal Defect					

Date:11/26/01ISR Number: 3828507-6Report Type:Expedited (15-DaCompany Report #B0126796A  
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Confusional State		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged per day	5 DAY	Feeling Of Despair Heart Rate Increased Oxygen Saturation Decreased					

Date:11/26/01ISR Number: 3828508-8Report Type:Expedited (15-DaCompany Report #B0126849A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 150MG Per day Initial or Prolonged UNKNOWN	Multiple Sclerosis	Zyban	PS	Glaxo Wellcome	ORAL
		Interferon Beta	SS		
		Lioresal	SS		
UNKNOWN					

Date:11/26/01ISR Number: 3828513-1Report Type:Expedited (15-DaCompany Report #A0167897A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:11/26/01ISR Number: 3828519-2Report Type:Expedited (15-DaCompany Report #B0126564A  
Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly UNKNOWN		Asthma		Zyban	PS	Glaxo Wellcome	

Complications Of Maternal  
Exposure To Therapeutic  
Drugs  
Eczema  
Neonatal Disorder  
Pregnancy  
Ventricular Septal Defect

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/26/01ISR Number: 3828521-0Report Type:Expedited (15-DaCompany Report #B0126796A  
Age:47 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice Initial or Prolonged per day 5 DAY	Confusional State Feeling Of Despair Heart Rate Increased Oxygen Saturation Decreased		Zyban	PS	Glaxo Wellcome	

Date:11/26/01ISR Number: 3828522-2Report Type:Expedited (15-DaCompany Report #B0126849A  
Age:42 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Per day Initial or Prolonged UNKNOWN	Multiple Sclerosis		Zyban	PS	Glaxo Wellcome	ORAL
UNKNOWN			Interferon Beta	SS		
			Lioresal	SS		

Date:11/26/01ISR Number: 3828606-9Report Type:Expedited (15-DaCompany Report #301831  
Age:45 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Drug Interaction Mood Swings Tobacco Abuse		Klonopin Zyban Topamax Zyprexa Geodon Divalproex	PS I I I I I	Roche	

Date:11/27/01ISR Number: 3829634-XReport Type:Expedited (15-DaCompany Report #A0155904A  
Age:65 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Myeloid Metaplasia		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
150MG Twice							
Other		Pancytopenia					
per day	2 YR			Depakote	C		ORAL
500MG Twice							
per day							

Date:11/27/01ISR Number: 3829640-5Report Type:Expedited (15-DaCompany Report #B0101350A  
Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Chronic Obstructive		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day		Pulmonary Disease					
60MG Twice		Respiratory Failure		Isosorbide	C		
per day		Sudden Death					
20MG Twice				Nifedipine	C	Glaxo Wellcome	
per day							
				Alverine	C		
				Aspirin	C		

Date:11/27/01ISR Number: 3829642-9Report Type:Expedited (15-DaCompany Report #B0121383A  
Age:39 YR Gender:Female I/FU:F

Outcome	PT
Death	Drug Level Above Therapeutic

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Overdose Pulmonary Congestion Renal Disorder Respiratory Failure		Zyban	PS	Glaxo Wellcome	ORAL
UNKNOWN				Desipramine	SS		
				Dihydrocodeine	SS		ORAL
				Imipramine	SS		
UNKNOWN				Dothiepin	C		

Date:11/27/01ISR Number: 3829643-0Report Type:Expedited (15-DaCompany Report #B0121689A  
Age:62 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice		Chronic Obstructive Pulmonary Disease		Zyban	PS	Glaxo Wellcome	ORAL
	per day		Respiratory Failure		Isosorbide	C		
	120MG per day		Sudden Death		Nifedipine	C	Glaxo Wellcome	
	40MG per day				Alverine	C		
					Aspirin	C		

Date:11/27/01ISR Number: 3829646-6Report Type:Expedited (15-DaCompany Report #B0126090A  
Age:57 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150MG Per day	12 DAY	Hypoaesthesia Metastases To Central Nervous System Monoparesis		Zyban	PS	Glaxo Wellcome	ORAL

Date:11/27/01ISR Number: 3829655-7Report Type:Expedited (15-DaCompany Report #B0127002A  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day	4 DAY	Facial Palsy	Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY				Salbutamol	C	Glaxo Wellcome	
(INHALATION)	100MCG As						
required				Indomethacin	C		ORAL
25MG Three							
times per day							

Date:11/27/01ISR Number: 3829656-9Report Type:Expedited (15-DaCompany Report #B0127123A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Twice		Feeling Abnormal	Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -	per day	101 DAY	Suicidal Ideation				
Initial or Prolonged			Suicide Attempt	Herbicide	SS		

Date:11/27/01ISR Number: 3830476-XReport Type:Direct Company Report #  
Age:46 YR Gender:Male I/FU:I

Outcome	PT
Other	Depression Hearing Impaired Panic Attack Paranoia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Sensory Disturbance Suicidal Ideation Visual Disturbance	Report Source	Product	Role	Manufacturer	Route
1 TABLE TWICE				Zyban 150mg Glaxosmithkline	PS	Glaxosmithkline	ORAL
DAILY ORAL							

Date:11/27/01ISR Number: 3830973-7Report Type:Expedited (15-DaCompany Report #HQ8752026NOV2001  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule,	PS		ORAL
ORAL				Remeron (Mirtazapine,) Wellbutrin (Amfebutamone Hydrochloride)	SS SS		

Date:11/28/01ISR Number: 3830608-3Report Type:Expedited (15-DaCompany Report #B0113971A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Per day		Overdose		Bupropion	PS	Glaxo Wellcome	ORAL
UNKNOWN		Pulmonary Congestion Renal Disorder		Dihydrocodeine Imipramine	SS SS		ORAL
UNKNOWN		Respiratory Failure		Desipramine	SS		
UNKNOWN				Prothiaden	C		



Date:11/28/01ISR Number: 3831358-XReport Type:Expedited (15-DaCompany Report #301831  
Age:45 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Condition Aggravated	Consumer	Klonopin			
Initial or Prolonged	Drug Interaction	Other	(Clonazepam)	PS		
	Mood Swings		Zyban (Bupropion			
	Tobacco Abuse		Hydrochloride)	SS		ORAL
ORAL						
			Topamax (Topiramate)	SS		
			Zyprexa (Olanzapine)	SS		
			Geodon (Ziprasidone			
			Hydrochloride)	SS		
			Divalproex			
			(Divalproex Sodium)	SS		

Date:11/28/01ISR Number: 3831812-0Report Type:Expedited (15-DaCompany Report #NSADSS2001034911  
Age:45 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Mood Swings	Consumer	Topamax			
Initial or Prolonged	Suicidal Ideation		(Unspecified)			
			(Topiramate)	PS		
			Zyban (Amfebutamone			
			Hydrochloride)	SS		ORAL
			Olanzapine			
			(Olanzapine)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Clonazepam  
(Clonazepam) SS

Date:11/28/01ISR Number: 3842958-5Report Type:Periodic Company Report #A0129608A  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required		Condition Aggravated	Health Professional Company	Wellbutrin Sr Tablet-Controlled Release	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage	100 MG / PER DAY / ORAL	Eye Disorder Migraine	Representative				

Date:11/28/01ISR Number: 3842959-7Report Type:Periodic Company Report #A0129610A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Blood Pressure Increased	Health Professional Company	Wellbutrin Sr Tablet-Controlled Release	PS		ORAL
	75 MG / ORAL	Urticaria	Representative	Sertraline Hydrochloride	C		

Date:11/28/01ISR Number: 3842960-3Report Type:Periodic Company Report #A0129338A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dystonia	Consumer	Wellbutrin Sr Tablet-Controlled Release	PS		ORAL
	150 MG / TWICE PER DAY	Muscle Tightness					

/ ORAL

Date:11/28/01ISR Number: 3842961-5Report Type:Periodic  
Age:11 YR Gender:Male I/FU:I

Company Report #A0129391A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion	Health Professional Company	Wellbutrin Sr Tablet-Controlled Release	PS		ORAL
150 MG / TWICE PER DAY / ORAL			Representative				

Date:11/28/01ISR Number: 3842962-7Report Type:Periodic  
Age:84 YR Gender:Female I/FU:I

Company Report #A0129304A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abnormal Behaviour Overdose	Health Professional Company	Wellbutrin Sr Tablet-Controlled Release	PS		ORAL
150 MG / SEE DOSAGE TEXT / ORAL			Representative				
		WK		Digoxin Frusemide Warfarin Sodium	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/28/01ISR Number: 3842963-9Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #A0130333A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Drug Interaction	Health	Wellbutrin Sr			
Other		Loss Of Consciousness	Professional	Tablet-Controlled			
ORAL		Petit Mal Epilepsy	Company	Release	PS		ORAL
			Representative	Fluoxetine Hydrochloride (Formation Unknown) (Fluoxetine Hydrochloride)	SS		

Date:11/28/01ISR Number: 3842964-0Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0131024A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Pain Constipation	Consumer	Wellbutrin Sr Tablet-Controlled	PS		ORAL
150 MG / FOUR TIMES PER DAY / ORAL	1	Headache Medication Error Overdose Paranoia Sleep Disorder					

Date:11/28/01ISR Number: 3842965-2Report Type:Periodic  
Age:54 YR Gender:Male I/FU:I

Company Report #A0131053A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cardiac Enzymes Increased Chest Pain Leukocytosis	Consumer	Wellbutrin Sr Tablet-Controlled Release	PS		ORAL
150 MG / TWICE PER DAY		Pruritus					

/ ORAL

Pyrexia  
Rash  
Sensation Of Foreign Body  
Toothache  
Urticaria  
Urticaria Papular

Date:11/28/01ISR Number: 3842966-4Report Type:Periodic Company Report #A0135817A  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Chest Pain	Health Professional	Wellbutrin Sr Tablet-Controlled Release	PS		ORAL
200MG / TWICE							
PER DAY/ ORAL 1 YR							

Date:11/28/01ISR Number: 3842967-6Report Type:Periodic Company Report #A0134688A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Overdose Somnolence	Health Professional	Wellbutrin Sr Tablet-Controlled Arelease (Bupropion Hydrochloride)	PS		ORAL
89 TABLET /							
SINGLE DOSE /							

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Freedom Of Information (FOI) Report

ORAL

Trazodone  
(Formulation  
Unknown) (Trazodone) SS

Date:11/28/01ISR Number: 3843813-7Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #A0134384A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Other	150 MG/TWICE	Convulsion	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
Required	PER DAY/ ORAL 6 MON	Road Traffic Accident					
Intervention to Prevent Permanent Impairment/Damage				Omeprazole Iron Salt	C		

Date:11/28/01ISR Number: 3843814-9Report Type:Periodic  
Age:16 YR Gender:Female I/FU:I

Company Report #A0137064A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300 MG/ TWICE	Grand Mal Convulsion	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
PER DAY/ ORAL				Venlafaxine Hydrochloride	C		
				Citalopram Hydrobromide	C		
				Clonazepam	C		
				Quetiapine Fumarate	C		

Date:11/28/01ISR Number: 3843816-2Report Type:Periodic Company Report #A0137857A  
Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abnormal Behaviour Convulsion Headache Overdose	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
20							
TABLET/ORAL							

Date:11/28/01ISR Number: 3843820-4Report Type:Periodic Company Report #A0137635A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Libido Decreased	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/PER							
DAY/ORAL				Orlistat	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/28/01  
 ISR Number: 3843822-8  
 Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #A0139117A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Anaphylactic Reaction Arthralgia Blood Pressure Abnormal Dizziness	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE PER DAY/ ORAL	Pruritus Urticaria Vomiting					

Date:11/28/01  
 ISR Number: 3843825-3  
 Report Type:Periodic  
 Age:30 YR Gender:Female I/FU:I

Company Report #A0138783A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Accommodation Disorder Amnesia Asthenia Constipation	Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE PER DAY/ORAL	Depression Disturbance In Attention Pain In Extremity Tremor		Topiramate Venlafaxine Hydrochloride Claritin-D Zaleplon	C C C C		

Date:11/28/01  
 ISR Number: 3843827-7  
 Report Type:Periodic  
 Age:9 YR Gender:Male I/FU:I

Company Report #A0138545A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Electroencephalogram Abnormal Epilepsy Fall	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
100 MG/PER						



DAY/ORAL 4 YR  
Hemiplegia  
Hypoaesthesia  
Movement Disorder  
Simple Partial Seizures  
Tremor

Methylphenidate C

Date:11/28/01ISR Number: 3843828-9Report Type:Periodic Company Report #A0138336A  
Age:68 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Anxiety Blood Creatine Phosphokinase Increased Dyspnoea	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE PER DAY/ORAL			Benazepril Rofecoxib	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/28/01ISR Number: 3843831-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0138131A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Cardiac Arrest Convulsion Overdose	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
1000 MG/SINGLE DOSE/ORAL						

Date:11/28/01ISR Number: 3843833-2Report Type:Periodic  
Age:77 YR Gender:Female I/FU:I

Company Report #A0138069A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Disability Required Intervention to 150 MG/TWICE Prevent Permanent PER DAY/ORAL Impairment/Damage	Bradykinesia Convulsion Crying Depression Drug Toxicity Electroencephalogram Abnormal Emotional Disorder Loss Of Consciousness Memory Impairment Mental Impairment	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)  Phenytoin (Formulation Unknown) (Phenytoin) Nitroglycerin Thyroxine Sodium Famotidine Conjugated Estrogens Fluvastatin Sodium Metoprolol Tartrate Spironolactone Verapamil Perdnisone	PS    SS C C C C C C C C C		ORAL

Date:11/28/01ISR Number: 3844209-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0152863A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required		Burning Sensation Drug Eruption Dry Skin Hypersensitivity	Health Professional	Wellbutrin Sr Tablet-Controlled Released (Bupropion Hydrochloride)	PS		ORAL
Intervention to ORAL							
Prevent Permanent Impairment/Damage		Pain Pruritus Rash Maculo-Papular Urticaria					

Date:11/28/01ISR Number: 3844210-0Report Type:Periodic Company Report #A0152864A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required		Burning Sensation Drug Eruption Dry Skin Hypersensitivity	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
Intervention to ORAL							
Prevent Permanent Impairment/Damage		Pain Pruritus Rash Maculo-Papular Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/28/01ISR Number: 3844211-2Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0152865A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged Required	Burning Sensation Drug Eruption Dry Skin	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage	Hypersensitivity  Pain Pruritus Rash Maculo-Papular Urticaria					

Date:11/28/01ISR Number: 3844212-4Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0152866A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged Required	Burning Sensation Drug Eruption Dry Skin	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage	Hypersensitivity  Pain Pruritus Rash Maculo-Papular Urticaria					

Date:11/28/01ISR Number: 3844213-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #A0152867A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged Required	Burning Sensation Drug Eruption Dry Skin	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage	Hypersensitivity  Pain Pruritus Rash Maculo-Papular					

Urticaria

Date:11/28/01ISR Number: 3844214-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0152868A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Burning Sensation	Health	Wellbutrin Sr			
Initial or Prolonged	Drug Eruption	Professional	Tablet-Controlled			
Required	Dry Skin		Release (Bupropion			
Intervention to	Hypersensitivity		Hydrochloride)	PS		ORAL
ORAL						
Prevent Permanent	Pain					
Impairment/Damage	Pruritus					
	Rash Maculo-Papular					
	Urticaria					

Date:11/28/01ISR Number: 3844215-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0152869A

Outcome  
Hospitalization -  
Initial or Prolonged

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Required Intervention to Prevent Permanent Dose Duration Impairment/Damage	PT  Burning Sensation Drug Eruption Dry Skin Hypersensitivity	Report Source  Health Professional	Product  Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	Role   PS	Manufacturer	Route   ORAL
ORAL	Pain Pruritus Rash Maculo-Papular Urticaria					

Date:11/28/01ISR Number: 3844216-1Report Type:Periodic Company Report #A0152870A  
Age: Gender:Male I/FU:I

Outcome Dose Duration Hospitalization - Initial or Prolonged Required Intervention to ORAL Prevent Permanent Impairment/Damage	PT  Burning Sensation Drug Eruption Dry Skin Hypersensitivity	Report Source  Health Professional	Product  Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	Role   PS	Manufacturer	Route   ORAL
	Pain Pruritus Rash Maculo-Papular Urticaria					

Date:11/28/01ISR Number: 3844217-3Report Type:Periodic Company Report #A0152871A  
Age: Gender:Male I/FU:I

Outcome Dose Duration Hospitalization - Initial or Prolonged Required Intervention to ORAL Prevent Permanent Impairment/Damage	PT  Burning Sensation Drug Eruption Dry Skin Hypersensitivity	Report Source  Health Professional	Product  Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	Role   PS	Manufacturer	Route   ORAL
	Pain Pruritus Rash Maculo-Papular Urticaria					

Date:11/28/01ISR Number: 3844218-5Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #A0154808A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Arthralgia	Health	Wellbutrin Sr			
Initial or Prolonged	Coordination Abnormal	Professional	Tablet-Controlled			
Required	Dyskinesia		Release (Bupropion			
Intervention to	Fatigue		Hydrochloride)	PS		ORAL
150 MG PER						
Prevent Permanent	Feeling Hot					
DAY ORAL						
Impairment/Damage	Headache		Vitamin E	C		
	Nervousness		Conjugated Estrogens	C		
	Tardive Dyskinesia		Thyroid	C		
			Ascorbic Acid	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/28/01ISR Number: 3844219-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0155864A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - ORAL Initial or Prolonged	Infectious Mononucleosis  Neutropenia Pyrexia	Health  Professional Company Representative	Wellbutrin	PS		ORAL

Date:11/28/01ISR Number: 3844220-3Report Type:Periodic  
Age:74 YR Gender:Male I/FU:I

Company Report #A0155718A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 100 MG / PER Initial or Prolonged DAY / ORAL Required Intervention to Prevent Permanent Impairment/Damage	Convulsion	Health  Professional  Company Representative	Wellbutrin   Fluoxetine Hydrochloride Metoprolol Succinate Atorvastatin Calcium Isosorbide Dinitrate Ranitidine Hydrochloride Aspirin Theophylline Fluticasone Propionate Combivent Ascorbic Acid Vitamin E	       C C C C  C C C C C C C		ORAL

Date:11/28/01ISR Number: 3844221-5Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #A0155624A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG /	Amnesia	Health	Wellbutrin	PS		ORAL



Initial or Prolonged TWICE PER DAY	Convulsion	Professional			
/ ORAL	Fall	Company			
	Grand Mal Convulsion Irritability	Representative	Levofloxacin	C	

Date:11/28/01ISR Number: 3844222-7Report Type:Periodic Company Report #A0156488A  
 Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300 MG / ORAL Initial or Prolonged	Convulsion	Health	Wellbutrin	PS		ORAL
		Professional Company Representative	Citalopram Hydrobromide	C		

Date:11/28/01ISR Number: 3844223-9Report Type:Periodic Company Report #A0157244A  
 Age:80 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Agitation Aphasia Pyrexia	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL			Venlafaxine			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride  
(Formulation  
Unknown)  
(Venlafaxine SS

Date:11/28/01ISR Number: 3844224-0Report Type:Periodic Company Report #A0157023A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG / PER	Convulsion	Health	Wellbutrin	PS		
Hospitalization - DAY / Initial or Prolonged			Professional  Company Representative				

Date:11/28/01ISR Number: 3844225-2Report Type:Periodic Company Report #A0156900A  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	200 MG / PER	Hallucination, Auditory Hallucination, Visual Infection	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
DAY / ORAL				Lithium Salt Haloperidol Nortriptyline Hcl	C C C		

Date:11/28/01ISR Number: 3844226-4Report Type:Periodic Company Report #A0157660A  
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL Initial or Prolonged Required		Arthralgia  Red Blood Cell Sedimentation Rate	Company Representative	Wellbutrin  Atorvastatin Calcium	PS  C		ORAL

Intervention to            Increased  
Prevent Permanent        Serum Sickness  
Impairment/Damage        Urticaria

Date:11/28/01ISR Number: 3844227-6Report Type:Periodic            Company Report #A0159578A  
Age:21 YR    Gender:Female        I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose            Duration Hospitalization - TWICE PER DAY Initial or Prolonged / ORAL	Convulsion	Health  Professional	Wellbutrin	PS		ORAL
		Company Representative	Claritin-D Hydroxyzine Inhaler Oral Contraceptive	C C C C		

Date:11/28/01ISR Number: 3844228-8Report Type:Periodic            Company Report #A0159581A  
Age:3 YR    Gender:Female        I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose            Duration Hospitalization - Initial or Prolonged  100 MG / ORAL	Accidental Exposure Hypoglycaemia Vomiting	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/28/01ISR Number: 3844332-4Report Type:Periodic  
Age:58 YR Gender:Male I/FU:I

Company Report #A0159463A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Agoraphobia Amnesia Back Pain Compression Fracture Convulsion Night Sweats	Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
200 MG / TWICE PER DAY/ ORAL	2	YR		Lovastatin Actifed Aspirin	C C C		

Date:11/28/01ISR Number: 3844333-6Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0160489A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other		Deafness Dizziness Ear Disorder Nausea Tinnitus	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER DAY / ORAL				Pravastatin Sodium Aspirin Fexofenadine Hydrochlorid Metoprolol	C C C C		

Date:11/28/01ISR Number: 3844334-8Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0160503A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Crying Depression	Consumer	Wellbutrin Sr Tablet-Controlled			

150 MG /	Feeling Abnormal	Release (Bupropion		
	Hypoaesthesia	Hydrochloride)	PS	ORAL
TWICE PER DAY	Hypoaesthesia Oral			
/ ORAL	Sedation			
	Suicide Attempt	Paracetamol		
		(Formulation		
		Unknown)		
		(Acetaminophen)	SS	
		Sertraline		
		Hydrochloride		
		(Formulation		
		Unknown) (Sertraline		
		Hydrochloride)	SS	
100 MG / PER				
DAY		Methylphenidate Hcl		
		(Formulation		
		Unknown)		
		(Methylphenidate		
20 MG		Hcl)	SS	
		Disulfiram		
		(Formulation		
		Unknown)		
250 MG / PER		(Disulfiram)	SS	
DAY				
		Lorazepam		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Formulation  
Unknown) (Lorazepam) SS

ORAL

5 MG / AS

REQUIRED /

ORAL

Date:11/28/01ISR Number: 3844335-XReport Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0161251A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Atrioventricular Block Bradycardia	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

150 MG / PER

DAY / ORAL

Venlafaxine  
Hydrochloride C

Date:11/28/01ISR Number: 3844336-1Report Type:Periodic  
Age:22 YR Gender:Female I/FU:I

Company Report #A0161748A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged		Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

150 MG /

TWICE PER DAY

/ ORAL

Date:11/28/01ISR Number: 3844337-3Report Type:Periodic  
Age:36 YR Gender:Male I/FU:F

Company Report #A0118454A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anxiety Asthenia Blood Pressure Increased Depersonalisation	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER DAY / ORAL	4	WK Dissociation Disturbance In Attention Dizziness Drug Withdrawal Syndrome Palpitations Panic Attack Simple Partial Seizures Tachycardia		Citalopram Hydrobromide (Formulation Unknown) (Citalopram Hydrobromide) Buspirone Hydrochloride	SS C		

Date:11/28/01ISR Number: 3844338-5Report Type:Periodic Company Report #A0124798A  
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthritis Pyrexia Serum Sickness Urticaria	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / TWICE PER DAY / ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/28/01ISR Number: 3844339-7Report Type:Periodic  
Age:35 YR Gender:Male I/FU:I

Company Report #A0139382A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Abdominal Discomfort Abdominal Pain Confusional State Convulsion		Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/SEE	Fall					
DOSAGE	Nausea					
TEXT/ORAL	Overdose					

Date:11/28/01ISR Number: 3844340-3Report Type:Periodic  
Age:17 YR Gender:Female I/FU:I

Company Report #A0139566A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Other Required Intervention to 150 MG/TWICE Prevent Permanent PER DAY/ORAL Impairment/Damage	Convulsion	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

Date:11/28/01ISR Number: 3844342-7Report Type:Periodic  
Age:10 YR Gender:Female I/FU:I

Company Report #A0141090A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Other	Convulsion Overdose	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
SINGLE DOSE /						
ORAL						



Date:11/28/01ISR Number: 3844350-6Report Type:Periodic  
Age:19 YR Gender:Male I/FU:I

Company Report #A0140813A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Confusional State	Health	Wellbutrin Sr			
Other		Dizziness	Professional	Tablet-Controlled			
		Fall	Company	Release (Bupropion			
		Grand Mal Convulsion	Representative	Hydrochloride)	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL		Headache					
		Hyperhidrosis		Biaxin	C		
		Laceration		Terbinafine	C		
		Loss Of Consciousness					
		Pallor					
		Postictal State					
		Vision Blurred					
		Vomiting					
		Weight Increased					

Date:11/28/01ISR Number: 3844352-XReport Type:Periodic  
Age:59 YR Gender:Male I/FU:I

Company Report #A0140570A

Outcome	PT
Hospitalization -	Dizziness
Initial or Prolonged	Hypoglycaemia
	Loss Of Consciousness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Syncope

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
100 MG/TWICE PER DAY/ORAL		Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
			Insulin	C		

Date:11/28/01ISR Number: 3844353-1Report Type:Periodic Company Report #A0139845A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Urinary Retention	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE PER DAY/ ORAL							

Date:11/28/01ISR Number: 3844354-3Report Type:Periodic Company Report #A0141485A  
Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthralgia Burning Sensation Dyspnoea Face Oedema	Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
100 MG/TWICE PER DAY/ORAL		Fear  Hypersensitivity Oedema Peripheral Pain Pain In Extremity Pharyngeal Oedema					

Pruritus  
Rash  
Swelling  
Tenderness  
Unemployment  
Urticaria

Date:11/28/01ISR Number: 3844355-5Report Type:Periodic  
Age:51 YR Gender:Male I/FU:I

Company Report #A0141554A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Grand Mal Convulsion Loss Of Consciousness Mouth Haemorrhage Tongue Biting	Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE  PER DAY/ORAL			Centrum Silver Iron Salt Ibuprofen Lansoprazole Alprazolam	C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/28/01ISR Number: 3844356-7Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #A0141899A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/TWICE PER DAY/ORAL	Grand Mal Convulsion Petit Mal Epilepsy Tremor Yawning	Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
			Vicodin	C		
			Prednisone	C		
			Co-Trimoxazole	C		
			Fluconazole	C		
			Nystatin	C		
			Medroxyprogesterone Ace.	C		

Date:11/28/01ISR Number: 3844357-9Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0142036A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Required Intervention to Prevent Permanent 150 MG/PER Impairment/Damage DAY/ORAL	Tremor	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

Date:11/28/01ISR Number: 3844358-0Report Type:Periodic  
Age:64 YR Gender:Male I/FU:I

Company Report #A0142590A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 100 MG / ORAL	Grand Mal Convulsion Tremor	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

ORAL

Wellbutrin Tablet (Bupropion Hydrochloride)	SS
Citalopram Hydrobromide	C
Olanzapine	C
Paroxetine Hydrochloride	C
Venlafaxine Hydrochloride	C
Verapamil Hydrochloride	C
Metoprolol Tartrate	C
Warfarin Sodium	C
Atorvastatin Calcium	C
Tylenol No. 3	C
Oxycodone Hydrochloride	C
Fiorinal	C
Tylenol W/ Codeine No. 4	C

ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/28/01ISR Number: 3844359-2Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0143252A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Cardiac Arrest	Health	Wellbutrin Sr			
Initial or Prolonged	Myocardial Infarction	Professional	Tablet-Controlled			
		Company	Release (Bupropion			
		Representative	Hydrochloride)	PS		ORAL
ORAL						

Date:11/28/01ISR Number: 3844360-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0143904A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Convulsion	Health	Wellbutrin Sr			
Initial or Prolonged		Professional	Tablet-Controlled			
Other		Company	Release (Bupropion			
		Representative	Hydrochloride)	PS		ORAL
ORAL						

Date:11/28/01ISR Number: 3844361-0Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #A0144030A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Convulsion	Health	Wellbutrin Sr			
Initial or Prolonged		Professional	Tablet-Controlled			
		Company	Release (Bupropion			
		Representative	Hydrochloride)	PS		ORAL
150 MG /						

TWICE PER DAY

/ ORAL

Date:11/28/01ISR Number: 3844362-2Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #A0143948A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion Electroencephalogram Abnormal	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL	DAY						

Date:11/28/01ISR Number: 3844363-4Report Type:Periodic Company Report #A0145187A  
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Disability		Accommodation Disorder Amnesia Confusional State Headache	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Date:11/28/01ISR Number: 3844364-6Report Type:Periodic Company Report #A0144973A  
 Age:35 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Anxiety Cold Sweat Coordination Abnormal Drug Ineffective Screaming Speech Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Weight Increased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
100 MG / TWICE PER DAY / ORAL		Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
12.5 MG / PER DAY / ORAL			Lamictal Tablet (Lamotrigine)	SS		ORAL
20 MG / PER DAY			Fluoxetine Hydrochloride (Formulation Unknown) (Fluoxetine Hydrochloride)	SS		
			Gabapentin	C		
			Quetiapine Fumarate	C		

Date:11/28/01  
Age:42 YR  
Gender:Male  
ISR Number: 3844365-8  
Report Type:Periodic  
I/FU:I

Company Report #A0144927A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anxiety Feeling Abnormal Lethargy	Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
100 MG / PER DAY / ORAL				Paroxetine Hydrochloride (Formulation Unknown) (Paroxetine			



Date:11/28/01ISR Number: 3844366-XReport Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0144931A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Stevens-Johnson Syndrome	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL						

Date:11/28/01ISR Number: 3844367-1Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #A0144731A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Required Intervention to 150 MG / Prevent Permanent TWICE PER DAY Impairment/Damage / ORAL	Erythema Pruritus Rash Respiratory Distress Urticaria	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/28/01ISR Number: 3844368-3Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #A0145663A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Duration Intentional Misuse Lethargy Nausea	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

SEE DOSAGE

TEXT / ORAL

Date:11/28/01ISR Number: 3844369-5Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #A0146394A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Required Intervention to 150 MG/ TWICE Prevent Permanent PER DAY/ ORAL Impairment/Damage	Duration Euphoric Mood Fall Fatigue Grand Mal Convulsion Libido Decreased Tooth Loss	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
			Paroxetine Hydrochloride Tablet (Paroxetine Hydrochloride)	SS		ORAL

20 MG/ PER

DAY/ ORAL

ORAL

Date:11/28/01ISR Number: 3844370-1Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #A0145969A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Duration Drug Effect Decreased Stress	Health Professional	Wellbutrin Sr Tablet-Controlled			

Required Intervention to 150 MG/ TWICE Prevent Permanent PER DAY/ ORAL Impairment/Damage	Urticaria		Release (Bupropion Hydrochloride)	PS		ORAL
			Methylprednisolone (Formulation Unknown) (Methylprednisolone)	SS		ORAL
ORAL						

Date:11/28/01ISR Number: 3844371-3Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #A0145970A

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required	Headache Pruritus Stress	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
Intervention to 150 MG/ TWICE Prevent Permanent PER DAY/ ORAL Impairment/Damage	Urticaria		Methylprednisolone (Formulation Unknown) (Methylprednisolone)	SS		ORAL
			Me-Prednisolone Na Succ. (Formulation Unknown) (Me-Prednisolone Na Succ.)	SS		
INTRAVENOUS	125 MG/ FOUR					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

TIMES PER

DAY/ INTRAV

ORAL				Prednisone (Formulation Unknown) (Prednisone)	SS		ORAL
				Ortho Tri-Cyclen	C		

Date:11/28/01ISR Number: 3844372-5Report Type:Periodic Company Report #A0147190A  
Age:21 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Colitis Gastrointestinal Infection Haematochezia	Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
400 MG/ THREE	Stomach Discomfort					

TIMES PER

DAY/ ORAL

Date:11/28/01ISR Number: 3844373-7Report Type:Periodic Company Report #A0147579A  
Age:83 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Hyponatraemia Inappropriate Antidiuretic Hormone Secretion	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE	Urinary Tract Infection					

PER DAY/ ORAL

Date:11/28/01ISR Number: 3844374-9Report Type:Periodic Company Report #A0150071A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required		Burning Sensation Drug Eruption Dry Skin Hypersensitivity	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
Intervention to ORAL							
Prevent Permanent Impairment/Damage		Pain Pruritus Rash Maculo-Papular Skin Ulcer Urticaria					

Date:11/28/01ISR Number: 3844375-0Report Type:Periodic Company Report #A0150103A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE							
PER DAY/ ORAL	4	YR					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/28/01ISR Number: 3844376-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0152014A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Hypersensitivity	Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL						

Date:11/28/01ISR Number: 3844377-4Report Type:Periodic  
Age:20 YR Gender:Female I/FU:I

Company Report #A0153398A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Convulsion Psychotic Disorder	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL 2 YR			Paroxetine Hydrochloride	C		

Date:11/29/01ISR Number: 3831200-7Report Type:Expedited (15-DaCompany Report #A0159004A  
Age:57 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice Initial or Prolonged per day	Bradycardia Dehydration Syncope		Wellbutrin	PS	Glaxo Wellcome	ORAL
81MG Per day			Aspirin	C		ORAL
			Glucophage	C		
			Nitroglycerin	C	Glaxo Wellcome	
			Altace	C		
			Zocor	C		
			Niaspan	C		

Date:11/29/01ISR Number: 3831203-2Report Type:Expedited (15-DaCompany Report #A0168050A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
400MG per day	3	YR	Drug Effect Decreased	Wellbutrin	PS	Glaxo Wellcome	ORAL
			Insomnia	Pepcid Ac	C		
			Medication Residue	Albuterol	C	Glaxo Wellcome	
				Combivent	C		
				Flovent	C	Glaxo Wellcome	

Date:11/29/01ISR Number: 3831210-XReport Type:Expedited (15-DaCompany Report #B0127463A  
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
300MG Per day	42	DAY	Abdominal Pain	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged			Drug Effect Decreased	Climaston	C		
			Malaise				
			Stress				
			Syncope Vasovagal				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/29/01ISR Number: 3831211-1Report Type:Expedited (15-DaCompany Report #B0127467A

Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxo Wellcome	ORAL
300MG Twice							
per day	26	DAY					
		Depression					
		Mood Swings		Salbutamol	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)							
				Beclomethasone	C	Glaxo Wellcome	
UNKNOWN							

Date:11/29/01ISR Number: 3832493-2Report Type:Expedited (15-DaCompany Report #HQ8542216NOV2001

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Anxiety	Consumer	Effexor (Venlafaxine Hydrochloride, Tablet)	PS		
		Injury		Ativan (Lorazepam)	SS		
				Buspar (Buspirone Hydrochloride)	SS		
				Inderal (Propranolol Hydrochloride)	SS		
				Paxil (Paroxetine Hydrochloride)	SS		
				Prozac (Fluoxetine Hydrochloride)	SS		
				Wellbutrin (Amfebutamone Hydrochloride)	SS		

Date:11/30/01ISR Number: 3831960-5Report Type:Expedited (15-DaCompany Report #B0102458A

Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Death  
 150MG Per day 41 DAY  
 Bronchitis Chronic  
 Zyban  
 PS Glaxo Wellcome ORAL

Date:11/30/01ISR Number: 3831961-7Report Type:Expedited (15-DaCompany Report #B0117985A  
 Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	8 WK	Bronchospasm		Zyban	PS	Glaxo Wellcome	
		Dizziness					
		Dyspnoea					
		Pain					

Date:11/30/01ISR Number: 3831962-9Report Type:Expedited (15-DaCompany Report #B0120692A  
 Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day 41 DAY	Bronchitis Chronic		Zyban	PS	Glaxo Wellcome	ORAL
		Circulatory Collapse					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/01ISR Number: 3831963-0Report Type:Expedited (15-DaCompany Report #B0120884A  
 Age:73 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aortic Aneurysm		Zyban	PS	Glaxo Wellcome	
150MG As		Blister					
directed	9 DAY	Dizziness					
		Insomnia					
		Tremor					

Date:11/30/01ISR Number: 3831970-8Report Type:Expedited (15-DaCompany Report #B0127466A  
 Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Angioneurotic Oedema		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day	20 DAY	Joint Swelling					
Initial or Prolonged		Lung Disorder					
		Oedema Peripheral					
		Urticaria					

Date:11/30/01ISR Number: 3833720-8Report Type:Direct Company Report #  
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Pruritus		Wellbutrin Sr 1150			
ONCE QD		Rash		Mg	PS		
		Urticaria		Vitamins	C		

Date:11/30/01ISR Number: 3833847-0Report Type:Direct Company Report #  
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Hypoaesthesia		Wellbutrin Sr 150mg			

Intervention to	Rash	Initially	PS
1 BID			
Prevent Permanent	Urticaria	Wellbutrin Sr 100mg	
Impairment/Damage		(Dose Increased)	SS
2 BID		Celexa	C

Date:12/03/01ISR Number: 3832651-7Report Type:Expedited (15-DaCompany Report #A0093320A  
Age:54 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Agitation		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice						
Initial or Prolonged	Asthenia					
per day						
307 DAY						
	Carpal Tunnel Syndrome		Azithromycin	C		
	Crying		Zantac	C	Glaxo Wellcome	
	Drug Ineffective		Nasonex	C		
	Emotional Distress					
	Eye Disorder					
	Facial Palsy					
	Hypoaesthesia					
	Insomnia					
	Irritability					
	Nervousness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/03/01ISR Number: 3832653-0Report Type:Expedited (15-DaCompany Report #A0154004A

Age:26 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angiopathy		Zyban	PS	Glaxo Wellcome	ORAL
150MG See		Blister					
dosage text	3	DAY					
		Burning Sensation		No Concurrent			
		Dermatitis Bullous		Medication	C		
		Erythema					
		Hypersensitivity					
		Insomnia					
		Pain In Extremity					
		Penile Pain					
		Penis Disorder					
		Pruritus					
		Scab					
		Scar					
		Skin Discolouration					
		Swelling					
		Ulcer					

Date:12/03/01ISR Number: 3832654-2Report Type:Expedited (15-DaCompany Report #A0161553A

Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	2	DAY					
		Hemiparesis					
		Migraine					

Date:12/03/01ISR Number: 3832658-XReport Type:Expedited (15-DaCompany Report #B0100684A

Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardiac Arrest		Zyban	PS	Glaxo Wellcome	
				Lovan	C		
				Alcohol	C		

Date:12/03/01ISR Number: 3832669-4Report Type:Expedited (15-DaCompany Report #D0023165A  
Age:30 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Psychotic Disorder		Zyban	PS	Glaxo Wellcome	ORAL

Date:12/03/01ISR Number: 3833744-0Report Type:Direct Company Report #  
Age:29 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening	Convulsion Loss Of Consciousness		Wellbutrin Sr 150 Mg Glaxo Smith Kline	PS	Glaxo Smith Kline	ORAL

150 MG TWICE

DAILY ORAL

Bupropion 150mg  
Lemmon

150MG TWICE

DAILY ORAL

19-Aug-2005 12:44 PM

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/04/01ISR Number: 3833080-2Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Wellbutrin Sr 150 Mg	PS		
150 MG BID							
		Dyspnoea		Ziprasidone (Geodon)	SS		
150 MG BID							
SEE IMAGE				Bupropion	SS		
				Albuterol Inhaler	C		
				Guanfacine	C		

Date:12/04/01ISR Number: 3834052-4Report Type:Direct  
 Age:11 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Arthralgia		Wellbutrin Sr			
Initial or Prolonged		Dermatitis		(100mg)	PS		
100MG ONE IN							
		Difficulty In Walking					
ADAY							
		Oedema					

Date:12/05/01ISR Number: 3834165-7Report Type:Expedited (15-DaCompany Report #A0167282A  
 Age:40 YR Gender:Male I/FU:F

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day 8 DAY							
Hospitalization -		Conjunctival Hyperaemia		No Concurrent			
Initial or Prolonged		Dry Mouth		Medications	C		
Disability		Dyspnoea					
		Neck Pain					
		Palpitations					
		Pharyngolaryngeal Pain					
		Pulmonary Embolism					
		Rash Erythematous					

Date:12/05/01ISR Number: 3834178-5Report Type:Expedited (15-DaCompany Report #B0127547A  
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		C-Reactive Protein  Increased  Pain Phlebitis Thrombophlebitis Superficial Varicose Vein		Zyban	PS	Glaxo Wellcome	

Date:12/05/01ISR Number: 3834183-9Report Type:Expedited (15-DaCompany Report #B0127963A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebrovascular Accident Hypertension		Zyban	PS	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/05/01ISR Number: 3834184-0Report Type:Expedited (15-DaCompany Report #B0127964A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Pain Agitation Confusional State Fatigue Hyperhidrosis Lymphadenopathy Pyrexia		Zyban	PS	Glaxo Wellcome	ORAL

Date:12/05/01ISR Number: 3834185-2Report Type:Expedited (15-DaCompany Report #B0127965A

Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day 6 DAY	Cardiac Arrest		Zyban	PS	Glaxo Wellcome	
	20MG per day	Cardiomyopathy		Enalapril	C		
	40MG per day			Omeprazole	C		
	20MG per day			Acenocoumarol	C		
	25MG per day			Simvastatin	C		
				Carvedilol	C	Glaxo Wellcome	

Date:12/05/01ISR Number: 3834186-4Report Type:Expedited (15-DaCompany Report #B0128139A

Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice per day	Haemorrhagic Stroke		Zyban	PS	Glaxo Wellcome	ORAL
	71 DAY	Renal Impairment		Atenolol	C		ORAL
	50MG per day			Felodipine	C		ORAL
	10MG per day						



2.5MG per day		Bendrofluazide	C	Glaxo Wellcome	ORAL
		Tibolone	C		ORAL
2.5MG per day					

Date:12/05/01ISR Number: 3834188-8Report Type:Expedited (15-DaCompany Report #B0128148A  
 Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthma		Zyban	PS	Glaxo Wellcome	ORAL
4 DAY				Salmeterol	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION) 1PUFF Twice							
per day							

Date:12/05/01ISR Number: 3834189-XReport Type:Expedited (15-DaCompany Report #B0128149A  
 Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Bronchospasm		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Initial or Prolonged		Localised Infection					
per day	78 DAY						
		Oedema					
		Proteinuria					
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/05/01ISR Number: 3834190-6Report Type:Expedited (15-DaCompany Report #B0128151A  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Hepatic Failure		Zyntabac	PS	Glaxo Wellcome	ORAL
150MG Per day							
Hospitalization -		Renal Failure		Sertraline	C		
Initial or Prolonged		Respiratory Failure		Alprazolam	C		
Disability							
Other							

Date:12/05/01ISR Number: 3834191-8Report Type:Expedited (15-DaCompany Report #B0128157A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Depression		Zyban	PS	Glaxo Wellcome	
150MG Twice							
per day	18 DAY	Enzyme Abnormality					
		Obsessive-Compulsive Disorder		Clomipramide	C		
		Suicidal Ideation		Alprazolam	C		

Date:12/05/01ISR Number: 3834193-1Report Type:Expedited (15-DaCompany Report #B0128228A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Csf Protein Increased		Zyban	PS	Glaxo Wellcome	ORAL
10 DAY							
		Loss Of Consciousness					
		Muscle Twitching					
		Toxicologic Test Abnormal					

Date:12/05/01ISR Number: 3834195-5Report Type:Expedited (15-DaCompany Report #D0023445A  
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 150MG In the Initial or Prolonged morning	2 WK	Blood Creatine Phosphokinase Increased Psychotic Disorder	Zyban  Pantozol Leponex  Tenormin  Rivotril	PS  C C C C	Glaxo Wellcome	ORAL  ORAL ORAL ORAL ORAL
40MG Per day 50MG Twice per day 25MG Twice per day 500MCG As required						

Date:12/05/01ISR Number: 3836267-8Report Type:Direct Company Report #  
Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin (100mg)	PS		ORAL
100MG 1 BID				Ortho-Cyclen	C		
ORAL							

Date:12/06/01ISR Number: 3834767-8Report Type:Expedited (15-DaCompany Report #A0168510A  
Age:58 YR Gender:Male I/FU:F

Outcome	PT
Other	Depression Dysphonia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	56 DAY	Mood Swings Psychiatric Symptom Psychomotor Hyperactivity Suicidal Ideation Thinking Abnormal Tremor		Wellbutrin Sr  Neurontin Anti-Hypertensive Unspecified Medications	PS  C C C	Glaxo Wellcome	ORAL

Date:12/06/01ISR Number: 3834776-9Report Type:Expedited (15-DaCompany Report #B0098813A  
Age:57 YR Gender:Female I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	16 DAY	Antimitochondrial Antibody Positive		Zyban	PS	Glaxo Wellcome	ORAL
20MG per day		Cutaneous Vasculitis		Lisinopril	C		
1TAB per day		Dermatitis Exfoliative		Dytenzide	C	Glaxo Wellcome	
		Leukocytosis Rash Erythematous Rash Maculo-Papular Rash Pruritic					

Date:12/06/01ISR Number: 3834787-3Report Type:Expedited (15-DaCompany Report #B0128344A  
Age:69 YR Gender:Male I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Life-Threatening Hospitalization - Initial or Prolonged		Zyban	PS	Glaxo Wellcome	ORAL
2.5MG Per day				Bendrofluazide	C	Glaxo Wellcome	ORAL

Disability 20MG Twice			Nifedipine	C	Glaxo Wellcome	ORAL
per day						
10MG Per day			Atenolol	C		ORAL
20MG Twice			Nicorandil	C		ORAL
per day						
RESPIRATORY (INHALATION)			Becotide	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)			Ventolin	C	Glaxo Wellcome	

Date:12/07/01ISR Number: 3835595-XReport Type:Expedited (15-DaCompany Report #A0169154A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Epistaxis Retinal Detachment		Zyban	PS	Glaxo Wellcome	ORAL

Date:12/07/01ISR Number: 3835608-5Report Type:Expedited (15-DaCompany Report #B0128267A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 6000MG Per Initial or Prolonged day 1 DAY		Agitation Mydriasis Suicide Attempt Tachycardia Tremor		Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/07/01ISR Number: 3836551-8Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG 1X		Confusional State Crying Dysphemia Fear Feeling Abnormal		Zyban (150 Mg) Catalytica Pharmaceuticals 1 1x Oral	PS	Catalytica Pharmaceuticals	ORAL
ORAL		Hallucination  Headache Hyperacusis Nausea Paranoia Suicidal Ideation Tinnitus Vision Blurred					

Date:12/07/01ISR Number: 3837149-8Report Type:Expedited (15-DaCompany Report #2001082086US  
 Age:45 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
INTRA VENOUS	600 MG, BID,	Blood Pressure Decreased Delirium Delusion	Health Professional	Linezolid(Linezolid) Solution, Sterile	PS		
IV		Drug Interaction Dry Mouth Haematocrit Decreased		Flagyl (Metronidazole) Tablet	SS		ORAL
500 MG, QID,		Haemoglobin Decreased					
ORAL		Hallucination, Visual Mydriasis		Sertraline(Sertraline) e)	SS		
200 MG, QD,		Respiratory Rate Increased		Risperidone(Risperidone) one)	SS		
1 MG, BID,		Serotonin Syndrome		Bupropion			

75 MG, BID	Vomiting	(Amfebutamone)	SS
	White Blood Cell Count Increased	Trazodone	C
		Litiumkarbonat (Lithium Carbonate)	C
		Baclofen	C
		Promethazine(Promethazine)	C
		Docusate	
		Sodium(Docusate Sodium)	C
		Bisacodyl(Bisacodyl)	C
		Megestrol(Megestrol)	C
		Lansoprazole	C

Date:12/10/01ISR Number: 3838049-XReport Type:Direct Company Report #  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Zyban 150mg Sr	PS		ORAL
150MG 1 TAB							
PO QD X3 D							
THEN BID							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/11/01ISR Number: 3836830-4Report Type:Expedited (15-DaCompany Report #A0163666A  
 Age:33 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Three Initial or Prolonged times per day	Depression		Wellbutrin	PS	Glaxo Wellcome	ORAL
Other 225MG Twice per day	Grand Mal Convulsion		Lamictal	C	Glaxo Wellcome	ORAL
35MG per day			Zyprexa	C		ORAL
			Luvox	C		ORAL
			Remeron	C		
			Lithium	C		
300MG Twice per day			Risperdal	C		ORAL
8MG Per day			Antabuse	C		ORAL

Date:12/11/01ISR Number: 3836835-3Report Type:Expedited (15-DaCompany Report #B0123297A  
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG As directed	Arrhythmia		Zyban	PS	Glaxo Wellcome	
	Circulatory Collapse					

Date:12/11/01ISR Number: 3836842-0Report Type:Expedited (15-DaCompany Report #B0128343A  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 7 DAY	Asthma		Zyban	PS	Glaxo Wellcome	ORAL



RESPIRATORY ( INHALATION)	Salbutamol	C	Glaxo Wellcome	
RESPIRATORY ( INHALATION)	Salmeterol	C	Glaxo Wellcome	
RESPIRATORY ( INHALATION)	Beclomethasone	C	Glaxo Wellcome	
	Co-Proxamol	C		ORAL
	Arthrotec	C		

Date:12/11/01ISR Number: 3836843-2Report Type:Expedited (15-DaCompany Report #B0128345A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia		Zyban	PS	Glaxo Wellcome	ORAL
4 DAY							
1UNIT Twice		Dysarthria		Mometasone	C		NASAL
per day							

Date:12/11/01ISR Number: 3836844-4Report Type:Expedited (15-DaCompany Report #B0128468A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Aggression		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day 13 DAY							
Initial or Prolonged		Attention Deficit/Hyperactivity Disorder Meningitis					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/11/01ISR Number: 3836845-6Report Type:Expedited (15-DaCompany Report #B0128545A  
 Age:39 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Confusional State		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -	Haemolytic Anaemia					
Initial or Prolonged	Jaundice					
	Psoriasis					
	Pyrexia					
	Renal Impairment					
	Sepsis					
	Thrombotic					
	Thrombocytopenic Purpura					

Date:12/11/01ISR Number: 3836846-8Report Type:Expedited (15-DaCompany Report #B0128645A  
 Age:61 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Diabetes Mellitus		Zyban	PS	Glaxo Wellcome	
150MG Per day						
Initial or Prolonged			Albyl-E	SS		ORAL
160MG Per day						
			Selozok	SS		ORAL
100MG Per day						

Date:12/11/01ISR Number: 3836847-XReport Type:Expedited (15-DaCompany Report #B0128651A  
 Age:57 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Death	Aortic Valve Stenosis		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day						
8 DAY	Brain Oedema		Adepal	C		ORAL
	Cardiac Failure		Lisinopril	C		ORAL
	Haemorrhagic Stroke					
	Headache					
	Hypertensive Crisis					
	Intracranial Aneurysm					
	Mydriasis					
	Neck Pain					

Otitis Externa  
Palpitations  
Ventricular Extrasystoles

Date:12/12/01ISR Number: 3837381-3Report Type:Expedited (15-DaCompany Report #A0169561A  
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Visual Acuity Reduced		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day							

Date:12/12/01ISR Number: 3837382-5Report Type:Expedited (15-DaCompany Report #A0169565A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Abortion Induced		Wellbutrin	PS	Glaxo Wellcome	ORAL
200MG Per day							
		Complications Of Maternal Exposure To Therapeutic Drugs Trisomy 21					

Freedom Of Information (FOI) Report

Date:12/12/01ISR Number: 3837383-7Report Type:Expedited (15-DaCompany Report #B0114598A

Age:56 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150MG Twice	Abdominal Pain		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - per day 62 DAY	Aggression					
Initial or Prolonged	Amnesia					
Other	Anaphylactic Shock					
	Anxiety					
	Balance Disorder					
	Blister					
	Blood Bilirubin Increased					
	Blood Pressure Increased					
	Bone Density Decreased					
	Chest Pain					
	Confusional State					
	Convulsion					
	Dermatitis					
	Diarrhoea					
	Disorientation					
	Disturbance In Attention					
	Dizziness					
	Dry Skin					
	Dysphagia					
	Dyspnoea					
	Eyelid Oedema					
	Face Oedema					
	Feeling Abnormal					
	Feeling Drunk					
	Feeling Hot					
	Flatulence					
	Gastroenteritis					
	Hyperhidrosis					
	Irritability					
	Irritable Bowel Syndrome					
	Loss Of Consciousness					
	Malaise					
	Memory Impairment					
	Nausea					
	Ocular Hyperaemia					
	Oedema Peripheral					
	Pain					
	Pharyngeal Oedema					
	Pruritus					

Rhinitis  
Speech Disorder  
Tinnitus  
Tongue Oedema  
Urticaria

Date:12/12/01ISR Number: 3837395-3Report Type:Expedited (15-DaCompany Report #B0128808A  
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Burning Sensation		Bupropion			
		Dermatitis Exfoliative		Hydrochloride	PS	Glaxo Wellcome	
UNKNOWN		1 WK					
		Erythema					
		Psoriasis					
		Rash Generalised					
		Throat Irritation					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/12/01  
 Age: 12/12/01  
 Gender:Female  
 I/FU:I

Report Type:Direct  
 Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Intolerance		Bupropion	PS		ORAL
100MG 1 BID		Tongue Oedema					
PO							

Date:12/12/01  
 Age: 12/12/01  
 Gender:Male  
 I/FU:I

Report Type:Expedited (15-Da  
 Company Report #PHBS2001US12331

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest Neonatal Coma Cyanosis Neonatal	Literature Health Professional	Ritaline (Methylphenidate Hydrochloride)	PS		
TRANSPLACENTAL	TRANSPLACENTA	Hyporeflexia					
L		Hypotonia Neonatal Maternal Drugs Affecting Foetus		Wellbutrin (Amfebutamone Hydrochloride)	SS		
TRANSPLACENTAL	TRANSPLACENTA	Neonatal Apnoeic Attack					
L		Oliguria Tremor Neonatal		Benadryl (Diphenhydramine Hydrochloride)	SS		
TRANSPLACENTAL	TRANSPLACENTA						
L							

Date:12/13/01  
 Age:56 YR  
 Gender:Male  
 I/FU:F

Report Type:Expedited (15-Da  
 Company Report #B0105221A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Other		Abdominal Pain Amnesia Anaphylactic Shock Anxiety		Zyban	PS	Glaxo Wellcome	ORAL

Diarrhoea  
Disturbance In Attention  
Irritability  
Pruritus  
Rash Erythematous

Date:12/13/01ISR Number: 3838227-XReport Type:Expedited (15-DaCompany Report #B0128151A  
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death							
150MG Per day							
Hospitalization -							
Initial or Prolonged							
Disability							
Other							

Date:12/13/01ISR Number: 3838233-5Report Type:Expedited (15-DaCompany Report #B0128893A  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -							
150MG Per day 52 DAY							
Initial or Prolonged							
Disability							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/14/01ISR Number: 3839172-6Report Type:Expedited (15-DaCompany Report #A0169154A  
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Epistaxis Retinal Detachment Tension Headache		Zyban	PS	Glaxo Wellcome	ORAL

Date:12/14/01ISR Number: 3839177-5Report Type:Expedited (15-DaCompany Report #B0096692A  
Age:73 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG Per day Initial or Prolonged		Pyrexia Vasculitis		Zyban Magnyl	PS C	Glaxo Wellcome	ORAL

Date:12/14/01ISR Number: 3839181-7Report Type:Expedited (15-DaCompany Report #B0126564A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Congenital Anomaly UNKNOWN	300MG per day	Asthma Caesarean Section Complications Of Maternal Exposure To Therapeutic Drugs Congenital Anomaly Dermatitis Atopic Neonatal Disorder Pregnancy Ventricular Septal Defect Acquired		Zyban	PS	Glaxo Wellcome	

Date:12/14/01ISR Number: 3839182-9Report Type:Expedited (15-DaCompany Report #B0128084A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Disability                      Dizziness                      Bupropion  
                                    Nausea                                      Hydrochloride                      PS                      Glaxo Wellcome                      ORAL  
150MG Per day    6                      DAY  
                                    Vision Blurred

Date:12/17/01ISR Number: 3839283-5Report Type:Expedited (15-DaCompany Report #A0165224A  
Age:                      Gender:Male                      I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Acquired		Wellbutrin	PS	Glaxo Wellcome	ORAL
200MG Twice		Tracheo-Oesophageal					
per day	316	DAY					
		Fistula		Buspar	C		ORAL
15MG Three		Drug Exposure During					
times per day	424	DAY					
		Pregnancy					
		Neonatal Disorder					
		Oesophageal Atresia					

Date:12/17/01ISR Number: 3839290-2Report Type:Expedited (15-DaCompany Report #B0126849A  
Age:42 YR                      Gender:Male                      I/FU:F

Outcome	PT
Hospitalization -	Difficulty In Walking
Initial or Prolonged	Faecal Incontinence

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Multiple Sclerosis Urinary Incontinence		Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
150MG Per day			Zyban	PS	Glaxo Wellcome	ORAL
SUBCUTANEOUS			Interferon Beta	SS		
3UNIT per day			Lioresal	SS		ORAL
1000MG Twice			Paracetamol	C	Glaxo Wellcome	ORAL
per day						

Date:12/17/01ISR Number: 3839295-1Report Type:Expedited (15-DaCompany Report #B0128930A  
Age:34 YR Gender:Male I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Hospitalization -		Aphasia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Carotid Artery Thrombosis					
Initial or Prolonged		Cerebrovascular Accident		Ketoprofen	C		
per day		Erythema		Alcohol	C		
		Hemiplegia					
		Hypercholesterolaemia					
		Thrombosis					

Date:12/18/01ISR Number: 3840066-0Report Type:Expedited (15-DaCompany Report #A0162932A  
Age:41 YR Gender:Male I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Bradycardia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Cardiac Disorder					
per day		Hypotension		Tylenol #1	C	Glaxo Wellcome	ORAL
		Loss Of Consciousness					
		Syncope					
		Syncope Vasovagal					

Date:12/18/01ISR Number: 3840075-1Report Type:Expedited (15-DaCompany Report #A0169635A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Wellbutrin	PS	Glaxo Wellcome	ORAL
300MG Twice		Coma					
per day		Fall					
		Feeling Jittery					
		Medication Error					
		Urinary Incontinence					

Date:12/18/01ISR Number: 3840081-7Report Type:Expedited (15-DaCompany Report #B0126850A  
Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Myocardial Infarction		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day							
Hospitalization -				Bisoprolol	C		
Initial or Prolonged				Potassium	C		
				Pravastatin	C		
				Glipizide	C		
				Lysine	C		
				Frusemide	C	Glaxo Wellcome	
				Omeprazole	C		
				Nitroglycerin	C	Glaxo Wellcome	
				Repaglinide	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Clopidogrel C  
 Ramipril C

Date:12/18/01ISR Number: 3840088-XReport Type:Expedited (15-DaCompany Report #B0128862A  
 Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Abdominal Pain Upper		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Chest Pain					
		Cholelithiasis					
		Hepatitis					
		Oesophagitis					

Date:12/18/01ISR Number: 3841902-4Report Type:Expedited (15-DaCompany Report #2015123  
 Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiomegaly	Health	Morphine Sulfate			
		Coma	Professional	(Similar To Andas			
		Pulmonary Congestion	Other	74-769 And 74-862)	PS		
		Pulmonary Oedema		Oxycodone			
		Therapeutic Agent		Hydrochloride			
		Toxicity		(Similar To Nda			
		Toxicologic Test Abnormal		20-553)	SS		
				Codeine	SS		ORAL
PO				Olanzapine	SS		ORAL
PO				Acetaminophen	SS		ORAL
PO				Paroxetine	SS		ORAL
PO				Bupropion	SS		ORAL
PO				Nicotine	SS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability							
150MG Twice		Burning Sensation		Zyban	PS	Glaxo Wellcome	ORAL
per day	11	Dysgeusia					
	DAY	Dyskinesia		Amitriptyline	C		
UNKNOWN	200MG	Per day					
		Hypoaesthesia		Unknown Medication	C		
		Insomnia					
		Muscle Injury					
		Neuralgia					
		Neuropathy Peripheral					
		Oral Pain					
		Panic Attack					
		Rash					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability							
150MG Per day	62	Corneal Disorder		Zyban	PS	Glaxo Wellcome	ORAL
	DAY	Epistaxis					
		Retinal Detachment					
		Tension Headache					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/19/01ISR Number: 3840976-4Report Type:Expedited (15-DaCompany Report #A0170133A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Petit Mal Epilepsy	Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice			Status Epilepticus				
per day				Anti-Epileptic Drug	C		

Date:12/19/01ISR Number: 3840977-6Report Type:Expedited (15-DaCompany Report #A0170310A

Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -			Cardiac Failure	Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
150MG Per day							
Initial or Prolonged			Congestive Coronary Artery Disease Dystonia				

Date:12/19/01ISR Number: 3840978-8Report Type:Expedited (15-DaCompany Report #A0170371A

Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -			Convulsion	Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day							
Initial or Prolonged			Speech Disorder Weight Decreased				

Date:12/19/01ISR Number: 3840990-9Report Type:Expedited (15-DaCompany Report #B0125004A

Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -			Aggression	Zyban	PS	Glaxo Wellcome	
Initial or Prolonged			Depression	Mefanamic Acid	C		ORAL
100UNIT							

Unknown	Emotional Distress				
40UNIT Twice	Fatigue		Gamolenic Acid	C	ORAL
per day	Loss Of Consciousness				
	Memory Impairment				
	Weight Decreased				

Date:12/19/01ISR Number: 3840994-6Report Type:Expedited (15-DaCompany Report #B0129161A  
Age:66 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Cardiac Failure		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Pneumonia					
	Pulmonary Oedema					
	Ventricular Hypokinesia					

Date:12/20/01ISR Number: 3841759-1Report Type:Expedited (15-DaCompany Report #A0107071A  
Age:78 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Amnesia
Initial or Prolonged	Coma
	Confusional State
	Convulsion
	Delusion
	Dementia Alzheimer'S Type
	Depressed Level Of

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 DAY		Zyban	PS	Glaxo Wellcome	ORAL
			Celexa	SS		
			Hormones	C		
			Vitamins	C		

Date:12/20/01  
 Age:65 YR  
 Gender:Female  
 I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice		Anxiety		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -	per day	9 DAY	Aortic Valve Sclerosis					
Initial or Prolonged	150MG Twice		Asthenia		Wellbutrin	SS	Glaxo Wellcome	ORAL
per day			Atrioventricular Block					
.188MG Twice			Cardiac Failure		Hyoscyamine	SS		
per day			Cardiac Failure					
TOPICAL	1IN Twice per		Congestive Cardiomegaly		Relafen Nitroglycerine	C C	Glaxo Wellcome Glaxo Wellcome	
day			Cerebral Infarction					
			Cerebrovascular Accident		Synthroid	C	Glaxo Wellcome	
			Confusional State		Lasix	C	Glaxo Wellcome	
4MG Per day			Dysarthria		Risperdal	C		
250MG Twice			Dyspnoea		Depakote	C		
per day			Ejection Fraction					
			Decreased		Sinemet Cr	C		
2MG Twice per			Electrocardiogram Qrs		Benztropine	C		



day	Complex Prolonged				
	Electrocardiogram T Wave Abnormal	Zestril	C		
		Azmacort	C		
2PUFF Twice	Facial Palsy				
per day	Headache	Atrovent	C	Glaxo Wellcome	
2PUFF Twice	Hemiparesis				
per day	Hypoaesthesia	Peri-Colace	C		
100MG Per day	Mitral Valve Incompetence	Cipro	C		ORAL
500MG Twice	Tongue Dry				
per day	Ventricular Extrasystoles				
	Ventricular Hypertrophy				
	Ventricular Hypokinesia				
	Vision Blurred				
	Vomiting				

Date:12/20/01ISR Number: 3841765-7Report Type:Expedited (15-DaCompany Report #A0169595A  
Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Apathy		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Hallucination, Auditory		Synthroid	C	Glaxo Wellcome	ORAL
.075MCG Per		Memory Impairment					
day		Paranoia		Ritalin	C		ORAL
25MG Twice		Rash					
per day		Scar		Vitamin	C		ORAL
15 DAY		Suicidal Ideation		Depakote	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/20/01ISR Number: 3841780-3Report Type:Expedited (15-DaCompany Report #B0127463A  
Age:54 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300MG Per day 11 DAY	Abdominal Pain		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Malaise Syncope Vasovagal		Climaston	C		

Date:12/20/01ISR Number: 3841784-0Report Type:Expedited (15-DaCompany Report #B0129486A  
Age:31 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice	Confusional State		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day 3 DAY	Dysarthria Feeling Drunk Vomiting					

Date:12/20/01ISR Number: 3841785-2Report Type:Expedited (15-DaCompany Report #B0129668A  
Age:52 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice	Hypothyroidism		Zyban	PS	Glaxo Wellcome	ORAL
per day	Urticaria					

Date:12/20/01ISR Number: 3846396-0Report Type:Periodic Company Report #HQ9690310AUG2000  
Age:35 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Abortion Spontaneous Headache Pregnancy Vision Blurred	Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended			

1) 75 MG 1X

Release)

PS

ORAL

PER 1 DAY,

ORAL; 2)

ORAL

Tegretol  
(Carbamazepine, )

SS

ORAL

ORAL

Wellbutrin  
(Amfebutamone  
Hydrochloride, )

SS

ORAL

ORAL

Zyrtec (Cetirizine  
Hydrochloride,)

SS

ORAL

ORAL

Date:12/20/01ISR Number: 3848232-5Report Type:Periodic  
Age:83 YR Gender:Male I/FU:I

Company Report #HQ4476109AUG2001

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Aphasia Pyrexia	Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule,	PS		ORAL
150 MG 1X PER							
1 DAY, ORAL							
				Wellbutrin (Amfebutamone Hydrochloride, )	SS		ORAL
150 MG 2X PER							
1 DAY, ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/21/01ISR Number: 3842767-7Report Type:Expedited (15-DaCompany Report #A0170813A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Intentional Misuse		Unspecified			
		Suicide Attempt		Medications	SS		
UNKNOWN							

Date:12/21/01ISR Number: 3842788-4Report Type:Expedited (15-DaCompany Report #B0126402A

Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Atrioventricular Block		Zyban	PS	Glaxo Wellcome	ORAL
23 DAY							
Hospitalization -		Second Degree					
Initial or Prolonged		Bradycardia					
Disability		Coronary Artery Surgery					
Other		Electrocardiogram					
		Repolarisation					
		Abnormality					
		Feeling Cold					
		Flushing					
		Headache					
		Myocardial Infarction					
		Pallor					

Date:12/21/01ISR Number: 3842804-XReport Type:Expedited (15-DaCompany Report #B0128266A

Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Initial or Prolonged		Blood Pressure Decreased					
per day	13 DAY						
		Electrocardiogram St		Fexofenadine	SS		
UNKNOWN	180MG per day 7 DAY						
		Segment Depression					
		Syncope					
		Urticaria					

Date:12/21/01ISR Number: 3842813-0Report Type:Expedited (15-DaCompany Report #B0129422A  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Duodenal Ulcer Gastric Haemorrhage Haematocrit Decreased Malaise		Zyntabac Pantoprazole	PS C	Glaxo Wellcome	

Date:12/21/01ISR Number: 3842818-XReport Type:Expedited (15-DaCompany Report #B0129835A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Overdose Somnolence Suicide Attempt		Zyban Zolpidem Antidepressant	PS SS SS	Glaxo Wellcome	ORAL ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/21/01ISR Number: 3842827-0Report Type:Expedited (15-DaCompany Report #B0130322A

Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Fall		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Food Craving		Aspirin	C		
		Hypoglycaemia		Nicotine	C	Glaxo Wellcome	
		Loss Of Consciousness					
		Muscle Rigidity					
		Scratch					
		Trismus					

Date:12/21/01ISR Number: 3842830-0Report Type:Expedited (15-DaCompany Report #B0130344A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day				Dothiepin	SS		ORAL
50MG At night				Prempak-C	C		ORAL
.625MG							
Unknown				Salbutamol	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)	100MCG As						
required				Diazepam	C		
UNKNOWN	5MG As						
required				Fluoxetine	C		
UNKNOWN	40MG Per day			Zopiclone	C		
UNKNOWN	7.5MG At						
night				Beclomethasone	C	Glaxo Wellcome	
RESPIRATORY							

(INHALATION) 100U Unknown

Date:12/21/01ISR Number: 3842836-1Report Type:Expedited (15-DaCompany Report #B0130602A

Age:56 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice Initial or Prolonged per day 63 DAY	Angina Unstable		Zyban	PS	Glaxo Wellcome	ORAL
UNKNOWN 1U Twice per day			Ranitidine	C	Glaxo Wellcome	
RESPIRATORY (INHALATION) 2PUFF As required			Beclomethasone Dipropionate	C	Glaxo Wellcome	
UNKNOWN 1U As directed			Aspirin	C		
UNKNOWN 1U Per day			Simvastatin	C		
UNKNOWN 1U Per day			Lansoprazole	C		

Date:12/26/01ISR Number: 3844395-6Report Type:Expedited (15-DaCompany Report #2001027085-1

Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 30 MILLIGRAMS Disability 1.0 DAILY Required ORAL	Drug Ineffective Fracture Grand Mal Convulsion Joint Dislocation Sedation	Health Professional	Paxil Glaxosmithkline	PS	Glaxosmithkline	ORAL
Intervention to Prevent Permanent Impairment/Damage SEE IMAGE 3 DAY			Wellbutrin Sr (Bupropion) Glaxowellcome	SS	Glaxowellcome	ORAL





FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/26/01ISR Number: 3844757-7Report Type:Expedited (15-DaCompany Report #001-0945-M0101302

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG Other (TID), PER ORAL	Accommodation Disorder Astigmatism Dyspepsia Hypoaesthesia Malaise Memory Impairment Mental Disorder	Health Professional	Neurontin (Gabapentin)	PS		ORAL
40 MG (QID), PER ORAL			Wellbutrin (Amfebutamone Hydrochloride) Methadone (Methadone)	SS SS		ORAL
			(Paracetamol, Hydrocodone Bitartrate)	C		

Date:12/26/01ISR Number: 3844848-0Report Type:Direct

Age:57 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200MG Q12 ORAL	Medication Error		Wellbutrin Sr 100mg Glaxo-Wellcome	PS	Glaxo-Wellcome	ORAL

Date:12/26/01ISR Number: 3844893-5Report Type:Direct

Age:29 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose 100 MG DAILY	Heart Rate Increased		Wellbutrin Sr 100 Mg	PS		ORAL

ORAL

Palpitations

Tremor

Zoloft

C

Synthroid

C

Prevacid

C

Date:12/26/01ISR Number: 3850322-8Report Type:Periodic

Company Report #A0136960A

Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent SEE DOSAGE Impairment/Damage TEXT/ ORAL		Eosinophilia Leukopenia	Health Professional Company Representative	Wellbutrin Tablet (Bupropionhydrochloride)	PS		ORAL
SEE DOSAGE TEXT / ORAL				Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	SS		ORAL
				Valproic Acid	C		

Date:12/27/01ISR Number: 3844300-2Report Type:Expedited (15-DaCompany Report #A0162401A

Age:45 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Condition Aggravated Drug Effect Decreased Loss Of Employment

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Mania Mood Swings Suicidal Ideation					
150MG Per day	1 WK	Tobacco Abuse		Zyban	PS	Glaxo Wellcome	ORAL
UNKNOWN				Topamax	SS		
UNKNOWN				Zyprexa	SS		
UNKNOWN	1MG Unknown			Klonopin	SS		

Date:12/27/01ISR Number: 3844301-4Report Type:Expedited (15-DaCompany Report #A0165315A  
Age:40 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Twice	per day	Deafness		Wellbutrin	PS	Glaxo Wellcome	ORAL
		10 DAY	Tinnitus		Paxil	C	Glaxo Wellcome	
					Klonopin	C		

Date:12/27/01ISR Number: 3844303-8Report Type:Expedited (15-DaCompany Report #A0170847A  
Age:60 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	400MG Per day		Depressed Level Of Consciousness		Wellbutrin Sr Uniphyl	PS C	Glaxo Wellcome	ORAL
	325MG Per day		Inappropriate Antidiuretic Hormone		Lipidil Novasen	C C		
	.5MG Three times per day		Secretion		Rivotril	C		
	2.5MG Per day		Polydipsia		Norvasc	C		

.5MG At night

Risperdal	C	
Zoloft	C	
Kemadrin	C	Glaxo Wellcome
Norvasen	C	

Date:12/27/01ISR Number: 3855970-7Report Type:Periodic Company Report #WAES 01091620  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Tab Vioxx 25 Mg	PS		ORAL
25		Asthenia					
MG/DAILY/PO		Fatigue		Meridia	SS		ORAL
15		Nervousness					
MG/DAILY/PO				Synthroid	SS		ORAL
50							
MICROGM/DAILY							
/PO				Lorazepam	SS		ORAL
1 MG/BID/PO				Wellbutrin	SS		ORAL
150							
MG/DAILY/PO							

Date:01/02/02ISR Number: 3846136-5Report Type:Expedited (15-DaCompany Report #B0128152A  
 Age:54 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Fall
Initial or Prolonged	Fatigue
	Gamma-Glutamyltransferase
	Increased
	Grand Mal Convulsion

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
40	DAY	Hyperhidrosis Loss Of Consciousness Malaise		Zyban	PS	Glaxo Wellcome	ORAL
160MG per day		Orthostatic Hypotension		Valsartan	SS		ORAL
200MG per day		Pallor		Celiprolol	SS		ORAL
40MG per day	8 DAY	Syncope Vasovagal		Frusemide	SS	Glaxo Wellcome	ORAL
		Tooth Injury		Atorvastatin	C		ORAL

Date:01/02/02ISR Number: 3846140-7Report Type:Expedited (15-DaCompany Report #B0130243A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Antinuclear Antibody Positive Asthenia Bilirubin Conjugated Increased Hepatic Enzyme Increased Hepatitis Cholestatic Hepatocellular Damage Jaundice Nausea		Bupropion	PS	Glaxo Wellcome	

Date:01/02/02ISR Number: 3846146-8Report Type:Expedited (15-DaCompany Report #B0130748A  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN	150MG Twice	Circulatory Collapse		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged per day	10 DAY	Ventricular Tachycardia		Diuretic	C		

Date:01/02/02ISR Number: 3847519-XReport Type:Direct  
Age:48 YR Gender:Female I/FU:I

Company Report #CTU 158144

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash Generalised Rash Pruritic		Bupropian 100 Mg Kaiser Permanente	PS	Kaiser Permanente	ORAL
100 MIG		Skin Tightness					
TWICE PER							
ORAL							

Date:01/02/02ISR Number: 3847957-5Report Type:Direct  
Age:23 YR Gender:Female I/FU:I

Company Report #CTU 158146

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Confusional State Crying		Paxil 40mg Smithklinebeechem	PS	Smithklinebeechem	ORAL
40 MG ORAL		Depression		Wellbutrin 50-150 Mg	SS		ORAL
50-15 MG ORAL		Drug Withdrawal Syndrome					
0 MG		Feeling Abnormal Feeling Hot Irritability Nervousness Paraesthesia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/04/02ISR Number: 3848973-XReport Type:Expedited (15-DaCompany Report #2014816

Age:32 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Bronchopneumonia Drug Toxicity Toxicologic Test Abnormal	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553) Fluoxetine Hydrochloride Chlordiazepoxide Bupropion Hydrochloride Cocaine Trazodone Hcl Insulin Lithium Risperdal (Risperidone)	PS SS SS SS SS C C C		

Date:01/07/02ISR Number: 3848657-8Report Type:Expedited (15-DaCompany Report #B0130993A

Age:55 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 15 DAY			Abdominal Pain		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged 1UD per day		87 DAY	Balance Disorder		Amlodipine Besylate	C		ORAL
			Flatulence Gastrointestinal Disorder Neoplasm Skin		Lithium Carbonate	C	Glaxo Wellcome	ORAL

Date:01/07/02ISR Number: 3849470-8Report Type:Expedited (15-DaCompany Report #2014857

Age:37 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Atherosclerosis Drug Toxicity Toxicologic Test Abnormal	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553)	PS		

Carbamazepine SS  
Propoxyphene Hcl SS  
Bupropion  
Hydrochloride SS  
Citalopram SS

Date:01/07/02ISR Number: 3850142-4Report Type:Direct  
Age:39 YR Gender:Female I/FU:I

Company Report #CTU 158536

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2 WEEKS PTA Initial or Prolonged	2 WK	Angioneurotic Oedema Dyspnoea Face Oedema Rash		Wellbutrin	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/08/02ISR Number: 3849090-5Report Type:Expedited (15-DaCompany Report #A0167755A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cough		Wellbutrin	PS	Glaxo Wellcome	ORAL
100MG Twice							
per day	24	DAY					
		Dysphonia					
		Urticaria					
		Wheezing					

Date:01/08/02ISR Number: 3849093-0Report Type:Expedited (15-DaCompany Report #A0171241A  
Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin	PS	Glaxo Wellcome	ORAL
300MG Per day							
		Maternal Drugs Affecting					
		Foetus					

Date:01/08/02ISR Number: 3849095-4Report Type:Expedited (15-DaCompany Report #B0116763A  
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	6	DAY					
		Overdose					
				Alcohol	C		ORAL

Date:01/08/02ISR Number: 3849104-2Report Type:Expedited (15-DaCompany Report #B0131087A  
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Overdose		Zyban	PS	Glaxo Wellcome	
UNKNOWN	12000MG						

Initial or Prolonged Suicide Attempt  
cumulative

dose

Date:01/08/02ISR Number: 3849873-1Report Type:Expedited (15-DaCompany Report #A130158  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Pain In Extremity	Health	Procardia Capsules	PS		
Intervention to Prevent Permanent		Peripheral Coldness	Professional	Amlodipine	SS		
SUBCUTANEOUS	8.00	Peripheral Vascular Disorder		Betaseron	SS		
Impairment/Damage							
TOTAL:EVERY							
OTHER DAY:		Pulse Abnormal					
SUBCUTANEOUS		Skin Discolouration					
				Wellburtrin	SS		
				Vivelle	C		
				Neurontin	C		
				Prozac	C		
				Zanaflex	C		
				Darvocet	C		
				Xanax	C		

Date:01/08/02ISR Number: 3850083-2Report Type:Direct Company Report #CTU 158601  
Age:27 YR Gender:Female I/FU:I

Outcome	PT
	Grand Mal Convulsion
	Vertigo

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Visual Disturbance

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
PO 80-160MG			Septra (Co-Trimaxazole )	PS		ORAL
BID 150MG (PO)	2 DAY		Zyban	SS		ORAL
BID	9 MON		Norplant	C		
			Prozac	C		
			Loratadine	C		
			Trimeth	C		
			Benzamide 250mg	C		
			Loperamide	C		

Date:01/08/02ISR Number: 3850518-5Report Type:Direct  
Age:45 YR Gender:Female I/FU:I

Company Report #CTU 158619

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Other 150MG BID PO	2 YR	Asthma Cardiomyopathy		Bupropion (Welbutrin Sr)	PS		ORAL

Date:01/09/02ISR Number: 3849503-9Report Type:Expedited (15-DaCompany Report #B0128084A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150MG Per day	6 DAY	Asthenia Dizziness Nausea Vision Blurred		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL

Date:01/09/02ISR Number: 3850947-XReport Type:Direct Company Report #CTU 158777  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Blood Pressure Increased Palpitations		Wellbutrin Sr 150mg. Glax	PS	Glax	

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Date:01/10/02ISR Number: 3850005-4Report Type:Expedited (15-DaCompany Report #A0164944A  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG See		Amnesia		Wellbutrin	PS	Glaxo Wellcome	ORAL
dosage text		Blood Potassium Decreased					
60MG Per day		Dizziness		Prozac	C		ORAL
		Grand Mal Convulsion Stress					

Date:01/10/02ISR Number: 3850018-2Report Type:Expedited (15-DaCompany Report #A0171493A  
Age:32 YR Gender:Female I/FU:F

Outcome	PT
Other	Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Drugs					
		Maternal Drugs Affecting					
		Foetus					
		Placental Disorder	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Pregnancy		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Per day							

Date:01/10/02ISR Number: 3850020-0Report Type:Expedited (15-DaCompany Report #A0171611A  
 Age: Gender:Female I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Vaginal Haemorrhage		Wellbutrin	PS	Glaxo Wellcome	ORAL
				Zyban	SS	Glaxo Wellcome	ORAL

Date:01/10/02ISR Number: 3850026-1Report Type:Expedited (15-DaCompany Report #B0131129A  
 Age: Gender:Male I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Myocardial Infarction		Zyban	PS	Glaxo Wellcome	
UNKNOWN		Renal Failure					

Date:01/10/02ISR Number: 3850974-2Report Type:Expedited (15-DaCompany Report #2014808  
 Age:46 YR Gender:Female I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Death		Coma	Health	Oxycontin Cr			
		Drug Toxicity	Professional	Tablets, 40 Mg			
		Ecchymosis	Other	(Oxycodone			
		Laceration		Hydrochloride)	PS		ORAL
40 MG PO		Toxicologic Test Abnormal		Doxepin Hcl	SS		
				Bupropion			
				Hydrochloride	SS		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150 MG 3X/DAY	Abnormal Behaviour		Wellbutrin Sr	PS		
Initial or Prolonged Disability	Aggression Agitation Delusion Gender Identity Disorder Insomnia Mania Mental Impairment Paranoia Pressure Of Speech Psychomotor Hyperactivity Thinking Abnormal Tinnitus Transvestism		Lipitor Thorazine	C C		

Outcome	PT
Other	Anxiety Confusional State

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Convulsion Depression Insomnia Somnolence	Report Source	Product	Role	Manufacturer	Route
300 MG BID PO, 900 MG DAILY PO, 1200 MG DAILY PO	1 DAY		Literature Health Professional	Lithium (Lithium)	PS		ORAL
150 MG BID PO				Venlafaxine (Venlafaxine)	SS		ORAL
100 MG BID PO, 300 MG DAILY PO, 200 MG DAILY PO				Bupropion (Bupropion)	SS		ORAL
				Gabapentin (Gabapentin)	C		
				Clonazepam (Clonazepam)	C		
				Glycopyrrolate (Glycopyrrolate)	C		
				Methohexital (Methohexital)	C		
				Succinylcholine (Succinylcholine Chloride)	C		
				Oxygen (Oxygen)	C		

Date:01/11/02ISR Number: 3850913-4Report Type:Expedited (15-DaCompany Report #A0170847A  
Age:60 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization -	Coma	Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
100MG Per day 9 DAY					
Initial or Prolonged	Convulsion	Zoloft	SS		ORAL
100MG Per day YR					
	Inappropriate	Risperdal	SS		ORAL
1MG Per day					
	Antidiuretic Hormone	Uniphyl	C		
400MG Per day					
	Secretion	Lipidil	C		
	Loss Of Consciousness	Novasen	C		
325MG Per day					
	Polydipsia	Rivotril	C		ORAL
.5MG Twice					
	Respiratory Tract				
per day					
	Infection	Norvasc	C		
2.5MG Per day					
		Kemadrin	C	Glaxo Wellcome	

Date:01/14/02ISR Number: 3852218-4Report Type:Expedited (15-DaCompany Report #B0130243A  
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Foreign	Bupropion			
		Hepatitis Cholestatic	Literature	Hydrochloride			
		Jaundice	Health	(Bupropion			
		Nausea	Professional	Hydrochloride)	PS		

Date:01/17/02ISR Number: 3854205-9Report Type:Expedited (15-DaCompany Report #B0125399A  
Age:42 YR Gender:Female I/FU:I

Outcome  
Hospitalization -  
Initial or Prolonged



Freedom Of Information (FOI) Report

Disability

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG PER DAY ORAL		Balance Disorder Cerebellar Syndrome Cerebrovascular Accident  Confusional State  Coordination Abnormal Difficulty In Walking Dizziness Dysarthria Fall Fear Fumbling Hallucination Hallucination, Visual Hypoaesthesia Lung Consolidation Paresis Pharyngolaryngeal Pain Swelling Tenderness Tongue Paralysis Vision Blurred	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)  Ferrous Sulfate Promethazine Propiomazine	PS  C C C		ORAL

Date:01/17/02ISR Number: 3854978-5Report Type:Expedited (15-DaCompany Report #A0155826A  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged  150 MG / TWICE PER DAY  / ORAL		Abdominal Pain Diarrhoea Haemorrhagic Gastritis Gastrointestinal Disorder  Haemorrhoids  Hyperplasia  Hypersensitivity Rectal Haemorrhage	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)  Famotidine Hrt	PS  C C		ORAL

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Aggression	Consumer	Wellbutrin			
Initial or Prolonged	Anxiety		Unspecified Tablet			
	Contusion		(Bupropion			
	Convulsion		Hydrochloride)	PS		
VARIALE DOSE	Depression					
	Homicidal Ideation					
	Injury					
	Memory Impairment					
	Mental Disorder					
	Premenstrual Syndrome					
	Suicidal Ideation					
	Swelling					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/17/02ISR Number: 3855045-7Report Type:Expedited (15-DaCompany Report #A0165315A

Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Deafness Neurosensory Tinnitus	Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
150 MG TWICE							
PER DAY ORAL							
				Paroxetine Hydrochloride	C		
				Vitamin E	C		
				Ascorbic Acid	C		
				Clonazepam	C		
				Calcium Carbonate	C		
				Multivitamins + Iron	C		
				Hydrochlorothiazide	C		

Date:01/17/02ISR Number: 3855861-1Report Type:Expedited (15-DaCompany Report #B0126090A

Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged		Hypoaesthesia Metastases To Central Nervous System	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG PER							
DAY ORAL		Monoparesis					
		Neurological Symptom					

Date:01/18/02ISR Number: 3854120-0Report Type:Expedited (15-DaCompany Report #B0120704A

Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Agitation		Zyban	PS	Glaxo Wellcome	
150MG Twice							
per day	21 DAY	Anxiety					

Breast Pain  
Confusional State  
Depression  
Disturbance In Attention  
Dizziness  
Exanthem  
Feeling Abnormal  
Hallucinations, Mixed  
Insomnia  
Irritability  
Malaise  
Mental Disorder  
Mood Altered  
Nausea  
Pain  
Pollakiuria  
Pruritus  
Psoriasis  
Seborrhoea  
Tremor  
Visual Disturbance

Zopiclone C  
Oxazepam C  
Naproxen C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/18/02ISR Number: 3854130-3Report Type:Expedited (15-DaCompany Report #B0132126A

Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour		Zyban	PS	Glaxo Wellcome	
150MG Twice		Automatism					
per day	10	DAY		Carbamazepine	C		
2TAB Twice		Complex Partial Seizures					
per day		Confusional State					
1.5TAB Twice		Deja Vu		Lithium	C		
per day		Echopraxia					
1TAB Twice		Epilepsy		Phenytoin	C		
per day		Hallucination, Auditory					
1TAB Per day		Thinking Abnormal		Amitriptyline	C		
1TAB Three				Amoxicillin	C		
times per day	10	DAY					

Date:01/18/02ISR Number: 3855669-7Report Type:Expedited (15-DaCompany Report #HQ9911115JAN2002

Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Consumer	Effexor Xr			
		Drug Withdrawal Syndrome		(Venlafaxine			
		Hepatitis		Hydrochloride,			
		Nausea		Capsule, Extended			
ORAL		Vomiting		Release)	PS		ORAL
				Wellbutrin			
				(Amfebutamone			
ORAL				Hydrochloride)	SS		ORAL

Date:01/18/02ISR Number: 3856814-XReport Type:Expedited (15-DaCompany Report #B0131464A  
Age:25 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Myocardial Infarction Purpura	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ORAL			Methylenedioxymetham phet (Formulation Unknown) (Methylenedioxymetha mphet)	SS		

Date:01/21/02ISR Number: 3854442-3Report Type:Expedited (15-DaCompany Report #A0172430A  
Age:59 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Anxiety Asthenia Blood Pressure Increased Constipation Depression Dizziness Excitability Immobile Insomnia Irritability Nervousness Oral Intake Reduced Rash Pruritic

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice			Lamictal	PS	Glaxo Wellcome	ORAL
per day			Wellbutrin	SS	Glaxo Wellcome	ORAL
			Paxil	SS	Glaxo Wellcome	ORAL
			Nardil	SS		
UNKNOWN			Zoloft	SS		
UNKNOWN			Ativan	C		
20 YR			Norvasc	C		
			Avapro	C		
			Klonopin	C		

Date:01/21/02ISR Number: 3854449-6Report Type:Expedited (15-DaCompany Report #B0131906A  
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Gastrointestinal Disorder		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Intestinal Dilatation					
per day	12 DAY	Intestinal Obstruction					

Date:01/22/02ISR Number: 3858046-8Report Type:Expedited (15-DaCompany Report #B0131893A  
 Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Atrial Fibrillation	Foreign	Zyban Tablet			
Initial or Prolonged		Cerebral Infarction		(Bupropion			
150MG/ORAL		Dysphasia		Hydrochloride)	PS		ORAL
		Memory Impairment					
		Nausea					
		Personality Disorder					

Ventricular Extrasystoles  
Visual Disturbance

Date:01/22/02ISR Number: 3858052-3Report Type:Expedited (15-DaCompany Report #B0131129A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dehydration Dizziness Dyspepsia Heart Rate Increased	Foreign Consumer	Bupropion Hydrochloride Tablet - Zyban (Bupropion Hydrochloride)	PS		
150 MG		Myocardial Infarction Renal Failure Ventricular Extrasystoles					

Date:01/22/02ISR Number: 3858057-2Report Type:Expedited (15-DaCompany Report #B0132111A

Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Testis Cancer	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG TWICE PER DAY ORAL				Salbutamol Sulphate	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/22/02ISR Number: 3858183-8Report Type:Expedited (15-DaCompany Report #A0169635A

Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Health	Wellbutrin Sr			
Required		Coma	Professional	Tablet-Controlled			
Intervention to		Fall		Release (Bupropion			
Prevent Permanent		Feeling Jittery		Hydrochloride)	PS		ORAL
200 MG/ TWICE							
Impairment/Damage		Medication Error					
PER DAY/ ORAL							
		Overdose		Conjugated Estrogens	C		
		Urinary Incontinence		Valaciclovir			
				Hydrochlorid	C		
				Gabapentin	C		

Date:01/23/02ISR Number: 3856112-4Report Type:Expedited (15-DaCompany Report #A0169561A

Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Vision Blurred		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	22 DAY	Visual Acuity Reduced					

Date:01/23/02ISR Number: 3856115-XReport Type:Expedited (15-DaCompany Report #A0173197A

Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Circulatory Collapse		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day		Myocardial Infarction					
		Nausea		Entex Pse	C		ORAL
		Urinary Incontinence					
		Vomiting					

Date:01/23/02ISR Number: 3856127-6Report Type:Expedited (15-DaCompany Report #B0132864A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Antinuclear Antibody		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Positive					
per day	6	DAY					
		Erythema					
		Oedema Peripheral					
		Swelling					
		Tenderness					

Date:01/23/02ISR Number: 3856157-4Report Type:Expedited (15-DaCompany Report #301407  
Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Disturbance In Attention		Lariam	PS	Roche	
36	DAY						
		Dry Mouth		Zyban	SS		
7	DAY						
		Energy Increased		Trinordiol	C		
		Hyperhidrosis					
		Insomnia					
		Tremor					
		Vertigo					
		Vision Blurred					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/23/02ISR Number: 3859513-3Report Type:Expedited (15-DaCompany Report #A130158

Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Chest Pain	Health	Procardia Capsules	PS		
Intervention to		Pain In Extremity	Professional	Amlodipine	SS		
Prevent Permanent		Peripheral Coldness		Betaseron	SS		
SUBCUTANEOUS	8.00						
Impairment/Damage		Peripheral Vascular					
TOTAL:EVERY		Disorder					
OTHER DAY:							
SUBCUTANEOUS				Wellbutrin	SS		
				Vivelle	C		
				Neurontin	C		
				Prozac	C		
				Zanaflex	C		
				Xanax	C		
				Darvocet	C		

Date:01/23/02ISR Number: 3859628-XReport Type:Direct

Company Report #CTU 159951

Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Anxiety		Wellbutrin 100 Mg			
Hospitalization -		Asthenia		B.I.D.	PS		
100 MG							
Initial or Prolonged		Blood Pressure Decreased					
B.I.D.							
Disability		Catatonia					
Required		Cognitive Disorder					
Intervention to		Confusional State					
Prevent Permanent		Decreased Activity					
Impairment/Damage		Depression					
		Disorientation					
		Hallucination					
		Hypoaesthesia					
		Malaise					
		Palpitations					
		Speech Disorder					
		Suicidal Ideation					

Date:01/23/02ISR Number: 3859836-8Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #NSADSS2001014108

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tardive Dyskinesia	Consumer	Risperdal (2 Mg Tablet) (Risperidone)	PS		ORAL
2 MG, 1 IN 1 DAY(S), ORAL							
ORAL				Prozac (Fluoxetine Hydrochloride)	SS		ORAL
ORAL				Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL

Date:01/23/02ISR Number: 3859988-XReport Type:Direct  
Age:33 YR Gender:Female I/FU:I

Company Report #CTU 159889

Outcome	PT
Other	Agitation Dizziness Headache

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Tremor Vision Blurred				
Dose	Duration		Report Source	Product	Role	Manufacturer
				Wellbutrin Sr 150mg Glaxo	PS	Glaxo
150 MG 2X/DAY						
ORAL						ORAL

Date:01/23/02ISR Number: 3860089-5Report Type:Direct Company Report #CTU 160078  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Confusional State		Zyban Catalytica	PS	Catalytica	ORAL
1 TAB 150 MG							
Hospitalization -		Depression					
2X DAY ORAL							
Initial or Prolonged		Insomnia		Paxil	C		
Disability		Suicidal Ideation					
Required		Syncope					
Intervention to		Tremor					
Prevent Permanent							
Impairment/Damage							

Date:01/24/02ISR Number: 3856839-4Report Type:Expedited (15-DaCompany Report #B0120523A  
Age:61 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardiac Failure Acute		Zyban	PS	Glaxo Wellcome	ORAL
42 DAY							
UNKNOWN		Myocardial Fibrosis		Sotalol	C		
UNKNOWN		Sudden Death		Warfarin	C	Glaxo Wellcome	

Date:01/24/02ISR Number: 3856844-8Report Type:Expedited (15-DaCompany Report #B0132587A  
Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day 8 DAY	Arterial Rupture		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Depressed Level Of Consciousness Haemorrhage Headache Hypertension Meningeal Disorder Musculoskeletal Stiffness Vomiting					

Date:01/24/02ISR Number: 3856845-XReport Type:Expedited (15-DaCompany Report #B0133010A  
Age:27 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day 24 DAY	Gastric Ulcer		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Haematemesis Haematocrit Decreased Middle Insomnia Nausea Oesophagitis Ulcerative Sleep Disorder Vomiting					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/25/02ISR Number: 3857846-8Report Type:Expedited (15-DaCompany Report #A0164667A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	4 MON			Zyban	PS	Glaxo Wellcome	ORAL
		Muscle Twitching					
		Nerve Injury					
		Nervous System Disorder					
		Pain In Extremity					
		Tremor					

Date:01/25/02ISR Number: 3857850-XReport Type:Expedited (15-DaCompany Report #A0169595A  
Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	.075MCG Per day			Wellbutrin	PS	Glaxo Wellcome	ORAL
		Anxiety		Synthroid	C	Glaxo Wellcome	ORAL
		Blister					
		Depression					
	25MG Twice per day			Ritalin	C		ORAL
		Erythema					
		Hallucination, Auditory					
	15 DAY			Vitamin	C		ORAL
		Memory Impairment		Depakote	C		ORAL
		Nervousness					
		Paranoia					
		Pruritus					
		Rash Generalised					
		Scratch					
		Suicidal Ideation					

Date:01/25/02ISR Number: 3857851-1Report Type:Expedited (15-DaCompany Report #A0170813A  
Age:17 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - YR				Wellbutrin	PS	Glaxo Wellcome	ORAL
		Convulsion					

Initial or Prolonged	Euphoric Mood	Soma	SS
UNKNOWN			
	Overdose	Neurontin	SS
UNKNOWN			
	Suicide Attempt	Effexor	C

Date:01/25/02ISR Number: 3857856-0Report Type:Expedited (15-DaCompany Report #A0173850A  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia		Wellbutrin	PS	Glaxo Wellcome	ORAL
100MG Per day		Cyst		Ssri	C		

Date:01/25/02ISR Number: 3860071-8Report Type:Expedited (15-DaCompany Report #301407  
 Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Disturbance In Attention	Foreign	Lariam			
		Dry Mouth	Health	(Mefloquine			
		Energy Increased	Professional	Hydrochloride)	PS		ORAL
1 DOSE FORM 1		Hyperhidrosis					
PER WEEK ORAL		Insomnia		Zyban (Bupropion			
		Tremor		Hydrochloride)	SS		ORAL
1 DOSE FORM		Vertigo					
DAILY ORAL		Vision Blurred		Trinordiol (Ethinyl			
				Estradiol/Levonorges			
				trel)	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/25/02ISR Number: 3860273-0Report Type:Expedited (15-DaCompany Report #B0130243A

Age:49 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged  150 MG /  TWICE PER DAY  /	Antinuclear Antibody Positive Asthenia Cholestasis  Chromaturia  Faeces Discoloured  Hepatitis Acute Hepatitis Cholestatic Hepatomegaly Hepatotoxicity Jaundice Liver Tenderness Nausea Prothrombin Time Prolonged	Foreign Literature Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS         C C		

Date:01/28/02ISR Number: 3858975-5Report Type:Expedited (15-DaCompany Report #B0131464A

Age:25 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Unknown Initial or Prolonged  UNKNOWN	Blood Creatinine Increased Drug Abuser  Ecchymosis Myocardial Infarction Toxicologic Test Abnormal		Zyban  Methylenedioxymetham phetamine	PS  SS	Glaxo Wellcome	ORAL

Date:01/28/02ISR Number: 3858991-3Report Type:Expedited (15-DaCompany Report #A0166999A

Age:37 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Other	Abortion Spontaneous	Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice					
	Maternal Drugs Affecting				
per day					
	Foetus	Synthroid	C	Glaxo Wellcome	ORAL
.75MG Per day					

Date:01/28/02ISR Number: 3858998-6Report Type:Expedited (15-DaCompany Report #B0131129A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG	Atrial Flutter		Zyban	PS	Glaxo Wellcome	
UNKNOWN	150MG	Unknown 7 DAY					
Initial or Prolonged		Condition Aggravated					
		Dehydration					
		Dizziness					
		Dyspepsia					
		Electrolyte Imbalance					
		Myocardial Infarction					
		Palpitations					
		Renal Failure					
		Ventricular Tachycardia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/02ISR Number: 3859009-9Report Type:Expedited (15-DaCompany Report #B0133231A  
 Age:63 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice Initial or Prolonged per day	Duration 9 DAY	Blood Pressure Systolic Increased Chest X-Ray Abnormal Dyspnoea	Zyban	PS	Glaxo Wellcome	ORAL
8MG Unknown		Hypoxia	Candesartan Cilexetil	SS		ORAL
12.5MG		Infection	Hydrochlorothiazide	SS		ORAL
Unknown		Spirometry Abnormal	Thyroxine Sodium Oestradiol	SS SS	Glaxo Wellcome	ORAL
UNKNOWN			Progesterone	SS		
UNKNOWN			Fluoxetine Hydrochloride	SS		ORAL
20MG Unknown			Dirithromycin	C		
UNKNOWN			Toplexil	C		
UNKNOWN			Erdosteine	C		

Date:01/28/02ISR Number: 3859010-5Report Type:Expedited (15-DaCompany Report #B0133703A  
 Age:52 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Unknown Initial or Prolonged	Duration	Chest Pain Condition Aggravated Coronary Artery Disease	Zyban	PS	Glaxo Wellcome	ORAL
			Simvastatin	SS		ORAL

Date:01/28/02ISR Number: 3859011-7Report Type:Expedited (15-DaCompany Report #B0133707A  
 Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Thrombotic		Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown							
Hospitalization -		Thrombocytopenic Purpura					
Initial or Prolonged							

Date:01/28/02ISR Number: 3859012-9Report Type:Expedited (15-DaCompany Report #B0133710A  
Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Angioneurotic Oedema		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day							
Initial or Prolonged		Arthralgia		Corticosteroid	SS		
UNKNOWN							
		Tendon Disorder					
		Urticaria					

Date:01/28/02ISR Number: 3860397-8Report Type:Direct Company Report #CTU 160229  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Erythema		Zyban 150mg Tbsr	PS	Tbsr	ORAL
150MG 1 TAB							
		Pruritus					
PO BID							
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/02ISR Number: 3862202-2Report Type:Expedited (15-DaCompany Report #A0172394A

Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety Confusional State Convulsion	Literature Health Professional	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL		Depression Drug Ineffective Insomnia Somnolence		Lithium Salt (Lithium Salt) Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	SS  SS		
150 MG/TWICE							
PER DAY				Electroconvulsive Therapy (Electroconvulsive Therapy)	SS		
SEE DOSAGE							
TEXT				Gabapentin Clonazepam Glycopyrronium Bromide Methohexitone Suxamethonium Oxygen Citalopram	C C C C C C		

Date:01/29/02ISR Number: 3860867-2Report Type:Expedited (15-DaCompany Report #B0101033A

Age:61 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardiac Failure Acute		Zyban	PS	Glaxo Wellcome	ORAL
42 DAY		Myocardial Fibrosis Sudden Cardiac Death		Sotalol Warfarin	C C	Glaxo Wellcome	
UNKNOWN							

Date:01/29/02ISR Number: 3860882-9Report Type:Expedited (15-DaCompany Report #B0134007A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Unknown	2 MON	Headache		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Hypertension Meningism Mental Disorder Musculoskeletal Stiffness Pain Vomiting Projectile					

Date:01/29/02ISR Number: 3860883-0Report Type:Expedited (15-DaCompany Report #B0134016A  
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death UNKNOWN	150MG Unknown	Sudden Death		Zyban	PS	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/29/02ISR Number: 3861794-7Report Type:Direct  
Age:37 YR Gender:Male I/FU:I

Company Report #CTU 160321

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG SR 1	Chills Coordination Abnormal		Wellbutrin 150 Mg Sr Tab Glaxo	PS	Glaxo	ORAL
TABLET BY		Depression					
MOUTH 2 X		Insomnia					
DAILY		Mania					
		Memory Impairment Suicidal Ideation Tremor					

Date:01/29/02ISR Number: 3862106-5Report Type:Direct  
Age:41 YR Gender:Female I/FU:I

Company Report #CTU 160282

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG BID PO		Dizziness Dyspnoea		Zyban (Bupropion) 150 Mg	PS		ORAL
		Headache		Albuterol Inhaler	C		
		Medication Error		Afrin Nasal Spray	C		
		Nausea		Motrin	C		
		Palpitations		Multivitamin	C		

Date:01/29/02ISR Number: 3903414-9Report Type:Periodic  
Age:66 YR Gender:Female I/FU:I

Company Report #A127209

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
250.00 MCG		Anxiety	Consumer	Tikosyn Capsules	PS		ORAL
TOTAL: BID: ORA		Depression					
		Diarrhoea					

225.00 MG Rash Wellbutrin SS  
 Tremor  
 TOTAL:TID Levoxyl C  
 Valium C

Date:01/29/02ISR Number: 3903459-9Report Type:Periodic Company Report #A128334  
 Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression	Consumer	Tikosyn Capsules	PS		ORAL
250.00 MCG		Rash					
TOTAL:PID							

:ORAL Wellbutrin SS ORAL  
 75:00 MG

TOTAL:DAILY:O

RAL Thyroid C

Date:01/30/02ISR Number: 3862121-1Report Type:Expedited (15-DaCompany Report #HQ0174226JAN2002  
 Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Arthralgia	Health	Efexor (Venlafaxine			
Initial or Prolonged		Leukocytoclastic	Professional	Hydrochloride,	PS		ORAL
Other		Vasculitis		Tablet)			
		Myalgia		Zyban (Amfebutamone	SS		
				Hydrochloride, )			
				Unspecified			



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Contraceptive  
(Unspecified  
Contraceptive) C

Date:01/30/02ISR Number: 3863609-XReport Type:Periodic Company Report #PHEH2001US01007  
Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 5 0MG QD, ORAL	2190 DAY	Convulsion Fall Loss Of Consciousness	Health Professional	Ritalin(Methylphenidate Hydrochloride) Tablet, 10mg	PS		ORAL
150 MG, BID, ORAL	30 DAY			Wellbutrin-Slow Release (Amfebutamone Hydrochloride) 150mg	SS		ORAL

Date:01/31/02ISR Number: 3861042-8Report Type:Expedited (15-DaCompany Report #A0174691A  
Age:8 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 450MG per day Initial or Prolonged 100MG Twice per day		Grand Mal Convulsion		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
10MG Per day 2MG Twice per day	6 MON	Infection Personality Change Respiratory Arrest Status Epilepticus		Depakote Celexa Risperdal	C C C		ORAL ORAL ORAL

Date:01/31/02ISR Number: 3861043-XReport Type:Expedited (15-DaCompany Report #B0101363A  
Age:37 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG See Initial or Prolonged dosage text 31 DAY Disability	Balance Disorder Blood Immunoglobulin G Increased Brain Scan Abnormal Coordination Abnormal Demyelination Diplopia Dysaesthesia Dysarthria Monoparesis Multiple Sclerosis Paraesthesia		Zyban	PS	Glaxo Wellcome	ORAL

Date:01/31/02ISR Number: 3861044-1Report Type:Expedited (15-DaCompany Report #B0124304A  
Age:31 YR Gender:Female I/FU:F

Outcome	PT
Life-Threatening Hospitalization - Initial or Prolonged	Abnormal Behaviour Aggression Dry Mouth Dry Skin Dysuria Excitability Heart Rate Increased Insomnia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG As directed	9 DAY	Irritability Logorrhoea Mydriasis Nausea		Zyban	PS	Glaxo Wellcome	ORAL
2700MG Single dose	1 DAY	Nervousness Overdose Somnolence Suicide Attempt Tearfulness Vomiting		Zyban  Alcohol	SS  C	Glaxo Wellcome	ORAL

Date:01/31/02ISR Number: 3861049-0Report Type:Expedited (15-DaCompany Report #B0133422A  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN	Initial or Prolonged	Chest Pain Coordination Abnormal Dizziness Malaise Trigeminal Neuralgia		Zyban	PS	Glaxo Wellcome	

Date:01/31/02ISR Number: 3861056-8Report Type:Expedited (15-DaCompany Report #B0134019A  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Mouth Haemorrhage		Zyban	PS	Glaxo Wellcome	ORAL
150MG As directed				Fluoxetine	SS		ORAL
20MG Per day				Olanzapine	SS		ORAL
7.5MG Unknown				Loxapine	SS		ORAL
50MG Per day							

2UNIT Per day

Date:01/31/02ISR Number: 3861579-1Report Type:Direct Company Report #CTU 160479  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG QD	Alcoholism		Wellbutrin 150mg	PS		ORAL
ORAL		Flushing					
		Paraesthesia Oral Swelling					

Date:01/31/02ISR Number: 3863242-XReport Type:Expedited (15-DaCompany Report #2020164  
 Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Brain Oedema	Health	Morphine Sulfate			
		Contusion	Professional	(Similar To Andas			
MG UNKNOWN PO		Depressed Level Of	Other	74-769 And 74-862)	PS		ORAL
		Consciousness		Oxycodone			
		Overdose		Hydrochloride			
MG UNKNOWN PO		Pulmonary Oedema		(Similar To Nda			
		Toxicologic Test Abnormal		20-553)	SS		ORAL
				Bupropion			
MG UNKNOWN PO				Hydrochloride	SS		ORAL
				Methadone	SS		ORAL
MG UNKNOWN PO				Diazepam	SS		ORAL
MG UNKNOWN PO							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/02ISR Number: 3861658-9Report Type:Expedited (15-DaCompany Report #A0173831A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ageusia Anosmia	Consumer	Zyban	PS	Glaxo Wellcome	ORAL

Date:02/01/02ISR Number: 3861659-0Report Type:Expedited (15-DaCompany Report #A0173931A

Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG Twice		Cholelithiasis	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day		Grand Mal Convulsion	Professional				
				Singulair	C		
				Levoxyl	C	Glaxo Wellcome	
				Prevacid	C		
				Luvox	C		

Date:02/01/02ISR Number: 3861672-3Report Type:Expedited (15-DaCompany Report #B0131127A

Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Chest Pain					
per day		Decreased Appetite		Acetylcysteine	C	Glaxo Wellcome	
UNKNOWN		Depression		Flunisolide	C		
UNKNOWN		Disturbance In Attention		Salbutamol	C	Glaxo Wellcome	
UNKNOWN		Dizziness Dry Mouth		Dexchlorpheniramine Maleate	C		
UNKNOWN		Dyspnoea Headache		Salmeterol + Fluticasone	C	Glaxo Wellcome	

UNKNOWN	Influenza Like Illness	Tenoxicam	C
UNKNOWN	Nausea	Piroxicam	C
	Oedema Peripheral		
	Palpitations		
	Sleep Disorder		

Date:02/01/02ISR Number: 3861679-6Report Type:Expedited (15-DaCompany Report #B0133944A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other							
150MG Unknown			Pulmonary Infarction	Zyban	PS	Glaxo Wellcome	

Date:02/01/02ISR Number: 3861683-8Report Type:Expedited (15-DaCompany Report #B0134202A  
 Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -							
300MG Per day 33 DAY			Abdominal Pain	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged			Flatulence				
			Intestinal Obstruction				
			Vomiting				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/02ISR Number: 3861685-1Report Type:Expedited (15-DaCompany Report #B0134232A  
 Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Unknown	Death		Zyban	PS	Glaxo Wellcome	ORAL
				Oxygen Therapy	C		

Date:02/01/02ISR Number: 3861686-3Report Type:Expedited (15-DaCompany Report #B0134348A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Unknown	Death		Zyban	PS	Glaxo Wellcome	ORAL

Date:02/01/02ISR Number: 3863259-5Report Type:Direct Company Report #CTU 160600  
 Age:47 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Constipation	1T TAB PO BID			Zyban 150mg Tbsr	PS		ORAL
				Habitrol	C		

Date:02/04/02ISR Number: 3862298-8Report Type:Expedited (15-DaCompany Report #B0134351A  
 Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Unknown 31 DAY	Abdominal Pain Upper		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Gastric Ulcer Gastrointestinal Haemorrhage Haematemesis Hiatus Hernia Oesophagitis Haemorrhagic		Alcohol	C		

Date:02/04/02ISR Number: 3862302-7Report Type:Expedited (15-DaCompany Report #B0134762A  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown							

Date:02/05/02ISR Number: 3863191-7Report Type:Expedited (15-DaCompany Report #A0175184A  
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Facial Palsy		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Per day 2 DAY							
				Celexa	C		
				Ziac	C		
				Spironolactone	C		
				Reglan	C	Glaxo Wellcome	
				Zyrtec	C	Glaxo Wellcome	

Date:02/05/02ISR Number: 3863198-XReport Type:Expedited (15-DaCompany Report #B0096692A  
Age:73 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Exanthem
Initial or Prolonged	Hepatic Steatosis
	Hepatomegaly

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Lung Infiltration Petechiae Red Blood Cell Sedimentation Rate Increased Vasculitis		Zyban	PS	Glaxo Wellcome	ORAL

Date:02/05/02ISR Number: 3863199-1Report Type:Expedited (15-DaCompany Report #B0102171A  
Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN		Alanine Aminotransferase DAY Increased Burning Sensation Dermatitis Exfoliative Fatigue Neutrophilia Oedema Peripheral Rash Generalised Rash Maculo-Papular Rash Pruritic Vision Blurred		Zyban	PS	Glaxo Wellcome	

Date:02/05/02ISR Number: 3863208-XReport Type:Expedited (15-DaCompany Report #B0126564A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly UNKNOWN	300MG per day	Asthma Caesarean Section Complications Of Maternal Exposure To Therapeutic Drugs Dermatitis Atopic Maternal Drugs Affecting Foetus Neonatal Disorder Pregnancy		Zyban	PS	Glaxo Wellcome	

Ventricular Septal Defect  
Acquired

Date:02/05/02ISR Number: 3863221-2Report Type:Expedited (15-DaCompany Report #B0135045A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Head Injury		Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown							

Date:02/05/02ISR Number: 3863225-XReport Type:Expedited (15-DaCompany Report #B0135166A  
Age:36 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Alanine Aminotransferase Increased Arthralgia Aspartate Aminotransferase Increased Blood Bilirubin Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Leukocytosis Neutrophil Count Increased	Report Source	Product	Role	Manufacturer	Route
150MG	Unknown	Pyrexia		Zyban	PS	Glaxo Wellcome	ORAL
		Urticaria					

Date:02/05/02ISR Number: 3865355-5Report Type:Expedited (15-DaCompany Report #A0174500A  
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Facial Palsy Lagophthalmos	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER							
DAY / ORAL							

Date:02/05/02ISR Number: 3865911-4Report Type:Expedited (15-DaCompany Report #A0159165A  
Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Deafness Depression Drug Interaction Liver Function Test	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
SEE DOSAGE		Abnormal					
TEXT/ORAL		Loss Of Consciousness Memory Impairment Scleral Discolouration		Metoprolol Succinate (Metoprolol Succinate)	SS		ORAL
50 MG PER		Tinnitus					
DAY/ORAL		Tooth Disorder Tooth Loss		Nitroglycerin Ranitidine Hydrochloride Hydroxyzine	C C C		

Date:02/05/02ISR Number: 3866408-8Report Type:Expedited (15-DaCompany Report #A0140463A  
Age:27 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia Cardiac Arrest Confusional State	Foreign Literature Health	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL	10 DAY	Euphoric Mood Feeling Abnormal Hypersensitivity Memory Impairment Myocardial Infarction Palpitations Psychomotor Hyperactivity Pyrexia Speech Disorder	Professional				

Date:02/05/02ISR Number: 3866618-XReport Type:Expedited (15-DaCompany Report #B0111327A  
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cardiomyopathy	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

150 MG / ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/05/02ISR Number: 3866620-8Report Type:Expedited (15-DaCompany Report #A0170371A  
 Age:43 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/ PER DAY / ORAL	Chills Convulsion Mental Disorder Pyrexia Speech Disorder Weight Decreased	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
			Varicella Virus Vaccine (Formulation Ukn) (Varicella Virus Vaccine)	SS		

Date:02/05/02ISR Number: 3866622-1Report Type:Expedited (15-DaCompany Report #B0134015A  
 Age:68 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 150 MG	Myocardial Infarction	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		

Date:02/05/02ISR Number: 3866682-8Report Type:Expedited (15-DaCompany Report #B0131464A  
 Age:25 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG ORAL	Angina Pectoris Blood Creatine Increased Blood Pressure Decreased Cardiac Failure Cutaneous Vasculitis Ecchymosis Hyperpyrexia Malaise Myocardial Infarction Toxicologic Test Abnormal	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
			Methylenedioxymetham phet (Methylenedioxymetha mphet. ) Ethanol Cannabis	SS C C		

Date:02/07/02ISR Number: 3866464-7Report Type:Direct Company Report #CTU 161205  
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased		Wellbutrin	PS		ORAL
150 MG 2		Stress					
TABLET TWICE							
A DAY ORAL							

Date:02/07/02ISR Number: 3866949-3Report Type:Expedited (15-DaCompany Report #001-0945-M0200153  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Asthma	Health	Neurontin			
Initial or Prolonged		Blood Glucose Fluctuation	Professional	(Gabapentin)	PS		ORAL
1500 MG (QID)							
Other		Hypoglycaemia					
PER ORAL		Incontinence		(Bupropion)	SS		ORAL
450 MG (TID)							
PER ORAL							
				(Advair)	C		
				(Prednisone)	C		
				(Diazepam)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/12/02ISR Number: 3866922-5Report Type:Expedited (15-DaCompany Report #A0175186A  
 Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dehydration		Wellbutrin	PS	Glaxo Wellcome	ORAL
200MG Twice		Grand Mal Convulsion					
per day		Stress		Zoloft	C		
UNKNOWN							

Date:02/13/02ISR Number: 3867728-3Report Type:Expedited (15-DaCompany Report #B0100348A  
 Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Zyban	PS	Glaxo Wellcome	ORAL
150MG Three		Confusional State					
times per day 9	DAY	Hallucination		Catapressan	C		
		Hallucination, Visual		Penicillin	C	Glaxo Wellcome	
		Psychotic Disorder		Netromycin			
		Restlessness		(Netilmicin)	C		
				Haloperidol	C		
				Morphine Chloride	C		

Date:02/13/02ISR Number: 3867733-7Report Type:Expedited (15-DaCompany Report #B0134349A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death		Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown							

Date:02/13/02ISR Number: 3867734-9Report Type:Expedited (15-DaCompany Report #B0134350A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death Death Zyban PS Glaxo Wellcome  
UNKNOWN 150MG Unknown

Date:02/13/02ISR Number: 3869419-1Report Type:Periodic Company Report #2013325  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553) Bupropion Hydrochloride Effexor (Venlafaxine) Diazepam Acetaminophen Ethanol (Alcohol, Anhydrous)	PS   SS SS SS SS		

Date:02/13/02ISR Number: 3869517-2Report Type:Periodic Company Report #2013220  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda			



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

20-553) PS  
 Alprazolam SS  
 Diazepam SS  
 Bupropion  
 Hydrochloride SS

Date:02/13/02ISR Number: 3869518-4Report Type:Periodic  
 Age:37 YR Gender:Male I/FU:I

Company Report #2013221

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse	Health Professional Other	Oxycontin Cr Tablets, 10 Mg (Oxycodone Hydrochloride)	PS		ORAL
PO				Alprazolam	SS		
				Bupropion			
				Hydrochloride	SS		
				Cyclobenzaprine Hcl	SS		
				Paroxetine Hcl	SS		

Date:02/13/02ISR Number: 3869590-1Report Type:Periodic  
 Age:37 YR Gender:Male I/FU:I

Company Report #2012330

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553)	PS		
				Hydrocodone Bitartrate	SS		
				Diazepam	SS		
				Bupropion	SS		
				Propoxyphene	SS		
				Diclofenac Sodium	SS		
				Caffeine	SS		
				Nicotine	SS		

Date:02/13/02ISR Number: 3870599-2Report Type:Periodic  
 Age:41 YR Gender:Female I/FU:I

Company Report #2013430

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Accidental Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553)	PS		
				Hydrocodone Bitartrate	SS		
				Chlorpheniramine	SS		
				Cyclobenzaprine	SS		
				Bupropion	SS		
				Propoxyphene	SS		
				Lidocaine	SS		
				Caffeine	SS		
				Nicotine	SS		
				Diazepam	SS		
				Oxazepam	SS		
				Temazepam	SS		
				Lorazepam	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/13/02ISR Number: 3871103-5Report Type:Periodic  
 Age:29 YR Gender:Male I/FU:I

Company Report #2015056

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional Other	Oxycodone Hydrochloride Morphine Sulfate (Similar To Nda 19-516) Codeine Olanzapine Acetaminophen Paroxetine Bupropion Nicotine	PS    SS SS SS SS SS SS		

Date:02/13/02ISR Number: 3872392-3Report Type:Periodic  
 Age:59 YR Gender:Female I/FU:I

Company Report #2014089

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553) Ethyl Alcohol Bupropion Diazepam Acetaminophen Ibuprofen	PS   SS SS SS SS SS		

Date:02/13/02ISR Number: 3875457-5Report Type:Periodic  
 Age:38 YR Gender:Male I/FU:I

Company Report #2012980

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553) Bupropion Hydrochloride	PS   SS		

Nicotine

SS

Date:02/13/02ISR Number: 3875463-0Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #2012972

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553)	PS		
				Propoxyphene	SS		
				Acetaminophen	SS		
				Bupropion	SS		
				Diazepam	SS		
				Trazodone	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/13/02ISR Number: 3876144-XReport Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #2014807

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Dependence	Health	Oxycodone			
Other		Intentional Misuse	Professional	Hydrochloride			
			Other	(Similar To Nda 20-553)	PS		
				Bupropion			
				Hydrochloride	SS		

Date:02/13/02ISR Number: 3876166-9Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #2013012

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health	Oxycodone			
			Professional	Hydrochloride			
			Other	(Similar To Nda 20-553)	PS		
				Ethanol (Alcohol, Anhydrous)	SS		
				Temazepam	SS		
				Acetaminophen	SS		
				Propoxyphene Hcl	SS		
				Bupropion			
				Hydrochloride	SS		
				Nicotine	SS		
				Caffeine Anhydride	SS		

Date:02/13/02ISR Number: 3876676-4Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #2012177

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health	Oxycodone			
			Professional	Hydrochloride	PS		ORAL
MG PO			Other	Hydrocodone			
				Bitartrate	SS		
				Bupropion	SS		ORAL

MG PO

MG PO	Meprobamate	SS	ORAL
MG PO	Diazepam	SS	ORAL
MG PO	Oxazepam	SS	ORAL
MG PO	Lorazepam	SS	ORAL
MG PO	Trazodone	SS	ORAL

Date:02/13/02ISR Number: 3876797-6Report Type:Periodic Company Report #2012673  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553)	PS		
				Bupropion	SS		
				Doxepin	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/13/02ISR Number: 3877723-6Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #2013135

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553)	PS		ORAL
MG, PO				Amitriptyline	SS		
				Bupropion			
				Hydrochloride	SS		
				Acetazolamide	C		
				Seroquel	C		
				Clonazepam	C		
				Nefazodone	C		
				Naprozen	C		
				Claritin (Loratadine)	C		
				Biaxin	C		

Date:02/13/02ISR Number: 3877736-4Report Type:Periodic  
Age:35 YR Gender:Male I/FU:I

Company Report #2013040

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health	Oxycontin Cr Tablets	PS		
MG			Professional Other	Diazepam	SS		
				Bupropion	SS		
				Fluconazole	SS		
				Oxazepam	SS		
				Temazepam	SS		

Date:02/13/02ISR Number: 3877738-8Report Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #2013023

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse	Health Professional Other	Oxycodone Hydrochloride	PS		
				Bupropion	SS		

Meperidine	SS
Acetaminophen	SS
Ketamine	
Hydrochloride	SS
Ritalin	
(Methylphenidate)	SS

Date:02/13/02ISR Number: 3878322-2Report Type:Periodic  
 Age:41 YR Gender:Male I/FU:I

Company Report #2013076

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycodone Hydrochloride Welbutrin (Bupropion) Dalmane (Flurazepam) Ethyl Alcohol Salicylates			
					PS		
					SS		
					SS		
					SS		
					SS		

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Freedom Of Information (FOI) Report

Date:02/13/02ISR Number: 3878382-9Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #2013697

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycontin Cr Tablets, 40 Mg (Oxycodone Hydrochloride)	PS		ORAL
40 MG, PO				Wellbutrin (Bupropion)	SS		
MG				Amitriptyline	SS		
MG				Alprazolam	SS		
MG				Alcohol	SS		

Date:02/14/02ISR Number: 3868750-3Report Type:Expedited (15-DaCompany Report #A0359101A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Respiratory Disorder		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:02/14/02ISR Number: 3868759-XReport Type:Expedited (15-DaCompany Report #B0132587A  
Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 8 DAY Initial or Prolonged		Aneurysm Ruptured		Zyban	PS	Glaxo Wellcome	ORAL
		Brain Scan Abnormal					
		Headache					
		Hydrocephalus					
		Hypertension					
		Loss Of Consciousness					
		Meningorrhagia					
		Musculoskeletal Stiffness					
		Nervous System Disorder					
		Osteoma					

Vomiting

Date:02/14/02ISR Number: 3868761-8Report Type:Expedited (15-DaCompany Report #B0135181A  
Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Unknown		Cardioactive Drug Level		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged INTRAVENOUS	1000IU	Increased per		Human Insulin	SS		
day	1	DAY	Clonic Convulsion				
22.5MG Per		Dyskinesia		Zopiclone	SS		ORAL
day	1	DAY	Feeling Abnormal				
1	DAY		Hypoglycaemic Coma	Digoxin	SS	Glaxo Wellcome	ORAL
		Overdose Tremor					

Date:02/14/02ISR Number: 3869747-XReport Type:Direct Company Report #CTU 161637  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Apnoea Grand Mal Convulsion		Wellbutrin Sr (Bupropion) 150 Mg Tab	PS		ORAL
150 MG PO BID				Prozac	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/02ISR Number: 3870033-2Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #CTU 161647

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Crying		Paxil 60mg			
		Depression		Glaxosmithkline	PS	Glaxosmithkline	ORAL
30MG TWICE							
DAILY ORAL		Diarrhoea					
		Dizziness		Welbutrin 125mg			
		Drug Withdrawal Syndrome		Glaxowellcome	SS	Glaxowellcome	ORAL
125MG TWICE							
DAILY ORAL		Ear Disorder					
		Feeling Abnormal					
		Headache					
		Influenza Like Illness					
		Nausea					
		Night Sweats					
		Nightmare					
		Paraesthesia					
		Suicidal Ideation					

Date:02/14/02ISR Number: 3870459-7Report Type:Expedited (15-DaCompany Report #A202614  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Blood Testosterone	Health	Zoloft Tablets	PS		
Intervention to		Decreased	Professional	Neurontin	SS		
Prevent Permanent		Hyponatraemia		Paxil	SS		
Impairment/Damage				Wellbutrin	SS		
				Ambien	C		
				Albuterol Inhaler	C		
				Atenolol	C		
				Folic Acid	C		
				Zocor	C		
				Hydrochlorothiazide	C		

Date:02/14/02ISR Number: 3870486-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200061  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Testosterone	Health	Neurontin			
Other		Decreased	Professional	(Gabapentin)	PS		
		Hyponatraemia		Sertraline			
				Hydrochloride	SS		
				Bupropion	SS		
				Paroxetine			
				Hydrochloride	SS		
				Zolpidem Tartrate	C		
				Salbutamol	C		
				Atenolol	C		
				Folic Acid	C		
				Simvastatin	C		
				Hydrochlorothiazide	C		

Date:02/15/02ISR Number: 3869246-5Report Type:Expedited (15-DaCompany Report #B0117368A  
Age:55 YR Gender:Male I/FU:F

Outcome PT  
Death Bronchial Carcinoma  
Cardiac Failure Acute  
Coronary Artery

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG As directed		Atherosclerosis Metastases To Lymph Nodes Myocardial Ischaemia Sudden Death	Zyban	PS	Glaxo Wellcome	ORAL
25MG per day			Captopril	C	Glaxo Wellcome	
40MG per day			Furosemide	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)			Salbutamol	C	Glaxo Wellcome	
			Fluticasone	C	Glaxo Wellcome	

Date:02/15/02ISR Number: 3869254-4Report Type:Expedited (15-DaCompany Report #B0135382A  
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 2 Initial or Prolonged Disability	2 WK	Dehydration		Zyban	PS	Glaxo Wellcome	ORAL
		Dysuria Eosinophil Count Increased Exanthem Mucosal Erosion Oedema Peripheral Pruritus Pyrexia Skin Exfoliation Toxic Skin Eruption		Nomegestrol	SS		ORAL

Date:02/15/02ISR Number: 3869257-XReport Type:Expedited (15-DaCompany Report #B0259229A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 15000MG		Overdose		Zyban	PS	Glaxo Wellcome	ORAL

Single dose

Date:02/15/02ISR Number: 3870837-6Report Type:Direct Company Report #CTU 161754  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS		ORAL
200MG PO BID				Lithium Carbonate	C		

Date:02/18/02ISR Number: 3869787-0Report Type:Expedited (15-DaCompany Report #A0173850A  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxo Wellcome	ORAL
Other		Cyst		Celexa	C		ORAL
100MG Per day							

Date:02/18/02ISR Number: 3869788-2Report Type:Expedited (15-DaCompany Report #A0175712A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Complex Partial Seizures		Wellbutrin	PS	Glaxo Wellcome	ORAL
Other		Convulsion		Fastin	SS	Glaxo Wellcome	
150MG Unknown		Drug Interaction		Vistaril	C		
30MG Per day							
50MG Twice							
per day							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

10MG At night	Ambien	C	
	Talwin	C	
	Imitrex	C	Glaxo Wellcome
	Phenergan	C	Glaxo Wellcome

Date:02/18/02ISR Number: 3869803-6Report Type:Expedited (15-DaCompany Report #B0259361A  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Unknown 21 DAY Initial or Prolonged		Psoriasis		Zyban	PS	Glaxo Wellcome	ORAL

Date:02/18/02ISR Number: 3869804-8Report Type:Expedited (15-DaCompany Report #B0259370A  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Per day 7 DAY		Psoriasis		Zyban	PS	Glaxo Wellcome	ORAL
250MG Three times per day				Naproxen	C		ORAL
1MG At night				Lorazepam	C		ORAL
60MG As required				Dihydrocodeine	C		ORAL
25MG As required				Hydroxyzine	C		ORAL

Date:02/18/02ISR Number: 3869805-XReport Type:Expedited (15-DaCompany Report #B0259495A  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Face Oedema		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Urticaria					
per day	31	DAY					
TOPICAL		Wheezing		Zovirax	SS	Glaxo Wellcome	
				Aspirin	C		ORAL
75MG Per day							

Date:02/20/02ISR Number: 3870925-4Report Type:Expedited (15-DaCompany Report #A0359237A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Aspiration		Wellbutrin	PS	Glaxo Wellcome	ORAL
300MG per day		Convulsion		Depakote	C		
Other		Medication Error					
		Overdose					

Date:02/20/02ISR Number: 3870927-8Report Type:Expedited (15-DaCompany Report #A0359335A  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT
Hospitalization -		Blindness
Initial or Prolonged		Coma
		Depression
		Disturbance In Attention
		Epileptic Aura
		Generalised
		Non-Convulsive Epilepsy
		Memory Impairment
		Optic Nerve Disorder



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Visual Acuity Reduced

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	YR		Wellbutrin	PS	Glaxo Wellcome	ORAL
200MG As required			Oxycontin Flexeril Acyclovir	C C C	Glaxo Wellcome	

Date:02/20/02ISR Number: 3870928-XReport Type:Expedited (15-DaCompany Report #A0359405A  
Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Condition Aggravated Fall Gait Disturbance Hip Fracture		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:02/20/02ISR Number: 3870935-7Report Type:Expedited (15-DaCompany Report #B0259584A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Unknown 1 Initial or Prolonged	1 WK	Alanine Aminotransferase Increased Blood Lactate Dehydrogenase Increased Condition Aggravated Hypertension Pancreatitis Tachycardia White Blood Cell Count Increased		Zyban Pancrelipase	PS SS	Glaxo Wellcome	ORAL ORAL

Date:02/20/02ISR Number: 3870936-9Report Type:Expedited (15-DaCompany Report #B0259585A  
Age:35 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Per day	Anxiety		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Confusional State Disturbance In Attention Drug Interaction Muscle Spasms Tachycardia		Proguanil Chloroquine	SS SS		ORAL ORAL

Date:02/20/02ISR Number: 3870937-0Report Type:Expedited (15-DaCompany Report #B0259831A  
Age:31 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG See	Chest Pain		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged dosage text 9 DAY	Electrocardiogram Repolarisation Abnormality					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/20/02ISR Number: 3872606-XReport Type:Direct  
Age:50 YR Gender:Female I/FU:I

Company Report #CTU 161881

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG		Arthralgia		Wellbutrin Sr	PS		
		Blood Pressure Increased Chronic Fatigue Syndrome Disturbance In Attention Fibromyalgia Insomnia Nervousness		Zyban	SS		

Date:02/20/02ISR Number: 3873029-XReport Type:Expedited (15-DaCompany Report #A202713  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 100.00 MG		Dry Skin	Consumer	Zoloft Tablets	PS		ORAL
Intervention to TOTAL:DAILY:0 Prevent Permanent RAL Impairment/Damage 50.00 MG		Energy Increased Fatigue		Amitriptyline	SS		ORAL
TOTAL:DAILY:0 RAL		Hernia Hunger Nausea					
		Night Sweats Paranoia Pollakiuria Productive Cough Sleep Disorder Tooth Abscess Tremor Viral Infection Weight Decreased		Welbutrin Tricor Lipitor Vitamin Unspecified Thioridazine	SS C C C C		

Date:02/21/02ISR Number: 3871529-XReport Type:Expedited (15-DaCompany Report #B0259698A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day		Amnesia		Alcohol	C		
		Confusional State					
		Disturbance In Attention					
		Dizziness					
		Loss Of Consciousness					
		Malaise					
		Postictal State					
		Road Traffic Accident					

Date:02/21/02ISR Number: 3871530-6Report Type:Expedited (15-DaCompany Report #B0259830A  
Age:60 YR Gender:Male I/FU:F

Outcome	PT
Life-Threatening	Asthenia
	Cyanosis
	Diabetes Mellitus
	Dyspnoea
	Hyperhidrosis
	Hypoglycaemic Coma
	Oxygen Saturation
	Decreased
	Pallor

Freedom Of Information (FOI) Report

Tachypnoea

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	6 DAY		Zyban	PS	Glaxo Wellcome	ORAL
3 YR			Metformin	SS		ORAL
3 YR			Glimepiride	SS		ORAL
5MG per day			Tramadol	C		
			Bisoprolol	C		ORAL
			Paracetamol	C	Glaxo Wellcome	
			Meprobamate	C		
			Buflomedil	C		
3 YR			Lysine Aspirin	C		ORAL
			Lansoprazole	C		
			Frusemide	C	Glaxo Wellcome	
			Ciprofibrate	C		
			Pepsane	C		
			Clorazepate	C		
1.25MG Per			Ramipril	C		ORAL
day						
			Vit B1 B6	C		

Date:02/21/02ISR Number: 3871531-8Report Type:Expedited (15-DaCompany Report #B0260146A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Dysgeusia		Zyban	PS	Glaxo Wellcome	ORAL
2TAB Twice							
Initial or Prolonged		Medication Error					
per day	10 DAY						
		Sputum Abnormal					

Date:02/21/02ISR Number: 3872709-XReport Type:Direct  
 Age:46 YR Gender:Female I/FU:I

Company Report #CTU 161965

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Convulsion		Wellbutrin Sr Blue			
Hospitalization -		Grand Mal Convulsion		Pill	PS		ORAL
1 PILL DAILY							
Initial or Prolonged							
ORAL							
Disability							
Required							
Intervention to							
Prevent Permanent							
Impairment/Damage							

Date:02/21/02ISR Number: 3872747-7Report Type:Direct  
Age:35 YR Gender:Female I/FU:I

Company Report #CTU 162017

Outcome	PT
Other	Agitation
	Balance Disorder
	Dizziness
	Dyspnoea
	Feeling Hot And Cold
	Flushing
	Influenza
	Insomnia
	Lower Respiratory Tract
	Infection
	Mood Swings
	Muscle Spasms

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG X 2/DAY		Nausea Paraesthesia Rash Pruritic Throat Tightness Tinnitus Urticaria Vision Blurred		Wellbutrin Sr 150 Mg X 2 Day = 300mg	PS		

Date:02/21/02ISR Number: 3873305-0Report Type:Expedited (15-DaCompany Report #B0259331A  
Age:32 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	TRANSDERMAL	14 MG/PER DAY/	Anxiety Convulsion Loss Of Consciousness Tongue Biting	Foreign Health Professional	Nicoderm Cq Patch 14 Mg (Nicotine)	PS		
	TRANSDERMAL		Urinary Incontinence		Zyban Tablet-Zyban (Bupropion Hydrochloride)	SS		ORAL

Date:02/22/02ISR Number: 3871912-2Report Type:Expedited (15-DaCompany Report #A0132245A  
Age:52 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged	150MG Twice per day	1 MON	Blindness Transient Convulsion	Health Professional	Zyban	PS	Glaxo Wellcome	ORAL
	150MG Twice per day		Deafness Dizziness		Zyban	SS	Glaxo Wellcome	ORAL
			Hyperglycaemia Hypoglycaemia Nervous System Disorder		Multivitamin Vitamin E Vitamin C	C C C		

Tinnitus

Aspirin

C

Date:02/22/02ISR Number: 3871915-8Report Type:Expedited (15-DaCompany Report #A0359236A  
Age:28 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Optic Nerve Disorder	Consumer	Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	41 DAY	Vision Blurred		No Concurrent Medications	C		

Date:02/22/02ISR Number: 3871916-XReport Type:Expedited (15-DaCompany Report #A0359405A  
Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated Fall Gait Disturbance	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome	ORAL

Date:02/22/02ISR Number: 3871921-3Report Type:Expedited (15-DaCompany Report #B0124405A  
Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Anaphylactic Shock	Health	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	17 DAY	Angina Pectoris	Professional				



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

INTRAMUSCULAR				Hepatyrix	C	Glaxo Wellcome	
				Ciprofloxacin	C		ORAL
				Norethisterone	C		ORAL
350UG Per day							

Date:02/22/02ISR Number: 3871923-7Report Type:Expedited (15-DaCompany Report #B0134062A  
 Age:24 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebral Artery Occlusion	Health	Quomem	PS	Glaxo Wellcome	ORAL
150MG As		Convulsion	Professional				
directed	29 DAY	Nuclear Magnetic		Diazepam	SS		
UNKNOWN	2MG per day	Resonance Imaging Brain		Alcohol	C		
		Abnormal					
		Photopsia					

Date:02/22/02ISR Number: 3871928-6Report Type:Expedited (15-DaCompany Report #B0259603A  
 Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Acne	Health	Zyban	PS	Glaxo Wellcome	
150MG Per day	6 DAY	Dysgeusia	Professional				
		Facial Pain					
		Facial Palsy					
		Pain					
		Sleep Disorder					
		Tongue Ulceration					

Date:02/22/02ISR Number: 3871929-8Report Type:Expedited (15-DaCompany Report #B0259722A  
 Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other			Hallucination, Auditory	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
300MG Unknown	10	DAY	Suicidal Ideation		Paracetamol	C	Glaxo Wellcome	ORAL
1000MG As								

required

Date:02/22/02ISR Number: 3871931-6Report Type:Expedited (15-DaCompany Report #B0259958A  
 Age:50 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Hypothyroidism		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	73	DAY	Oedema		Salbutamol	C	Glaxo Wellcome	
RESPIRATORY			Weight Increased					
(INHALATION)								

Date:02/22/02ISR Number: 3871932-8Report Type:Expedited (15-DaCompany Report #B0260109A  
 Age:22 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Urticaria		Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown	2	WK	Wheezing					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/22/02ISR Number: 3871933-XReport Type:Expedited (15-DaCompany Report #B0260117A  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Panic Attack		Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown		Rash					

Date:02/22/02ISR Number: 3871934-1Report Type:Expedited (15-DaCompany Report #B0260178A  
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination, Auditory		Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown	22 DAY						

Date:02/22/02ISR Number: 3875272-2Report Type:Expedited (15-DaCompany Report #200211202GDDC  
 Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Myocardial Infarction	Foreign Health	Indapamide (Dapa-Tabs) Tablets	PS		ORAL
2.5 MG PO	1 YR		Professional Other	Amfebutamone Hydrochloride (Zyban)	SS		
8 DAY				Quinapril Hydrochloride (Accupril)	C		
				Acetylsalicylate Calcium (Solprin)	C		

Date:02/22/02ISR Number: 3879462-4Report Type:Periodic Company Report #HQ2442222JUN2001  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Effexor (Venlafaxine)			

ORAL

Hydrochloride, Unspec)	PS	ORAL
Haldol (Haloperidol, )	SS	
Remeron (Mirtazapine, )	SS	
Wellbutrin (Amfebutamone Hydrochloride, )	SS	

Date:02/25/02ISR Number: 3873437-7Report Type:Direct  
Age:67 YR Gender:Female I/FU:I

Company Report #CTU 162124

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG 1T PO		Blood Pressure Increased		Zyban (150mg) Tbsr	PS		ORAL

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Date:02/25/02ISR Number: 3874220-9Report Type:Expedited (15-DaCompany Report #B0130905A  
Age:47 YR Gender:Male I/FU:I

Outcome Disability	PT Ageusia Dermatitis Bullous
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ TWICE PER DAY/ORAL		Difficulty In Walking Erythema Glossitis Oropharyngeal Swelling Pruritus Rash Vesicular Skin Exfoliation	Foreign Health Professional	Zyban Tablet - Zyban(Bupropion Hydrochloride)  Ketoprofen	PS  C		ORAL

Date:02/25/02ISR Number: 3874230-1Report Type:Expedited (15-DaCompany Report #B0259252A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG/ PER DAY/ ORAL		C-Reactive Protein Increased Cheilitis Dermatitis Bullous	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL		Hypopyon Neutrophilia Pyrexia		Propofan (Formulation Unknown) (Propofan)	SS		ORAL
		Rash Erythematous Rash Pustular		Prazepam Zolpidem Alprazolam	C C C		

Date:02/25/02ISR Number: 3874252-0Report Type:Expedited (15-DaCompany Report #B0259334A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG / TWICE PER DAY /		Dyspnoea Hyperventilation Muscle Spasms	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS	Zyban	ORAL

ORAL

Date:02/25/02ISR Number: 3874253-2Report Type:Expedited (15-DaCompany Report #B0133422A

Age:33 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG / PER DAY / ORAL	Chest Pain Coordination Abnormal Dizziness Hypoaesthesia Malaise Trigeminal Neuralgia Wheezing	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)  Oral Contraceptive	PS  C	Zyban	ORAL

Date:02/25/02ISR Number: 3874313-6Report Type:Expedited (15-DaCompany Report #B0129422A

Age:48 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - Initial or Prolonged 150 MG	Duodenal Ulcer Gastric Haemorrhage Haematocrit Decreased Irritability Malaise Mood Altered	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/02ISR Number: 3873670-4Report Type:Expedited (15-DaCompany Report #A0167897A  
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Treatment Noncompliance		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:02/26/02ISR Number: 3873675-3Report Type:Expedited (15-DaCompany Report #A0359984A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:02/26/02ISR Number: 3873687-XReport Type:Expedited (15-DaCompany Report #D0037982A  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Other		Deafness Neurosensory Intentional Misuse Suicide Attempt Tinnitus Vestibular Disorder		Zyban	PS	Glaxo Wellcome	ORAL

Date:02/26/02ISR Number: 3873688-1Report Type:Expedited (15-DaCompany Report #D0038006A  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Unknown		Sudden Cardiac Death		Zyban	PS	Glaxo Wellcome	ORAL

Date:02/27/02ISR Number: 3874199-XReport Type:Expedited (15-DaCompany Report #A0359318A  
Age:13 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death Convulsion Wellbutrin PS Glaxo Wellcome ORAL  
9 MON  
Hospitalization - Medication Error  
Initial or Prolonged Overdose

Date:02/27/02ISR Number: 3876517-5Report Type:Expedited (15-DaCompany Report #EMADSS2002001073  
Age:18 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - Initial or Prolonged 1.3 G, 1IN 1 Disability TIME (S), Required ORAL	Cardiac Arrest Cerebral Ischaemia Overdose  Status Epilepticus	Foreign Health Professional	Zydol (Tablet) (Tramadol Hydrochloride)	PS		ORAL
Intervention to Prevent Permanent 9 G, 1IN 1 Impairment/Damage TIME (S),  ORAL			Zyban (Amfebutamone Hydrochloride)	SS		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/02ISR Number: 3875251-5Report Type:Expedited (15-DaCompany Report #A0174691A

Age:8 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	450MG per day	19 MON	Health	Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
Hospitalization -	1000MG Twice		Professional	Depakote	C		ORAL
Initial or Prolonged	per day		Drug Level Increased				
Other	10MG Per day	6 MON	Grand Mal Convulsion	Celexa	C		ORAL
	2MG Twice per		Infection	Risperdal	C		ORAL
	day		Personality Change				
			Pyrexia				
			Respiratory Arrest				
			Status Epilepticus				

Date:02/28/02ISR Number: 3875252-7Report Type:Expedited (15-DaCompany Report #A0175021A

Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Twice		Health	Zyban	PS	Glaxo Wellcome	ORAL
	per day	86 DAY	Professional	Effexor	C		

Date:02/28/02ISR Number: 3875254-0Report Type:Expedited (15-DaCompany Report #A0175712A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Unknown		Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
	30MG Per day		Convulsion	Fastin	SS	Glaxo Wellcome	
	50MG Twice		Drug Interaction	Vistaril	C		

per day

10MG At night

Ambien	C		
Talwin	C		
Imitrex	C	Glaxo Wellcome	
Phenergan	C	Glaxo Wellcome	
Prozac	C		ORAL

Date:02/28/02ISR Number: 3875256-4Report Type:Expedited (15-DaCompany Report #A0360169A

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Overdose	Health Professional	Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:02/28/02ISR Number: 3875258-8Report Type:Expedited (15-DaCompany Report #B0126006A

Age: Gender:Unknown I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Myocardial Infarction	Health Professional	Zyban	PS	Glaxo Wellcome	

Date:02/28/02ISR Number: 3875264-3Report Type:Expedited (15-DaCompany Report #B0260668A

Age:31 YR Gender:Female I/FU:I

Outcome	PT
Other	Drug Interaction
	Headache
	Ill-Defined Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Malaise				
		Nausea				
		Tachycardia	Report Source	Product	Role	Manufacturer
Dose	Duration		Consumer	Zyntabac	PS	Glaxo Wellcome
300MG Per day	9 DAY			Flu Medication	SS	
4	DAY			Ethinylloestradiol	SS	
						Route
						ORAL
						ORAL
						ORAL

Date:02/28/02ISR Number: 3875265-5Report Type:Expedited (15-DaCompany Report #D0038028A  
 Age:31 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration		Health	Zyban	PS	Glaxo Wellcome	ORAL
Dose		Fear					
Life-Threatening			Professional				
7.5G per day	1 DAY	Intentional Misuse					
Other		Suicide Attempt					
		Tachycardia					
		Tremor					
		Visual Disturbance					
		Vomiting					

Date:02/28/02ISR Number: 3877113-6Report Type:Direct Company Report #CTU 162543  
 Age: Gender: I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration			Zyban 1 A Day	PS		ORAL
Dose		Abdominal Pain					
Life-Threatening							
1 A DAY PO		Alanine Aminotransferase		Prednisone	C		
		Increased		Solumedrol	C		
		Arthralgia		Hydroxyzine	C		
		Aspartate					
		Aminotransferase					
		Increased					
		Red Blood Cell					
		Sedimentation Rate					
		Increased					
		Subarachnoid Haemorrhage					
		Urticaria					

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG / ORAL	Agitation Aspiration Blood Pressure Increased  Clonic Convulsion Coma Convulsion Disorientation Dystonia Extrapyramidal Disorder Hyperreflexia Hypertonia Hypotension Overdose Tachycardia Tremor	Foreign Literature Health  Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)  Ethanol	PS  C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/02ISR Number: 3875966-9Report Type:Expedited (15-DaCompany Report #A0360184A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coma		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged		Convulsion					

Date:03/01/02ISR Number: 3875973-6Report Type:Expedited (15-DaCompany Report #B0259906A

Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 2 DAY Initial or Prolonged		Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
1%DS Per day		Fall		Atacand	C		ORAL
		Malaise Meningioma					

Date:03/01/02ISR Number: 3875974-8Report Type:Expedited (15-DaCompany Report #B0260007A

Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bronchospasm Pharyngolaryngeal Pain Respiratory Disorder		Zyban	PS	Glaxo Wellcome	ORAL

Date:03/01/02ISR Number: 3875977-3Report Type:Expedited (15-DaCompany Report #B0260166A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged RESPIRATORY (INHALATION)		Delirium Loss Of Consciousness		Zyban Illicit Drug(S)	PS C	Glaxo Wellcome	ORAL
		Medication Error Memory Impairment					

Date:03/01/02ISR Number: 3878242-3Report Type:Direct  
Age:46 YR Gender:Female I/FU:I

Company Report #CTU 162612

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Abdominal Pain		Wellbutrin (Generic			
Other		Diarrhoea		Form)	PS		
75 MG	BID						
		Pharmaceutical Product					
		Complaint					

Date:03/01/02ISR Number: 3878249-6Report Type:Direct  
Age:37 YR Gender:Male I/FU:I

Company Report #CTU 162615

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anger		Zyban (150 Mg) Tbsr	PS	150 Mg 1 Tab Qd X	
				Nicorette	C	3 Then Bid	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/02ISR Number: 3878807-9Report Type:Expedited (15-DaCompany Report #A204048

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 50.00 MG	Anxiety	Health	Zoloft Tablets	PS		ORAL
Initial or Prolonged TOTAL:DAILY:0	Depression	Professional				
Required RAL	Diabetes Mellitus	Company				
Intervention to 300.00 MG	Drug Effect Decreased	Representative	Wellbutrin	SS		ORAL
Prevent Permanent TOTAL:PID:ORA	Fatigue					
Impairment/Damage L	Fear					
10.00	Gait Disturbance		Zyprexa	SS		ORAL
TOTAL:DAILY:0	Hallucination, Visual					
RAL	Hemianopia					
	Insomnia					

Date:03/05/02ISR Number: 3877376-7Report Type:Expedited (15-DaCompany Report #A0171269A

Age:48 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150MG Twice	Pneumonitis	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
per day	Pulmonary Fibrosis	Professional				
3 YR			Celexa	C		
UNKNOWN	20MG Per day					

Date:03/05/02ISR Number: 3877379-2Report Type:Expedited (15-DaCompany Report #A0359960A

Age:64 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Hospitalization - 2 WK Initial or Prolonged	Coronary Artery Disease Disorientation Dyspnoea Hallucination, Visual Nervousness	Zyban No Concurrent Medication	PS C	Glaxo Wellcome	ORAL
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Date:03/05/02ISR Number: 3877383-4Report Type:Expedited (15-DaCompany Report #A0360114A  
Age:41 YR Gender:Female I/FU:F

Outcome Dose Other 4500MG	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Unknown		Arrhythmia Confusional State Convulsion Heart Rate Decreased Overdose Somnolence Tachycardia		Zyban	PS	Glaxo Wellcome	ORAL

Date:03/05/02ISR Number: 3877384-6Report Type:Expedited (15-DaCompany Report #A0360115A  
Age:62 YR Gender:Female I/FU:F

Outcome Dose Other 150MG Twice per day 100MG Per day 25MG Per day	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Convulsion Drug Interaction Drug Level Increased Loss Of Consciousness	Consumer	Bupropion Surmontil Electroconvulsive Therapy Liothyronine	PS SS C C	Glaxo Wellcome Glaxo Wellcome	ORAL ORAL ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/05/02ISR Number: 3877388-3Report Type:Expedited (15-DaCompany Report #A0360367A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Liver Disorder	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
		Pain	Professional	Ssri	C		
		Swelling					

Date:03/05/02ISR Number: 3877412-8Report Type:Expedited (15-DaCompany Report #B0134019A

Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Mouth Haemorrhage		Zyban	PS	Glaxo Wellcome	ORAL
150MG As		Overdose					
directed							
				Fluoxetine	SS		ORAL
20MG Per day				Olanzapine	SS		ORAL
7.5MG Twice							
per day				Aceprometazine + Meprobamate	SS		ORAL
1UNIT per day				Anethole	SS		ORAL
6UNIT per day				Zolpidem	SS		ORAL
1UNIT per day				Loxapine	SS		ORAL
50MG Per day				Cyamemazine	C		ORAL
25MG Three							
times per day				Pristinamycin	C		ORAL

Date:03/05/02ISR Number: 3877415-3Report Type:Expedited (15-DaCompany Report #B0259607A

Age:57 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Erythema		Zyban	PS	Glaxo Wellcome	
UNKNOWN		150MG Per day	39 DAY					
			Hypersensitivity		Fluvastatin	C		ORAL
	80MG Per day							
			Infarction		Nateglinide	C		ORAL
	120MG Per day							
			Malaise					

Date:03/05/02ISR Number: 3877417-7Report Type:Expedited (15-DaCompany Report #B0260160A  
Age:32 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Abnormal Behaviour		Zyban	PS	Glaxo Wellcome	ORAL
1UD Per day								
Initial or Prolonged			Confusional State		Ciblor	SS	Glaxo Wellcome	ORAL
2UD Unknown								
			Disorientation		Aspegic	SS		ORAL
1G Three								
times per day			Headache					
1UD Twice per			Muscular Weakness		Solupred	SS		ORAL
day								
			Musculoskeletal Stiffness					
1UD Three								
times per day			Nausea		Exomuc	SS	Glaxo Wellcome	ORAL
			Photophobia					

Date:03/05/02ISR Number: 3877418-9Report Type:Expedited (15-DaCompany Report #B0260324A  
Age:49 YR Gender:Male I/FU:I

Outcome	PT
Death	Brain Oedema
Hospitalization -	C-Reactive Protein
Initial or Prolonged	Increased
	Cardiac Arrest
	Cardio-Respiratory Arrest
	Cyanosis

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Fall Hepatic Failure Injury				
75MG per day		Laryngeal Oedema Leukocytosis	Zyban Kardegic	PS SS	Glaxo Wellcome	ORAL ORAL
2TAB per day	1 DAY	Malaise	Surgam	SS		ORAL
1 DAY		Mydriasis	Rulid	SS	Glaxo Wellcome	ORAL
UNKNOWN		Pyrexia	Elisor	C		
		Renal Failure Renal Impairment	Discotrine Sectral	C C	Glaxo Wellcome	ORAL
UNKNOWN		Syncope Ventricular Fibrillation	Artotec	C		ORAL

Date:03/05/02ISR Number: 3877421-9Report Type:Expedited (15-DaCompany Report #B0260698A  
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN		Anxiety		Quomem	PS	Glaxo Wellcome	
Initial or Prolonged UNKNOWN	150MG per day	Body Temperature Decreased		Levothyroxine	SS	Glaxo Wellcome	
Disability Other		Bowel Sounds Abnormal Cardiac Murmur Chest Pain Chills Diarrhoea Echocardiogram Abnormal Electrocardiogram St Segment Depression Fungal Infection Insomnia Malaise Palpitations Pco2 Decreased Red Blood Cells Urine Respiratory Rate Increased					

Thirst  
Vomiting

Date:03/05/02ISR Number: 3877426-8Report Type:Expedited (15-DaCompany Report #B0260870A  
Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Unknown 6 DAY	Cardiac Failure		Zyban	PS	Glaxo Wellcome	ORAL
Disability		Congestive		Salmeterol	C	Glaxo Wellcome	
RESPIRATORY							
Other		Chest Pain					
(INHALATION)							
RESPIRATORY		Dyspnoea Exacerbated		Salbutamol	C	Glaxo Wellcome	
(INHALATION)							

Date:03/05/02ISR Number: 3879324-2Report Type:Expedited (15-DaCompany Report #A204303  
Age: Gender:Male I/FU:I

Outcome	PT
Required	Blood Pressure Increased
Intervention to	Burns Third Degree
Prevent Permanent	Euphoric Mood
Impairment/Damage	Nausea
	Pruritus

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100.00 MG		Skin Disorder Somnolence Urine Analysis Abnormal Weight Increased	Consumer	Zoloft Tablets	PS		ORAL
TOTAL DAILY							
ORAL							
TOPICAL	TOPICAL			Benadryl (Diphenhydramine)	SS		
ORAL				Wellbutrin (Bupropion)	SS		ORAL
ORAL				Oxycodone	SS		ORAL
				Acetaminophen/Diphen hydramine	C		
				Darvon (Propoxyphene)	C		
				Vitamin C (Ascorbic Acid)	C		

Date:03/05/02ISR Number: 3879326-6Report Type:Expedited (15-DaCompany Report #A103444  
Age:79 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
50.00 MG		Anger	Consumer	Zoloft Tablets	PS		
Intervention to TOTAL DAILY		Autoimmune Thyroiditis					
Prevent Permanent 300.00 MG		Balance Disorder		Wellbutrin	SS		ORAL
Impairment/Damage TOTAL BID		Blood Oestrogen Decreased					
ORAL		Cerebral Haemorrhage					
		Cerebrovascular Accident		Coumadin	SS		
300.00 MG		Contusion		Effexor Sr	SS		ORAL
TOTAL BID		Convulsion					

ORAL

Drug Ineffective

Epistaxis	Toprol	C
Fall	Zebeta	C
Head Injury	Lanoxin	C
Heart Rate Decreased	Lasix	C
Heart Rate Increased	Synthroid	C
Hostility	Vitamin E	C
Hyperhidrosis	Estrogen	C
Illogical Thinking	Amantadine	C
Laceration	Zocor	C
Medication Error	Estradiol Vaginal	
Meningioma	Ring	C
Nuclear Magnetic Resonance Imaging Brain Abnormal		
Palpitations		
Parkinsonism		
Prothrombin Time Prolonged		
Syncope		
Vaginal Haemorrhage		

Date:03/05/02ISR Number: 3880013-9Report Type:Expedited (15-DaCompany Report #A203837

Age:24 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Dysphonia
Initial or Prolonged	Face Oedema
	Laryngeal Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	PT	Report Source	Product	Role	Manufacturer	Route
25.00 MG		Oedema Peripheral Urticaria		Foreign Health Professional	Zyrtec Tablets Hydroxyzine	PS SS		ORAL
TOTAL:DAILY:0								
150.00 MG					Prazepam Zyban	SS SS		ORAL
TOTAL:					.....	C		

Date:03/07/02ISR Number: 3879227-3Report Type:Expedited (15-DaCompany Report #A0360688A  
Age: Gender:Female I/FU:I

Dose	Duration	Outcome	PT	Report Source	Product	Role	Manufacturer	Route
		Congenital Anomaly	Dacryostenosis Congenital Drug Exposure Via Breast Milk Lethargy Maternal Drugs Affecting Foetus Neonatal Disorder		Wellbutrin Vitamins	PS C	Glaxo Wellcome	ORAL ORAL

Date:03/07/02ISR Number: 3879242-XReport Type:Expedited (15-DaCompany Report #B0261298A  
Age:20 YR Gender:Female I/FU:I

Dose	Duration	Outcome	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	5 DAY	Hospitalization -	Conversion Disorder		Zyban	PS	Glaxo Wellcome	ORAL
100MG Per day		Initial or Prolonged	Dyspnoea		Sertraline	C		ORAL
					Nitrazepam	C		ORAL

Propranolol	C	Glaxo Wellcome	ORAL
Ethinylloestradiol +			
Levonorgestrel	C		ORAL

Date:03/07/02ISR Number: 3880501-5Report Type:Direct Company Report #CTU 163005  
 Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Tinnitus		Wellbutrin Sr	PS		ORAL
150 MG BID PO							

Date:03/07/02ISR Number: 3881197-9Report Type:Expedited (15-DaCompany Report #A0359597A  
 Age:14 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Aggression
Initial or Prolonged	Agitation
	Confusional State
	Coordination Abnormal
	Difficulty In Walking
	Disorientation
	Dysarthria
	Grand Mal Convulsion
	Hallucination, Visual
	Intentional Misuse
	Nervous System Disorder



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Neurological Examination Abnormal Somnolence	Report Source	Product	Role	Manufacturer	Route
		Suicide Attempt Tachycardia Toxicologic Test Abnormal Vomiting	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
SEE DOSAGE							
TEXT/ORAL				Cannabis	C		

Date:03/07/02ISR Number: 3881362-0Report Type:Expedited (15-DaCompany Report #A0359376A  
Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Aggression Agitation Cardiotoxicity Conduction Disorder	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
1500 MG/ SEE DOSAGE		Dysarthria					
TEXT/ORAL		Electrocardiogram Qt Prolonged Grand Mal Convulsion Neurotoxicity Overdose Sinus Tachycardia Ventricular Tachycardia		Cannabis	C		

Date:03/08/02ISR Number: 3879868-3Report Type:Expedited (15-DaCompany Report #B0114598A  
Age:56 YR Gender:Male I/FU:F

Outcome	PT
Life-Threatening Hospitalization - Initial or Prolonged Other	Abdominal Pain Aggression Alcohol Poisoning Amnesia Anaphylactic Shock

Angioneurotic Oedema  
Balance Disorder  
Blister  
Blood Bilirubin Increased  
Blood Pressure Increased  
Chest Pain  
Circulatory Collapse  
Confusional State  
Convulsion  
Diarrhoea  
Disorientation  
Disturbance In Attention  
Dizziness  
Dry Skin  
Dysphagia  
Dyspnoea  
Feeling Hot  
Flatulence  
Gastroenteritis  
Irritability  
Loss Of Consciousness  
Malaise  
Nausea

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	62 DAY	Ocular Hyperaemia Oedema Peripheral Pain Pruritus Rash Erythematous Rhinitis Sensation Of Foreign Body Speech Disorder Tinnitus Urticaria	Health Professional	Zyban	PS	Glaxo Wellcome	ORAL

Date:03/08/02ISR Number: 3879875-0Report Type:Expedited (15-DaCompany Report #B0260318A  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 3 DAY Initial or Prolonged		Angina Unstable Anxiety Chest Pain Diabetes Mellitus Rash Erythematous Rash Pruritic		Zyban	PS	Glaxo Wellcome	ORAL

Date:03/11/02ISR Number: 3880684-7Report Type:Expedited (15-DaCompany Report #A0159874A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	27 DAY	Arthralgia Arthropathy Blister Difficulty In Walking Drug Hypersensitivity Dry Mouth Insomnia Malaise Muscle Spasms		Zyban	PS	Glaxo Wellcome	ORAL

Oedema Peripheral  
 Pain  
 Pain In Extremity  
 Pyrexia  
 Serum Sickness  
 Urticaria

Date:03/11/02ISR Number: 3880685-9Report Type:Expedited (15-DaCompany Report #A0360876A  
 Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Per day			Abortion Spontaneous	Wellbutrin	PS	Glaxo Wellcome	ORAL
UNKNOWN			Maternal Drugs Affecting	Depakote	C		
			Foetus	Prozac	C		ORAL

Date:03/11/02ISR Number: 3880692-6Report Type:Expedited (15-DaCompany Report #B0261434A  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
150MG Unknown			Arrhythmia	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged			Dyspnoea				
			Oedema Peripheral				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/11/02ISR Number: 3880693-8Report Type:Expedited (15-DaCompany Report #B0261457A  
Age:76 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Per day 43 DAY Initial or Prolonged		Volvulus Of Bowel	Zyban	PS	Glaxo Wellcome	ORAL
			Co-Proxamol	C		ORAL
			Vitamin D + Calcium	C		ORAL

Date:03/11/02ISR Number: 3880695-1Report Type:Expedited (15-DaCompany Report #D0038006A  
Age:46 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 150MG Unknown		Sudden Cardiac Death	Zyban	PS	Glaxo Wellcome	ORAL

Date:03/12/02ISR Number: 3882670-XReport Type:Expedited (15-DaCompany Report #A103444  
Age:79 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 50.00 MG Intervention to TOTAL:DAILY Prevent Permanent 300.00 MG Impairment/Damage TOTAL:PID:ORA		Anger	Zolofit Tablets	PS		
		Autoimmune Thyroiditis				
		Balance Disorder	Wellbutrin	SS		ORAL
		Blood Oestrogen Decreased				
		Blood Pressure Increased				
		Blood Thyroid Stimulating Hormone Decreased	Coumadin Effexor Sr	SS SS		ORAL
		Contusion				
		Convulsion				
		Drug Ineffective	Toprol	C		
		Epistaxis	Zebeta	C		
		Essential Tremor	Lanoxin	C		

Fall	Lasix	C
Haemorrhagic Stroke	Synthroid	C
Head Injury	Vitamin E	C
Headache	Estrogen	C
Heart Rate Decreased	Amantadine	C
Heart Rate Increased	Zocor	C
Hostility	Estradiol Vaginal	
Hyperhidrosis	Ring	C
Laceration		
Nuclear Magnetic		
Resonance Imaging Brain		
Abnormal		
Palpitations		
Parkinson'S Disease		
Prothrombin Time		
Prolonged		
Syncope		
Thinking Abnormal		
Thyroid Function Test		
Abnormal		
Transient Ischaemic		
Attack		
Treatment Noncompliance		
Vaginal Haemorrhage		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/13/02ISR Number: 3881848-9Report Type:Expedited (15-DaCompany Report #B0083476A

Age:38 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150MG Twice per day	Duration 7 DAY	Hemianopia Ventricular Hypertrophy Vision Blurred	Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
			Ethinylestradiol + Cyproterone	C		ORAL

Date:03/13/02ISR Number: 3881849-0Report Type:Expedited (15-DaCompany Report #B0092066A

Age:38 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150MG As Hospitalization - directed	Duration 7 DAY	Dizziness Hyperhidrosis	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Malignant Hypertension Ventricular Hypertrophy Visual Disturbance	No Concurrent Medication	C		

Date:03/13/02ISR Number: 3881853-2Report Type:Expedited (15-DaCompany Report #B0259427A

Age:31 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Unknown Initial or Prolonged	Duration	Agitation	Zyban	PS	Glaxo Wellcome	ORAL
		Aspiration Blood Pressure Increased Clonic Convulsion Coma Convulsion Disorientation Dystonia Extrapyramidal Disorder Hyperreflexia Hypertonia	Alcohol	C		

Hypotension  
Overdose  
Tachycardia  
Tremor  
Ventricular Tachycardia

Date:03/13/02ISR Number: 3881856-8Report Type:Expedited (15-DaCompany Report #B0260146A  
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2TAB Twice				Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	10 DAY	Medication Error					
Other		Sputum Abnormal					

Date:03/13/02ISR Number: 3881863-5Report Type:Expedited (15-DaCompany Report #B0261713A  
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice		Facial Palsy		Zyban	PS	Glaxo Wellcome	ORAL
per day	21 DAY						



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/13/02ISR Number: 3881864-7Report Type:Expedited (15-DaCompany Report #B0261714A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complications Of Maternal		Zyban	PS	Glaxo Wellcome	ORAL
150MG	Unknown	WK					
		Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus Stillbirth					

Date:03/13/02ISR Number: 3881865-9Report Type:Expedited (15-DaCompany Report #B0261801A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Angina Pectoris		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Initial or Prolonged per day		Anxiety					
		Myocardial Infarction					

Date:03/14/02ISR Number: 3882514-6Report Type:Expedited (15-DaCompany Report #A0162401A  
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Bipolar Disorder	Health	Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day 1	WK						
Initial or Prolonged UNKNOWN		Drug Ineffective	Professional	Topamax	SS		
		Psychotic Disorder		Zyprexa	SS		
UNKNOWN				Klonopin	SS		
UNKNOWN	1MG	Unknown					

Date:03/14/02ISR Number: 3882515-8Report Type:Expedited (15-DaCompany Report #A0172391A  
 Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged dosage text		Gamma-Glutamyltransferase Increased Lipase Increased	Health Professional	Wellbutrin	PS	Glaxo Wellcome	ORAL
UNKNOWN		Overdose		Trazodone	SS		

Date:03/14/02ISR Number: 3882517-1Report Type:Expedited (15-DaCompany Report #A0175186A  
Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 200MG Twice Initial or Prolonged per day		Dehydration Grand Mal Convulsion Stress	Health Professional	Wellbutrin Zoloft	PS C	Glaxo Wellcome	ORAL ORAL

Date:03/14/02ISR Number: 3882518-3Report Type:Expedited (15-DaCompany Report #A0359318A  
Age:14 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 9 MON Hospitalization - Initial or Prolonged		Convulsion Medication Error Overdose Poisoning	Health Professional	Wellbutrin	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/14/02ISR Number: 3882519-5Report Type:Expedited (15-DaCompany Report #A0360364A  
Age:12 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300MG Per day 6 MON	Grand Mal Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Influenza Like Illness		No Concurrent Medications	C		

Date:03/14/02ISR Number: 3882521-3Report Type:Expedited (15-DaCompany Report #A0361722A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 50MG Per day	Depression Grand Mal Convulsion	Health Professional	Wellbutrin Zoloft	PS SS	Glaxo Wellcome	ORAL

Date:03/14/02ISR Number: 3882537-7Report Type:Expedited (15-DaCompany Report #B0262044A  
Age:31 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - UNKNOWN 150MG	Grand Mal Convulsion Unknown		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged	Hallucination Ileus Paralytic Overdose		Alcohol	C		

Date:03/14/02ISR Number: 3882539-0Report Type:Expedited (15-DaCompany Report #B0262046A  
Age:20 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - UNKNOWN 150MG	Blood Pressure Decreased See		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged dosage text	Dizziness Electrocardiogram Qt Prolonged					

Flushing  
 Grand Mal Convulsion  
 Hallucination  
 Heart Rate Increased  
 Hypothermia  
 Overdose  
 Respiratory Rate  
 Increased  
 Tachycardia  
 Vomiting

Date:03/14/02ISR Number: 3882540-7Report Type:Expedited (15-DaCompany Report #B0262054A

Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day		Overdose		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged UNKNOWN	10TAB	Suicide Attempt		Paracetamol	SS	Glaxo Wellcome	
cumulative dose							
10TAB				Panadeine Forte	SS	Glaxo Wellcome	
cumulative dose							
UNKNOWN	10TAB			Mersyndol Forte	SS		
cumulative dose							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/18/02ISR Number: 3883871-7Report Type:Expedited (15-DaCompany Report #A0361528A  
Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Apnoea		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Convulsion		Zofran	SS	Glaxo Wellcome	
INTRAVENOUS							
		Depressed Level Of		Versed	SS		
INTRAVENOUS							
		Consciousness		Fentanyl	SS		
INTRAVENOUS							
		Drug Interaction		Toradol	SS		
INTRAVENOUS							
		Heart Rate Increased					
		Hyperventilation					

Date:03/18/02ISR Number: 3883872-9Report Type:Expedited (15-DaCompany Report #A0361723A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abnormal Behaviour		Wellbutrin	PS	Glaxo Wellcome	ORAL
200MG Twice							
Initial or Prolonged		Condition Aggravated					
per day	6 MON						
		Convulsion					

Date:03/18/02ISR Number: 3883873-0Report Type:Expedited (15-DaCompany Report #A0361761A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Amnesia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Initial or Prolonged		Bipolar Disorder					
per day	5 DAY						
RESPIRATORY		Convulsion		Inhalers	C		
(INHALATION)		Loss Of Consciousness					
				Imipramine	C		ORAL
				Lodine	C		ORAL

Vicodin Es  
Unspecified  
Medication

C  
C

ORAL  
ORAL

Date:03/19/02ISR Number: 3885899-XReport Type:Expedited (15-DaCompany Report #A103444  
Age:79 YR Gender:Female I/FU:F

Outcome	PT
Required	Anger
Intervention to	Balance Disorder
Prevent Permanent	Blood Oestrogen Decreased
Impairment/Damage	Blood Pressure Increased
	Blood Thyroid Stimulating
	Hormone Decreased
	Cerebral Haemorrhage
	Cerebrovascular Accident
	Contusion
	Convulsion
	Drug Ineffective
	Epistaxis
	Fall
	Head Injury
	Headache
	Heart Rate Decreased
	Heart Rate Increased
	Hostility
	Hyperhidrosis
	Illogical Thinking
	Laceration

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Meningioma Nuclear Magnetic Resonance Imaging Brain				
50.00 MG		Abnormal	Consumer	Zoloft Tablets	PS	
TOTAL:DAILY		Overdose				
300.00 MG		Palpitations		Wellbutrin	SS	ORAL
TOTAL:BID:ORA		Parkinsonism				
L		Prothrombin Time				
		Prolonged		Coumadin	SS	
300.00 MG		Syncope		Effexor Sr	SS	ORAL
TOTAL:BID:ORA		Treatment Noncompliance				
L		Tremor				
		Vaginal Haemorrhage		Toprol	C	
				Zebeta	C	
				Lanoxin	C	
				Lasix	C	
				Synthroid	C	
				Vitamin E	C	
				Estrogen	C	
				Amantadine	C	
				Zocor	C	
				Estradiol Vaginal Ring	C	

Date:03/19/02ISR Number: 3886238-0Report Type:Direct  
Age:41 YR Gender:Male I/FU:I

Company Report #CTU 163739

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Urticaria		Zyban	PS		ORAL
2 DAY ORAL							

Date:03/20/02ISR Number: 3885010-5Report Type:Expedited (15-DaCompany Report #B0262696A  
Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence		Quomem	PS	Glaxo Wellcome	ORAL
150MG Twice		Insomnia					
per day	7	DAY					
		Muscle Spasms		Loratadine +			
		Syncope		Pseudoephedrine	C		
UNKNOWN	1TAB	Twice					
per day							
				Unspecified Nasal			
				Spray	C		NASAL

Date:03/20/02ISR Number: 3885013-0Report Type:Expedited (15-DaCompany Report #D0037982A  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Deafness		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -		Intentional Misuse					
Initial or Prolonged		Suicide Attempt					
Other		Therapeutic Agent					
		Toxicity					
		Tinnitus					
		Vestibular Disorder					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/20/02ISR Number: 3885680-1Report Type:Direct  
Age:52 YR Gender:Female I/FU:I

Company Report #CTU 163802

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Pruritus Rash Erythematous		Zyban	PS		

Date:03/20/02ISR Number: 3892700-7Report Type:Direct  
Age:40 YR Gender:Male I/FU:I

Company Report #CTU 164583

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG BID		Arthralgia Myalgia		Zyban	PS		

Date:03/21/02ISR Number: 3885584-4Report Type:Expedited (15-DaCompany Report #B0262336A  
Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death UNKNOWN	150MG	As Abdominal Pain Upper		Zyban	PS	Glaxo Wellcome	
directed	42 DAY	Asthenia					
UNKNOWN	150MG	Per day Cerebral Haemorrhage		Venlafaxine	SS		
40MG Variable		Cerebrovascular Accident		Pantoprazole	C		
dose		Condition Aggravated					
25MG Per day		Confusional State		Rofecoxib	C		
10MG As		Disturbance In Attention		Temazepam	C		
required		Fall					
		Hypertension					
		Malaise					
		Snoring					
		Subdural Haematoma					
		Tremor					

Weight Decreased

Date:03/21/02ISR Number: 3885586-8Report Type:Expedited (15-DaCompany Report #B0262560A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Ectopic Pregnancy		Zyban	PS	Glaxo Wellcome	ORAL

Date:03/21/02ISR Number: 3885588-1Report Type:Expedited (15-DaCompany Report #B0262674A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abdominal Pain Aortic Aneurysm Rupture Malaise Nervousness Sudden Death		Zyban	PS	Glaxo Wellcome	ORAL

Date:03/21/02ISR Number: 3885589-3Report Type:Expedited (15-DaCompany Report #B0262805A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Zyban	PS	Glaxo Wellcome	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/21/02ISR Number: 3887372-1Report Type:Expedited (15-DaCompany Report #A204303

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required	100.00 MG	Blood Pressure Increased	Consumer	Zoloft Tablets	PS		ORAL
Intervention to	TOTAL:DAILY:0	Burns Third Degree	Health				
Prevent Permanent	RAL	Euphoric Mood	Professional				
Impairment/Damage	TOPICAL	Nausea		Benadryl			
		Pruritus		(Diphenhydramine)	SS		
		Somnolence		Wellbutrin(Bupropion			
		Weight Increased		)	SS		ORAL
ORAL				Oxycodone	SS		ORAL
ORAL				Acetaminophen/Diphen			
				hydramine	C		
				Darvon			
				(Propoxyphene)	C		
				Vitamin C (Ascorbic			
				Acid)	C		

Date:03/21/02ISR Number: 3888014-1Report Type:Direct Company Report #CTU 163962

Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Completed Suicide		Wellbutrin Sr 150 Mg	PS		
Death		Gait Disturbance		Effexor Xr 150 Mg	SS		
BID				Claritin	C		
ONE DAILY				Zyrtec	C		

Date:03/22/02ISR Number: 3886683-3Report Type:Expedited (15-DaCompany Report #A0362556A

Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 200MG Twice Initial or Prolonged per day	Apnoea Coma	Wellbutrin	PS	Glaxo Wellcome	ORAL
25MG In the morning	Hypotonia Mydriasis	Elavil	C	Glaxo Wellcome	

Date:03/22/02ISR Number: 3886691-2Report Type:Expedited (15-DaCompany Report #B0261810A  
Age:32 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Unknown Initial or Prolonged		Abnormal Behaviour		Zyban	PS	Glaxo Wellcome	ORAL
		Anxiety Condition Aggravated Confusional State Depression Hallucination, Auditory Head Injury Injury Insomnia Loss Of Consciousness Mental Disorder Paranoia Schizoaffective Disorder Suicide Attempt Thinking Abnormal		Benzodiazepines Nicotine	C C	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/22/02ISR Number: 3887103-5Report Type:Expedited (15-DaCompany Report #HQ2442222JUN2001  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression	Literature	Effexor (Venlafaxine Hydrochloride, Unspec)	PS		ORAL
ORAL		Drug Ineffective					
		Murder					
				Haldol (Haloperidol)	SS		
				Remeron (Mirtazapine)	SS		
				Wellbutrin (Amfebutamone Hydrochloride)	SS		

Date:03/25/02ISR Number: 3887336-8Report Type:Expedited (15-DaCompany Report #B0262751A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Respiratory Depression		Zyban	PS	Glaxo Wellcome	ORAL

Date:03/25/02ISR Number: 3888085-2Report Type:Direct Company Report #CTU 164102  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Alcoholism		Clonazepam 1mg Tablet Generic	PS		ORAL
4 DAY ORAL		Grand Mal Convulsion					
		Muscle Twitching		Wellbutrin 150mg Tablet Glaxo Wellcome	SS	Glaxo Wellcome	ORAL
2 DAY ORAL				Azmacort	C		
				Serevent	C		
				Albuteral	C		
				Zoloft	C		

Date:03/26/02ISR Number: 3888170-5Report Type:Expedited (15-DaCompany Report #A0363291A  
Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Carbon Monoxide Poisoning		Bupropion	PS	Glaxo Wellcome	
UNKNOWN							

Date:03/26/02ISR Number: 3888171-7Report Type:Expedited (15-DaCompany Report #A0363292A  
Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Bupropion	PS	Glaxo Wellcome	
UNKNOWN							

Date:03/26/02ISR Number: 3889776-XReport Type:Direct Company Report #CTU 164235  
Age:74 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Pruritus Rash Urticaria		Bupropion Aspirin Glipizide Lisinopril Metformin	PS C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/27/02ISR Number: 3890887-3Report Type:Expedited (15-DaCompany Report #EMADSS2002001073  
Age:18 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - Initial or Prolonged 1.3 G, 1 IN 1 Disability TIME(S), ORAL Required Intervention to 9 G, 1 IN 1 Prevent Permanent TIMES(S), Impairment/Damage ORAL	Cardiac Arrest Cerebral Ischaemia Medication Error Overdose Status Epilepticus	Foreign Health Professional	Zydol (Tablet) (Tramadol Hydrochlorde)  Zyban (Amfebutamone Hydrochloride)	PS  SS		ORAL  ORAL

Date:03/28/02ISR Number: 3890150-0Report Type:Expedited (15-DaCompany Report #A0363470A  
Age:23 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other	Alcohol Interaction Anaphylactic Reaction		Wellbutrin Ethanol Ortho Tri-Cyclen Neoral	PS SS C C	Glaxo Wellcome	ORAL

Date:03/28/02ISR Number: 3890167-6Report Type:Expedited (15-DaCompany Report #B0262849A  
Age:44 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other UNKNOWN directed 21 DAY	Dyspepsia Exanthem Headache Insomnia Nasopharyngeal Disorder Nervousness		Zyntabac	PS	Glaxo Wellcome	

Date:03/28/02ISR Number: 3891873-XReport Type:Expedited (15-DaCompany Report #B0262338A  
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage DAY / ORAL		Haematuria	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:03/28/02ISR Number: 3891892-3Report Type:Expedited (15-DaCompany Report #B0262209A  
Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN DOSAGE TEXT/UNKNOWN	150 MG/	SEE	Foreign Literature	Zyban (Bupropion Hydrochloride)	PS		
		Cardiac Arrest Convulsion	Health				
		Drug Toxicity	Professional				
		Loss Of Consciousness		Ethanol (Formulation Unknown) (Alcohol)	SS		
		Overdose Respiratory Depression Tachycardia					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/28/02ISR Number: 3892047-9Report Type:Expedited (15-DaCompany Report #B0122442A

Age:34 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Disability 150 MG / UNK / ORAL	Cheilitis Dermatitis Bullous Leukocytosis  Oral Soft Tissue Disorder	Foreign Health Professional	Zyban Tablet -(Bupropion Hydrochloride)	PS		ORAL
ORAL	Pyrexia Rash Rash Pustular		Propofan (Formulation Unknown) (Propofan)	SS		ORAL
UNKNOWN	Skin Lesion		Prazepam (Formulation Unknown) (Prazepam)	SS		
UNKNOWN			Zolpidem (Formulation Unknown) (Zolpidem)	SS		
ORAL			Alprazolam (Formulation Unknown)	SS		ORAL

Date:03/28/02ISR Number: 3892048-0Report Type:Expedited (15-DaCompany Report #B0262369A

Age:74 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG, ORAL	Choreoathetosis	Foreign	Zyban (Bupropion Hydrochloride)	PS		ORAL
			Naproxen	C		
			Atorvastatin Calcium	C		
			Clopidogrel	C		

Date:03/28/02ISR Number: 3892071-6Report Type:Expedited (15-DaCompany Report #B0262088A

Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage DAY / ORAL		Condition Aggravated Impaired Driving Ability Nystagmus Tremor	Foreign Health Professional	Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:03/28/02ISR Number: 3892073-XReport Type:Expedited (15-DaCompany Report #B0262087A  
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG/ AS DIRECTED / ORAL		Asthenia Lymphocyte Count Increased Pain Paralysis Pruritus Generalised Rash Generalised	Foreign Health Professional	Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:03/28/02ISR Number: 3892074-1Report Type:Expedited (15-DaCompany Report #B0261810A  
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT
Hospitalization - Initial or Prolonged		Abnormal Behaviour Aggression

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Anxiety Confusional State Depression Emotional Distress Energy Increased	Foreign Consumer	Zyban (Bupropion Hydrochloride)	PS		ORAL
		Hallucination, Auditory Head Injury Injury Insomnia Loss Of Consciousness Mental Disorder Paranoia Speech Disorder Suicide Attempt Tendon Injury Toxicologic Test Abnormal		Benzodiazepines Nicotine	C C		

Date:03/28/02ISR Number: 3892075-3Report Type:Expedited (15-DaCompany Report #B0261797A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG / TWICE PER DAY / ORAL		Diplopia Drug Ineffective Grand Mal Convulsion Status Epilepticus	Foreign	Zyban (Bupropion Hydrochloride) Multivitamin	PS C		ORAL

Date:03/28/02ISR Number: 3892077-7Report Type:Expedited (15-DaCompany Report #B0261782A  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150 MG / PER DAY/ ORAL		Suicide Attempt	Foreign	Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:03/28/02ISR Number: 3892085-6Report Type:Expedited (15-DaCompany Report #B0262370A  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Alcohol Problem Anxiety	Foreign	Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / ORAL		Decreased Activity Stress					

Date:03/28/02ISR Number: 3892125-4Report Type:Expedited (15-DaCompany Report #B0262329A  
Age: Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Agitation Anxiety Arthralgia Constipation Disturbance In Attention Dizziness Dry Mouth Dysgeusia Dyspnoea

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / ORAL		Face Oedema Fatigue Flushing Hallucination Headache	Foreign Consumer	Zyban (Bupropion Hydrochloride)	PS		ORAL
		Hostility Insomnia Irritability Mood Altered Nausea Oedema Mouth Pyrexia Tinnitus Tremor Urticaria Visual Acuity Reduced Wheezing					

Date:03/28/02ISR Number: 3892126-6Report Type:Expedited (15-DaCompany Report #B0262184A  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	3000 MG	Intentional Misuse Somnolence Suicide Attempt	Foreign Health Professional	Zyban (Bupropion Hydrochloride)	PS		ORAL
SINGLE DOSE	ORAL			Benzodiazepines (Formulation Unknown)	SS		ORAL
SINGLE DOSE	ORAL			(Benzodiazepines)			

Date:03/29/02ISR Number: 3891154-4Report Type:Expedited (15-DaCompany Report #A0170371A  
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day				Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Chills Convulsion Mental Disorder Pyrexia Speech Disorder Weight Decreased		Varivax	SS	Glaxo Wellcome	

Date:03/29/02ISR Number: 3891160-XReport Type:Expedited (15-DaCompany Report #A0363028A  
Age:8 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1 MON				Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Rash Red Blood Cell Sedimentation Rate Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/02ISR Number: 3891164-7Report Type:Expedited (15-DaCompany Report #A0364009A  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	61 DAY	Lung Neoplasm Malignant Therapeutic Response Unexpected		Zyban	PS	Glaxo Wellcome	ORAL

Date:03/29/02ISR Number: 3891165-9Report Type:Expedited (15-DaCompany Report #B0259229A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG As directed		Drug Level Above Therapeutic Overdose Toxicologic Test Abnormal	Health Professional	Zyban Alcohol Salicylate	PS C C	Glaxo Wellcome	ORAL

Date:03/29/02ISR Number: 3891167-2Report Type:Expedited (15-DaCompany Report #B0259698A  
 Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day 6MG Unknown	5 DAY	Agitation Amnesia Confusional State Disturbance In Attention Dizziness Excitability Loss Of Consciousness Malaise Road Traffic Accident Sleep Attacks	Health Professional	Zyban Bromazepam Alcohol	PS SS SS	Glaxo Wellcome	ORAL ORAL

Date:03/29/02ISR Number: 3891172-6Report Type:Expedited (15-DaCompany Report #B0263091A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 15 DAY		Drug Interaction	Health	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Intentional Misuse Somnolence	Professional	Trizivir Myolastan	SS SS	Glaxo Wellcome	ORAL ORAL
500MG per day		Suicide Attempt					

Date:03/29/02ISR Number: 3891173-8Report Type:Expedited (15-DaCompany Report #B0263361A  
Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Depression		Zyban	PS	Glaxo Wellcome	
UNKNOWN	150MG	Unknown					
Other		Suicidal Ideation		No Concurrent Medications	C		

Date:03/29/02ISR Number: 3891178-7Report Type:Expedited (15-DaCompany Report #D0038006A  
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Sudden Cardiac Death	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/02ISR Number: 3891179-9Report Type:Expedited (15-DaCompany Report #D0038185A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Unknown	WK		Zyban	PS	Glaxo Wellcome	ORAL
				Haemorrhagic Stroke Myocardial Infarction Renal Embolism			

Date:03/30/02ISR Number: 3892695-6Report Type:Direct Company Report #CTU 164579  
Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG PO BID			Bupropion (Wellbutrin)	PS		ORAL
				Urticaria			

Date:03/30/02ISR Number: 3892696-8Report Type:Direct Company Report #CTU 164580  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG PO BID Initial or Prolonged				Bupoprion (Zyban) Celexa	PS C		ORAL
				Urticaria			

Date:03/30/02ISR Number: 3892713-5Report Type:Direct Company Report #CTU 164588  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other Required Intervention to Prevent Permanent Impairment/Damage				Zyban	PS		
				Angle Closure Glaucoma Cataract			

Date:04/01/02ISR Number: 3892060-1Report Type:Expedited (15-DaCompany Report #B0095597A  
Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Aphasia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice						
Hospitalization -	Hemiplegia					
per day 5 DAY						
Initial or Prolonged						

Date:04/01/02ISR Number: 3892063-7Report Type:Expedited (15-DaCompany Report #B0263173A  
Age:42 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Areflexia
Initial or Prolonged	Arterial Disorder
	Blood Pressure
	Fluctuation
	Cerebral Perfusion
	Pressure Decreased
	Dysphasia
	Electroencephalogram
	Abnormal
	Hemiparesis
	Hemiplegia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
UNKNOWN	150MG Twice	Nervous System Disorder Single Photon Emission Computerised Tomogram	Zyban	PS	Glaxo Wellcome	
per day	5 DAY	Abnormal Ultrasound Scan Abnormal				

Date:04/01/02ISR Number: 3892773-1Report Type:Direct Company Report #CTU 164682  
 Age:29 YR Gender:Male I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
Other		Grand Mal Convulsion	Zyban, Glaxowellcome	PS	Glaxowellcome	
150 MG BID,						
047			Nicotine Patch	C		

Date:04/02/02ISR Number: 3892792-5Report Type:Expedited (15-DaCompany Report #A0135919A  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
Disability		Burning Sensation	Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Dysgeusia				
per day	12 DAY	Dyskinesia	Amitriptyline	C		
UNKNOWN	200MG Per day	Hypoaesthesia	Unknown Medication	C		
		Insomnia				
		Muscle Strain				
		Neuralgia				
		Neuropathy Peripheral				
		Oral Pain				
		Pain				
		Panic Attack				
		Paraesthesia				
		Rash				

Date:04/02/02ISR Number: 3892796-2Report Type:Expedited (15-DaCompany Report #A0359984A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Level Increased Overdose		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:04/02/02ISR Number: 3892799-8Report Type:Expedited (15-DaCompany Report #A0363247A  
Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day 5 DAY Initial or Prolonged		Convulsion Cyanosis Depression Grand Mal Convulsion Pneumonia		Zyban Plavix Altace Prozac Flexeril Lipitor Toprol Xl Advair Albuterol Duragesic	PS C C C C C C C C C	Glaxo Wellcome      Glaxo Wellcome Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/02/02ISR Number: 3892801-3Report Type:Expedited (15-DaCompany Report #A0363417A  
 Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 200MG Twice Initial or Prolonged per day			Abdominal Pain	Wellbutrin	PS	Glaxo Wellcome	ORAL
			Chest Pain				
	MON		Gallbladder Disorder	Topamax	SS		
			Liver Function Test Abnormal	Lipitor	C		

Date:04/02/02ISR Number: 3893886-0Report Type:Direct Company Report #CTU 164757  
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1 PILL 3 TIMES ORAL			Diarrhoea Gastroenteritis Viral	Wellbutrin Sr 150 Mg Glaxo	PS	Glaxo	ORAL
			Nausea				
			Pharmaceutical Product Complaint				

Date:04/02/02ISR Number: 3894237-8Report Type:Direct Company Report #CTU 164810  
 Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG BID ORAL			Pruritus	Zyban 150mg Glaxo-Wellcome	PS	Glaxo-Wellcome	ORAL
				Nicoderm	C		

Date:04/03/02ISR Number: 3894510-3Report Type:Expedited (15-DaCompany Report #A0356146A 2001029370-1  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anxiety Female Orgasmic Disorder Libido Decreased	Consumer	Paxil Tablet (Paroxetine Hydrochloride)	PS		ORAL
40 MG/ORAL		Pregnancy		Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride) Clonazepam	SS C		

Date:04/03/02ISR Number: 3896444-7Report Type:Direct Company Report #CTU 164881  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 PO QD X 3D THEN BID		Disturbance In Attention Lethargy Paraesthesia Sinusitis Tremor		Zyban 150mg Tbsr  Nicorette 2mg	PS  C	Tbsr	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/04/02ISR Number: 3894062-8Report Type:Expedited (15-DaCompany Report #B0262329A  
 Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Unknown Initial or Prolonged		Agitation Anxiety Constipation Disturbance In Attention Dizziness Dry Mouth Dysgeusia Dyspepsia Fatigue Flushing Hallucination Headache Hostility Insomnia Irritability Mood Altered Nausea Serum Sickness Tinnitus Tremor Visual Acuity Reduced		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/04/02ISR Number: 3894068-9Report Type:Expedited (15-DaCompany Report #B0263835A  
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 14 DAY		Discomfort Dysphagia Oedema Peak Expiratory Flow Rate Decreased Respiratory Disorder Urticaria Generalised		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/04/02ISR Number: 3894069-0Report Type:Expedited (15-DaCompany Report #B0264143A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Retinal Detachment		Zyntabac	PS	Glaxo Wellcome	ORAL
150MG	Unknown						

Date:04/05/02ISR Number: 3894837-5Report Type:Expedited (15-DaCompany Report #A0364473A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Disorder		Wellbutrin	PS	Glaxo Wellcome	ORAL
300MG	Per day						
		Constipation		Paxil	C	Glaxo Wellcome	ORAL
20MG	Per day 1 YR						
		Dyspnoea		Unspecified			
		Haemorrhoids		Medication	C		
		Libido Decreased					
		Night Sweats					
		Tremor					
		Vision Blurred					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/05/02ISR Number: 3894838-7Report Type:Expedited (15-DaCompany Report #B0111133A  
 Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death				Zyban	PS	Glaxo Wellcome	
150MG Single							
Hospitalization -	1 DAY						
dose		Lower Respiratory Tract					
Initial or Prolonged		Infection		Temazepam	C		ORAL
10MG Per day				Loperamide	C		ORAL
2MG Per day		Malaise					
300MG per day		Renal Failure		Allopurinol	C	Glaxo Wellcome	ORAL
		Urinary Tract Infection		Co-Proxamol	C		ORAL
7.5MG per day		Vomiting		Meloxicam	C		ORAL
				Salbutamol	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)		200MCG Twice					
per day							

Date:04/05/02ISR Number: 3894839-9Report Type:Expedited (15-DaCompany Report #B0112291A  
 Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death				Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	2 DAY						
Hospitalization -							
2MG Per day		Renal Failure		Loperamide	C		ORAL
Initial or Prolonged				Allopurinol	C	Glaxo Wellcome	ORAL
300MG Per day				Meloxicam	C		ORAL
7.5MG Per day				Salbutamol	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)		100MCG As					
required							

10MG Per day Temazepam C ORAL

Date:04/08/02ISR Number: 3898185-9Report Type:Direct Company Report #CTU 165150  
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
ONCE PO QD X		Hypersensitivity		Zyban 150 Mg Tbsr	PS		
3D, THEN BID		Rash					
TOPICAL	ONCE PATCH			Habitrol 21 Mg Patch	SS		
TOP QD							

Date:04/09/02ISR Number: 3896308-9Report Type:Expedited (15-DaCompany Report #B0131342A  
Age:72 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Creatinine		Zyban	PS	Glaxo Wellcome	ORAL
13 DAY		Increased		Piribedil	SS		
UNKNOWN		Hyperkalaemia		Gemfibrozil	SS		
UNKNOWN				Zopiclone	SS		
UNKNOWN				Buflomedil Hydrochloride	SS		
UNKNOWN				Metformin Hydrochloride	SS		
UNKNOWN				Clonazepam	C		

Date:04/09/02ISR Number: 3896311-9Report Type:Expedited (15-DaCompany Report #B0263833A  
Age:65 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Arthralgia
Initial or Prolonged	Dysphagia
	Hypersensitivity



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Myalgia				
		Rash				
		Rash Psoriaform	Report Source	Product	Role	Manufacturer
Dose	Duration					Route
150MG Per day	8 DAY	Weight Decreased		Zyban	PS	Glaxo Wellcome

Date:04/09/02ISR Number: 3896313-2Report Type:Expedited (15-DaCompany Report #B0264067A  
 Age:33 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose		Death		Zyban	PS	Glaxo Wellcome	ORAL
Death							
300MG per day							

Date:04/09/02ISR Number: 3896314-4Report Type:Expedited (15-DaCompany Report #B0264074A  
 Age: Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose		Myocardial Infarction		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -							
150MG Per day	1 MON						
Initial or Prolonged				Antihypertensive (Unspecified)	C		
				Statines	C		

Date:04/09/02ISR Number: 3899602-0Report Type:Direct Company Report #CTU 165372  
 Age:31 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose		Anaphylactic Reaction		Bupropion 75 Mg			
Hospitalization -		Joint Swelling		Generic	PS		ORAL
Initial or Prolonged							
75MG 2 TIMES		Oedema Peripheral					
ORAL							
		Pain		Serax-Oxezapam	C		
		Pruritus					
		Throat Tightness					
		Urticaria					

Date:04/10/02ISR Number: 3896976-1Report Type:Expedited (15-DaCompany Report #A0364404A  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Twice		Blindness Transient		Wellbutrin	PS	Glaxo Wellcome	ORAL
per day		Visual Field Defect					
				Zoloft	C		
				Trazodone	C		
				Atenolol	C		
				Micardis	C	Glaxo Wellcome	
				Hydrochlorothiazide	C		
				Estrace	C		
				Lipitor	C		
				Plavix	C		

Date:04/10/02ISR Number: 3898377-9Report Type:Expedited (15-DaCompany Report #2002099876FR  
Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Fall Inappropriate Antidiuretic Hormone	Foreign Health Professional	Aldactazine (Spironolactone, Althiazide) Tablet	PS		ORAL
ORAL		Secretion Weight Increased		Zyban (Amfebutamone Hydrochloride)	SS		ORAL

150 MG, BID,

19-Aug-2005 12:44 PM  
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL

Date:04/10/02ISR Number: 3899394-5Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #CTU 165452

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 150 MG QD Intervention to ORAL Prevent Permanent Impairment/Damage		Erythema Multiforme		Wellbutrin Sr 150mg	PS		ORAL

Date:04/10/02ISR Number: 3899460-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200269  
 Age:17 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Grand Mal Convulsion Overdose	Health Professional	Neurontin (Gabapentin)	PS		ORAL
PER ORAL				Amfebutamone Hydrochloride	SS		ORAL
PER ORAL				Carisoprodol	SS		ORAL
				Venlafaxine Hydrochloride	C		

Date:04/10/02ISR Number: 3899545-2Report Type:Direct  
 Age:9 YR Gender:Male I/FU:I

Company Report #CTU 165455

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG PO /QD		Dyskinesia Ear Disorder		Wellbutrin Sr 100mg Qd	PS		ORAL

Date:04/11/02ISR Number: 3897617-XReport Type:Expedited (15-DaCompany Report #A0359237A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300MG per day	Overdose		Wellbutrin	PS	Glaxo Wellcome	ORAL
				Depakote	C		

Date:04/11/02ISR Number: 3897618-1Report Type:Expedited (15-DaCompany Report #A0359335A  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Aura		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Blindness					
		Coma		Oxycontin	C		
		Depression		Flexeril	C		
		Disturbance In Attention		Acyclovir	C	Glaxo Wellcome	
200MG As required		Memory Impairment					
		Optic Nerve Disorder					
		Petit Mal Epilepsy					
		Sleep Attacks					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/11/02ISR Number: 3897620-XReport Type:Expedited (15-DaCompany Report #A0361528A  
 Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Apnoea		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Clonic Convulsion		Zofran	SS	Glaxo Wellcome	
INTRAVENOUS	1 DAY						
		Coma		Versed	C		
INTRAVENOUS							
		Drug Interaction		Fentanyl	C		
INTRAVENOUS							
		Heart Rate Increased		Toradol	C		
INTRAVENOUS							
		Hyperventilation		Sevoflurane	C		
RESPIRATORY							
(INHALATION)		Memory Impairment					
		Muscle Rigidity		Propofol	C		
INTRAVENOUS							
		Respiratory Rate Increased					

Date:04/11/02ISR Number: 3897621-1Report Type:Expedited (15-DaCompany Report #A0361722A  
 Age:13 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Depression		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	9 MON	Drug Interaction					
		Electroencephalogram		Zoloft	SS		
50MG Per day							
		Abnormal		Trazodone	C		
50MG At night							
		Grand Mal Convulsion					

Date:04/11/02ISR Number: 3897622-3Report Type:Expedited (15-DaCompany Report #A0361723A  
 Age:10 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Other Convulsion Wellbutrin PS Glaxo Wellcome ORAL  
 200MG Twice  
 per day 6 MON  
 Date:04/11/02ISR Number: 3897627-2Report Type:Expedited (15-DaCompany Report #A0365391A  
 Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pulmonary Hypertension		Zyban	PS	Glaxo Wellcome	ORAL
150MG See							
dosage text	8	DAY					

Date:04/11/02ISR Number: 3897642-9Report Type:Expedited (15-DaCompany Report #B0264055A  
 Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Grand Mal Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Respiratory Arrest		Antidepressant	C		

Date:04/11/02ISR Number: 3897646-6Report Type:Expedited (15-DaCompany Report #B0264147A  
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Leukocytoclastic		Zyban	PS	Glaxo Wellcome	
UNKNOWN	150MG As	Vasculitis					
directed	19	DAY					
UNKNOWN				Conjugated Oestrogens	C		
UNKNOWN	.5UNIT As			Lormetazepam	C		
required							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/11/02ISR Number: 3897647-8Report Type:Expedited (15-DaCompany Report #B0264148A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Formication		Zyban	PS	Glaxo Wellcome	
UNKNOWN	150MG	Twice					
Initial or Prolonged		Peripheral Coldness					
per day							
Other				Diltiazem			
				Hydrochloride	C	Glaxo Wellcome	
UNKNOWN	120MG	Per day					
				Isoflavone	C		
				Levothyroxine	C	Glaxo Wellcome	
UNKNOWN	50MG	Per day					

Date:04/11/02ISR Number: 3897649-1Report Type:Expedited (15-DaCompany Report #B0264529A  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Diabetes Mellitus		Zyban	PS	Glaxo Wellcome	
UNKNOWN	150MG	Per day 61 DAY					
Initial or Prolonged		Insulin-Dependent					
Disability		Headache					
		Pruritus					

Date:04/11/02ISR Number: 3897650-8Report Type:Expedited (15-DaCompany Report #B0264531A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Phlebothrombosis		Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown							
Initial or Prolonged							

Date:04/11/02ISR Number: 3897652-1Report Type:Expedited (15-DaCompany Report #B0264661A  
Age:78 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death UNKNOWN 150MG Per day 5 DAY Acute Myocardial Infarction Malaise Nausea Zyban Tramadol Hydrochloride PS C Glaxo Wellcome

Date:04/12/02ISR Number: 3898872-2Report Type:Expedited (15-DaCompany Report #A0364303A  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Angina Pectoris Hypertension		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL

Date:04/12/02ISR Number: 3898883-7Report Type:Expedited (15-DaCompany Report #B0262849A  
 Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Variable dose 21 DAY		Difficulty In Walking Dyspepsia Exanthem Headache Insomnia Nasopharyngeal Disorder Nervousness Pruritus Rash	Health Professional	Zyntabac	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/12/02ISR Number: 3898886-2Report Type:Expedited (15-DaCompany Report #B0264401A  
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	17 DAY	Overdose	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	200MG per day	Suicide Attempt		Bromazepam	SS		
	160MG Per day	Therapeutic Agent		Kardegic	C		ORAL
	2MG Per day	Toxicity		Amarel	C		ORAL
	40MG Per day			Monicor	C		ORAL

Date:04/15/02ISR Number: 3901257-3Report Type:Expedited (15-DaCompany Report #A0364109A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	TOPICAL	Application Site Pruritus	Health Professional	Nicotine Patch (Nicotine)	PS		
	TOPICAL	Convulsion		Bupropion Hydrochloride Tablet-Controlled Release (Bupropion Hydrochloride)	SS		ORAL
	150 MG//ORAL						

Date:04/16/02ISR Number: 3899837-7Report Type:Expedited (15-DaCompany Report #WAES 0204ESP00009  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	10 DAY	Deep Vein Thrombosis		Noroxin	PS	Merck & Co., Inc	ORAL
Initial or Prolonged	37 DAY			Bupropion Hydrochloride	SS		ORAL

Date:04/16/02ISR Number: 3900670-8Report Type:Direct  
Age:50 YR Gender:Female I/FU:I

Company Report #CTU 165767

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability CANT RECALL I		Dysgraphia		Wellbutrin	PS		
CUT IT IN		Nervous System Disorder					
HALF		Tremor					

Date:04/16/02ISR Number: 3901964-2Report Type:Expedited (15-DaCompany Report #200211202GDDC  
Age:71 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Foreign Health	Indapamide (Dapa-Tabs) Tablets	PS		ORAL
2.5 MG PO	436 DAY		Professional Other	Amfebutamone Hydrochloride (Zyban)	SS		
8 DAY				Quinapril Hydrochloride (Accupril)	C		
				Acetylsalicylate Calcium (Solprin)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/18/02ISR Number: 3901476-6Report Type:Expedited (15-DaCompany Report #A0364730A  
 Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	15	DAY					
		Crying					
		Hallucination, Visual		Norplant	C		
		Memory Impairment		Synthroid	C	Glaxo Wellcome	
		Panic Attack		Xanax	C		
		Schizoaffective Disorder					

Date:04/18/02ISR Number: 3901484-5Report Type:Expedited (15-DaCompany Report #B0259229A  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Drug Toxicity		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day							
		Excoriation		Alcohol	C		
		Mucous Membrane Disorder		Salicylate	C		
		Overdose					
		Pulmonary Haemorrhage					
		Pulmonary Oedema					

Date:04/18/02ISR Number: 3902435-XReport Type:Direct Company Report #CTU 166101  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Mallory-Weiss Syndrome		Bupropion 100 Mg Bid			
Initial or Prolonged		Nausea		Plus 75 Mg Hs	PS		
100 MG BID							
PLUS 75 MG HS		Pharmaceutical Product					
	YR	Complaint					
		Vomiting					

Date:04/19/02ISR Number: 3902287-8Report Type:Expedited (15-DaCompany Report #A0365819A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	13 DAY	Diarrhoea Haemorrhagic Pain		Zyban	PS	Glaxo Wellcome	ORAL
1 DAY				Aspirin	C		
2 DAY				Fexofenadine	C		
				Atorvastatin Levothyroxine	C C	Glaxo Wellcome	

Date:04/19/02ISR Number: 3902295-7Report Type:Expedited (15-DaCompany Report #B0134349A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 150MG Unknown		Death		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/19/02ISR Number: 3902296-9Report Type:Expedited (15-DaCompany Report #B0134350A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death UNKNOWN	150MG Unknown	Death		Zyban	PS	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/19/02ISR Number: 3902300-8Report Type:Expedited (15-DaCompany Report #B0265475A  
 Age:23 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Per day 4 DAY Initial or Prolonged	Pericarditis		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/22/02ISR Number: 3903730-0Report Type:Expedited (15-DaCompany Report #B0134348A  
 Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 150MG Unknown	Death		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/22/02ISR Number: 3903736-1Report Type:Expedited (15-DaCompany Report #B0265190A  
 Age:40 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Aggression Depression Dizziness Mouth Ulceration Muscle Rigidity Nausea Vision Blurred		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/22/02ISR Number: 3903738-5Report Type:Expedited (15-DaCompany Report #B0265412A  
 Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - UNKNOWN Initial or Prolonged	Aggression Agitation		Zyban	PS	Glaxo Wellcome	



Date:04/22/02ISR Number: 3903739-7Report Type:Expedited (15-DaCompany Report #B0265413A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN		Aggression		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged		Emotional Disorder					

Date:04/22/02ISR Number: 3904707-1Report Type:Expedited (15-DaCompany Report #A0141408A  
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN		Anger Delusion Depression	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG TWICE PER DAY ORAL		Dizziness Feeling Jittery Impulsive Behaviour Mania Nervousness Psychomotor Hyperactivity Suicidal Ideation Tremor Ventricular Extrasystoles		Semisodium Valproate Gabapentin Venlafaxine Hydrochloride Glipizide Vicodin	C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/22/02ISR Number: 3904734-4Report Type:Expedited (15-DaCompany Report #A204048

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 50.00 MG	Anxiety	Health	Zoloft Tablets	PS		ORAL
Initial or Prolonged TOTAL:DAILY:O Required RAL	Depression	Professional				
Intervention to 300.00 MG Prevent Permanent TOTAL:PID:ORA Impairment/Damage L	Diabetes Mellitus					
	Drug Ineffective		Wellbutrin	SS		ORAL
	Fatigue					
	Fear					
10.00 TOTAL:DAILY:O RAL	Gait Disturbance		Zyprexa	SS		ORAL
	Hallucination, Visual					
	Hemianopia					
	Insomnia					

Date:04/22/02ISR Number: 3904736-8Report Type:Expedited (15-DaCompany Report #A103444

Age:79 YR Gender:Female I/FU:F

Outcome	PT
Required	Akinesia
Intervention to	Anger
Prevent Permanent	Autoimmune Thyroiditis
Impairment/Damage	Balance Disorder
	Blood Oestrogen Decreased
	Blood Pressure Increased
	Blood Thyroid Stimulating
	Hormone Decreased
	Cerebral Haemorrhage
	Contusion
	Convulsion
	Dementia
	Drug Ineffective
	Dysphonia
	Epistaxis
	Fall

Head Injury  
Headache  
Heart Rate Decreased  
Heart Rate Increased  
Hostility  
Hyperhidrosis  
Laceration  
Medication Error  
Meningioma  
Mood Swings  
Nervous System Disorder  
Nuclear Magnetic  
Resonance Imaging  
Abnormal  
Overdose  
Palpitations  
Parkinson'S Disease  
Prothrombin Time  
Prolonged  
Staring  
Syncope  
Thinking Abnormal  
Thyroid Function Test  
Abnormal  
Transient Ischaemic  
Attack

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vaginal Haemorrhage

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
50.00 MG		Consumer	Zoloft Tablets	PS		
TOTAL:DAILY						
300.00 MG			Wellbutrin	SS		ORAL
TOTAL:BID:ORA						
L						
300.00 MG			Coumadin	SS		
			Effexor Xr	SS		ORAL
TOTAL:BID:ORA						
L						
			Toprol	C		
			Zebeta	C		
			Lanoxin	C		
			Lasix	C		
			Synthroid	C		
			Vitamin E	C		
			Estrogen	C		
			Amantadine	C		
			Zocor	C		
			Estradiol Vaginal			
			Ring	C		

Date:04/22/02ISR Number: 3904888-XReport Type:Expedited (15-DaCompany Report #WAES 0204ESP00009  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	400 MG/BID PO 10 DAY	Deep Vein Thrombosis	Other	Tab Noroxin (Norfloxacin)	PS		ORAL
150 MG/DAILY				Tab Bupropion Hcl	SS		ORAL
PO	37 DAY						

Date:04/22/02ISR Number: 3904902-1Report Type:Expedited (15-DaCompany Report #B0264913A

Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
1 TABLET/ TWICE PER DAY/ ORAL				Omeprazole Betahistine	C C		

Date:04/22/02ISR Number: 3904906-9Report Type:Expedited (15-DaCompany Report #B0264464A

Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/PER DAY/ORAL			Company Representative	Bromazepam Tablet (Bromazepam)	SS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/22/02ISR Number: 3904907-0Report Type:Expedited (15-DaCompany Report #B0264307A

Age:40 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 3000 MG/SINGLE DOSE/ORAL ORAL	Clonic Convulsion Intentional Misuse Muscle Contractions Involuntary Suicide Attempt	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
			Bromazepam Tablet (Bromazepam)	SS		ORAL
			Oxazepam	SS		

Date:04/22/02ISR Number: 3905066-0Report Type:Expedited (15-DaCompany Report #B0265019A

Age:60 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/AS DIRECTED/ UNKNOWN	Amnesia Anhedonia Convulsion Decreased Activity Dizziness Dyspnoea Fatigue Fear Gastrointestinal Disorder Joint Stiffness Loss Of Consciousness Pancreatitis Panic Attack Peritonitis Purulence Sexual Dysfunction Sleep Disorder	Foreign Health Professional	Zyban Tabltn - Zyban (Bupropion Hydrochloride)	PS		
			Simvastatin	C		
			Seloken	C		
			Renitec	C		
			Frusemide	C		
			Aspirin	C		

Date:04/22/02ISR Number: 3905146-XReport Type:Expedited (15-DaCompany Report #B0264762A  
Age:47 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Deep Vein Thrombosis	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ PER DAY/ ORAL						
800 MG / TWICE PER DAY/ ORAL			Norfloxacin (Formulation Unknown) (Norfloxacin)	SS		ORAL

Date:04/22/02ISR Number: 3905188-4Report Type:Expedited (15-DaCompany Report #B0264124A  
Age: Gender: I/FU:I

Outcome  
Death  
Congenital Anomaly  
Required  
Intervention to  
Prevent Permanent  
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 TABLET /		Congenital Diaphragmatic Hernia Maternal Drugs Affecting Foetus	Foreign	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL		Pregnancy Pulmonary Malformation		Folic Acid Amoxicillin Trihydrate	C C		

Date:04/22/02ISR Number: 3905190-2Report Type:Expedited (15-DaCompany Report #B0264762A  
Age:47 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG / PER DAY / ORAL		Deep Vein Thrombosis	Foreign	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
	800 MG / TWICE PER DAY / ORAL				Norfloxacin (Norfloxacin) (Formulation Unknown)	SS		ORAL

Date:04/22/02ISR Number: 3905407-4Report Type:Direct Company Report #CTU 166383  
Age:12 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 3X DAILY			Bipolar I Disorder Suicide Attempt		Wellbutrin 75mg Burroughs Wellcome	PS	Burroughs Wellcome	ORAL



Initial or Prolonged  
ORALLY 75MG  
Required  
EACH APPROX 1  
Intervention to  
YEAR 1 YR  
Prevent Permanent  
Impairment/Damage

Date:04/23/02ISR Number: 3904636-3Report Type:Expedited (15-DaCompany Report #A0359318A  
Age:14 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	9 MON	Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged		Drug Level Increased Medication Error Overdose Therapeutic Agent Toxicity					

Date:04/23/02ISR Number: 3904641-7Report Type:Expedited (15-DaCompany Report #A0366482A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Unknown	Unevaluable Event		Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/23/02ISR Number: 3904642-9Report Type:Expedited (15-DaCompany Report #B0101413A

Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Grand Mal Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Status Epilepticus					
per day	22	DAY		Diazepam	C		
UNKNOWN				Tramadol	C		ORAL
				Alcohol	C		

Date:04/23/02ISR Number: 3904644-2Report Type:Expedited (15-DaCompany Report #B0264307A

Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Clonic Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
3000MG Single		Completed Suicide					
Initial or Prolonged		Intentional Misuse		Lexomil	SS		ORAL
dose	1	DAY		Seresta	SS		ORAL
1	DAY			Alcohol	C		
1	DAY						
		Muscle Contractions					
		Involuntary					
		Somnolence					

Date:04/23/02ISR Number: 3904645-4Report Type:Expedited (15-DaCompany Report #B0264464A

Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	9	DAY		Lexomil	SS		ORAL
9	DAY						
		Insomnia					
		Nervousness					

Date:04/23/02ISR Number: 3904647-8Report Type:Expedited (15-DaCompany Report #B0264957A  
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Twice per day	40 DAY	Cardiac Arrest Ventricular Fibrillation		Zyban	PS	Glaxo Wellcome	ORAL
5MG Unknown				Digoxin	C	Glaxo Wellcome	ORAL
75MG Unknown				Lisinopril	C		ORAL
100MG Unknown				Aspirin	C		ORAL
				Thyroxine	C	Glaxo Wellcome	ORAL

Date:04/23/02ISR Number: 3904649-1Report Type:Expedited (15-DaCompany Report #B0265519A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day Initial or Prolonged		Cholestasis Focal Nodular Hyperplasia Liver Disorder Systemic Lupus Erythematous Rash		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/24/02ISR Number: 3905736-4Report Type:Expedited (15-DaCompany Report #A0062462A  
Age:61 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Diplopia Eye Disorder Vision Blurred	Health Professional	Zyban	PS	Glaxo Wellcome	ORAL
				Zestril	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Triamterene C Glaxo Wellcome  
Prinivil C

Date:04/24/02ISR Number: 3905741-8Report Type:Expedited (15-DaCompany Report #A0366054A  
Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	39 DAY	Completed Suicide		Wellbutrin	PS	Glaxo Wellcome	ORAL
	25MG Per day	Drug Ineffective		Effexor	C		ORAL
	400MG Twice per day	Libido Decreased		Neurontin	C		ORAL
		Overdose					
		Pulmonary Congestion					
		Sexual Dysfunction					
		Toxicologic Test Abnormal					
		Urinary Retention					

Date:04/24/02ISR Number: 3905742-XReport Type:Expedited (15-DaCompany Report #A0366092A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	42 DAY	Abortion Spontaneous		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Complications Of Maternal Exposure To Therapeutic Drugs					
		Maternal Drugs Affecting Foetus					

Date:04/24/02ISR Number: 3905743-1Report Type:Expedited (15-DaCompany Report #A0366462A  
Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Twice	Neutropenia		Wellbutrin	PS	Glaxo Wellcome	ORAL

Initial or Prolonged Pneumonia Necrotising  
per day 6 MON  
White Blood Cell Count  
Abnormal

Levothyroxine C Glaxo Wellcome ORAL

Date:04/24/02ISR Number: 3905744-3Report Type:Expedited (15-DaCompany Report #A0366463A  
Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Per day		Dystonia		Haldol	SS		ORAL
1 DAY		Fatigue		Zoloft	C		
UNKNOWN	100MG Per day	1 YR					
2 WK		Gaze Palsy		Oral Contraceptives	C		ORAL
		Headache					
		Muscle Spasms					
		Musculoskeletal Stiffness					
		Oculogyration					
		Pruritus					
		Rash					
		Viral Infection					
		Vision Blurred					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/25/02ISR Number: 3906203-4Report Type:Expedited (15-DaCompany Report #A0364704A

Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	47	DAY					
		Hair Disorder					
		Hair Growth Abnormal					
		Thyroid Disorder					

Date:04/25/02ISR Number: 3906212-5Report Type:Expedited (15-DaCompany Report #B0133710A

Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Angioneurotic Oedema		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	17	DAY					
Hospitalization -		Arthralgia					
Initial or Prolonged		Tendon Disorder					
Disability		Urticaria					
Other							

Date:04/25/02ISR Number: 3906213-7Report Type:Expedited (15-DaCompany Report #B0134007A

Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Csf Protein Increased		Zyban	PS	Glaxo Wellcome	ORAL
92	DAY						
Initial or Prolonged		Headache					
Disability		Hypertension					
		Meningeal Disorder					
		Mental Disorder					
		Musculoskeletal Stiffness					
		Pain					
		Vomiting Projectile					

Date:04/25/02ISR Number: 3906215-0Report Type:Expedited (15-DaCompany Report #B0265714A

Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged dosage text	8 DAY	Anxiety Blood Creatine Phosphokinase Increased Chest Pain Dry Mouth Tachycardia Tremor		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/25/02ISR Number: 3907615-5Report Type:Direct Company Report #CTU 166660  
Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 AM 1 NOON		Abdominal Pain Anxiety		Wellbutrin 100 Generic	PS		
1 5 PM		Cerebrovascular Accident Depression		Wellbutrin 75mg Generic	SS		
		Fear Hallucination, Visual Psychotic Disorder Vomiting		Cardiac Medications	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/26/02ISR Number: 3906912-7Report Type:Expedited (15-DaCompany Report #B0099525A  
 Age:60 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Grand Mal Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
300MG Per day 8 DAY						
Initial or Prolonged	Lung Neoplasm					

Date:04/26/02ISR Number: 3906915-2Report Type:Expedited (15-DaCompany Report #B0262805A  
 Age: Gender:Unknown I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Death		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/26/02ISR Number: 3908559-5Report Type:Expedited (15-DaCompany Report #B0261797A  
 Age:34 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Life-Threatening	Diplopia	Foreign	Zyban			
Other	Epilepsy	Other	(Bupropion			
	Status Epilepticus		Hydrochloride)	PS		ORAL
150 MG/ TWICE						
PER DAY/ ORAL						
			Multivitamin Capsule			
			(Multivitamin)	SS		ORAL
			Ephedrine			
			(Formulation			
			Unknown) (Ephedrine)	SS		
			Guarana (Formulation			
			Unknown) (Guarana)	SS		ORAL
ORAL			Neuroleptic	C		
			Antidepressant	C		
			Benzodiazepines	C		



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Dizziness Dysarthria Medication Error Vision Blurred	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS	Wellbutrin	ORAL
150 MG / SEE							
DOSAGE TEXT /							
ORAL							
				Thyroxine Sodium	C		
				Metformin			
				Hydrochloride	C		
				Irbesartan	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent RESPIRATORY Impairment/Damage (INHALATION)	150	Asthma Hypoaesthesia Hypoaesthesia Oral	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
MG/INHALED							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/26/02ISR Number: 3910106-9Report Type:Periodic  
Age:59 YR Gender:Male I/FU:I

Company Report #01P-163-0106856-00

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anxiety Circulatory Collapse Drug Level Increased	Other	Depakote (Divalproex Sodium) (Divalproex Sodium)	PS		ORAL
1000 MG, PER ORAL		Parkinson'S Disease					
150 MG, 2 IN 1 D				Bupropion Hydrochloride	SS		
20 MG, 1 IN 1 D, PER ORAL				Paroxetine Hydrochloride	SS		ORAL
20 MG, 1 IN 1 D, PER ORAL				Citalopram	SS		ORAL
				Paroxetine Hydrochloride	C		

Date:04/26/02ISR Number: 3923573-1Report Type:Periodic  
Age:66 YR Gender:Female I/FU:F

Company Report #A127209

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 250.00 MCG		Anxiety	Consumer	Tikosyn Capsules	PS		ORAL
TOTAL BID ORAL		Depression	Health				
75.00 MG		Diarrhoea	Professional				
TOTAL DAILY		Rash		Wellbutrin	SS		
		Tremor					
				Levoxyl	C		
				Valium	C		

Date:04/29/02ISR Number: 3907744-6Report Type:Expedited (15-DaCompany Report #A0366602A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Bundle Branch Block Left	Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
150MG Per day							
			Chest Pain	Celexa	C		
			Dyspnoea	Tricor	C		
			Electrocardiogram St	Ranitidine	C	Glaxo Wellcome	
			Segment Abnormal	Disopyramide	C		
25 YR							
				Aspirin	C		ORAL
81MG Per day							

Date:04/29/02ISR Number: 3907746-XReport Type:Expedited (15-DaCompany Report #A0366645A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Hallucination, Olfactory	Zyban	PS	Glaxo Wellcome	
			Temporal Lobe Epilepsy				

Date:04/29/02ISR Number: 3907749-5Report Type:Expedited (15-DaCompany Report #B0259252A  
Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT
Hospitalization -		Cheilitis
Initial or Prolonged		Dermatitis Bullous
		Hypopyon
		Neutrophilia
		Pyrexia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Rash Erythematous Rash Pustular Skin Inflammation	Report Source	Product	Role	Manufacturer	Route
150MG Per day	12 DAY			Zyban	PS	Glaxo Wellcome	ORAL
1UNIT per day	1 DAY			Propofan	SS		ORAL
40MG per day	277 DAY			Prazepam	C		ORAL
1UNIT per day	277 DAY			Zolpidem	C		ORAL
.25MG As required	277 DAY			Alprazolam	C		ORAL

Date:04/29/02ISR Number: 3907751-3Report Type:Expedited (15-DaCompany Report #B0262674A  
Age:39 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Abdominal Pain Aortic Aneurysm Rupture Chest Pain Dyspnoea Malaise Nervousness Sudden Death		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/29/02ISR Number: 3907752-5Report Type:Expedited (15-DaCompany Report #B0263361A  
Age:25 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide		Zyban	PS	Glaxo Wellcome	
UNKNOWN		150MG	Unknown					
Other			Depression Overdose		No Concurrent Medications	C		

Date:04/29/02ISR Number: 3907754-9Report Type:Expedited (15-DaCompany Report #B0265665A  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Unknown 4 DAY	Cardiac Arrest		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged		Coronary Artery Dissection Ventricular Fibrillation		Ethinylloestradiol + Norgestimate	C		ORAL

Date:04/29/02ISR Number: 3907755-0Report Type:Expedited (15-DaCompany Report #B0265912A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	30 DAY	Circulatory Collapse Epilepsy Intracranial Aneurysm Loss Of Consciousness		Zyban	PS	Glaxo Wellcome	ORAL

Date:04/29/02ISR Number: 3910761-3Report Type:Periodic Company Report #NSADSS2001031860  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Convulsion	Health	Ultram	PS		ORAL
ORAL		Drug Interaction	Professional	Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/30/02ISR Number: 3908288-8Report Type:Expedited (15-DaCompany Report #B0262751A  
Age:37 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 12 DAY Initial or Prolonged		Pharyngolaryngeal Pain	Zyban	PS	Glaxo Wellcome	ORAL
		Respiratory Disorder Respiratory Failure				

Date:04/30/02ISR Number: 3908289-XReport Type:Expedited (15-DaCompany Report #B0263835A  
Age:33 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 14 DAY Initial or Prolonged		Difficulty In Walking	Zyban	PS	Glaxo Wellcome	ORAL
		Discomfort Dysphagia Hypovolaemia Lymphocyte Count Decreased Neutrophil Count Decreased Oedema Peripheral Pain Red Blood Cell Sedimentation Rate Increased Respiratory Disorder Skin Lesion Urticaria Generalised White Blood Cell Count Increased				

Date:04/30/02ISR Number: 3908292-XReport Type:Expedited (15-DaCompany Report #B0264397A  
Age:46 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200MG Per day		Alanine Aminotransferase Increased	Zyban Lipanthyl	PS SS	Glaxo Wellcome	ORAL ORAL

Blood Cholesterol  
Increased  
Diabetes Mellitus  
Non-Insulin-Dependent  
Drug Interaction  
Gamma-Glutamyltransferase  
Increased

Date:04/30/02ISR Number: 3908296-7Report Type:Expedited (15-DaCompany Report #B0266376A  
Age:52 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - UNKNOWN Initial or Prolonged	Atrial Fibrillation Fatigue Malaise Sleep Disorder		Zyban Proton Pump Inhibitor	PS C	Glaxo Wellcome	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/30/02ISR Number: 3909116-7Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #CTU 166884

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 1 BID, 150MG SR ORAL	Crying Depression Emotional Disorder Mood Altered Suicidal Ideation		Bupropion 150mg 1 Bid , Glaxo  Vioxx Aleve	PS  C C	Glaxo	ORAL

Date:05/01/02ISR Number: 3909189-1Report Type:Expedited (15-DaCompany Report #A0170371A  
 Age:43 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Per day Initial or Prolonged	Cerebrovascular Accident Chills Convulsion Pyrexia Speech Disorder Weight Decreased	Health Professional	Zyban Influenza Vaccine	PS SS	Glaxo Wellcome Glaxo Wellcome	ORAL

Date:05/01/02ISR Number: 3909192-1Report Type:Expedited (15-DaCompany Report #A0366635A  
 Age:38 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 300MG per day	Atrial Flutter Blindness Transient Dyspnoea		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:05/01/02ISR Number: 3909194-5Report Type:Expedited (15-DaCompany Report #A0367027A  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						



Other Myasthenia Gravis Wellbutrin PS Glaxo Wellcome ORAL  
 150MG Twice  
 per day 10 DAY  
 No Concurrent Medication C

Date:05/01/02ISR Number: 3909195-7Report Type:Expedited (15-DaCompany Report #A0367126A  
 Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion Intentional Misuse Suicide Attempt Vomiting		Wellbutrin Sr Ritalin	PS C	Glaxo Wellcome	ORAL ORAL

Date:05/01/02ISR Number: 3909200-8Report Type:Expedited (15-DaCompany Report #B0266132A  
 Age: Gender:Female I/FU:I

Outcome	PT
Disability	Abdominal Pain Abdominal Pain Upper Chest Wall Pain Dizziness Drug Interaction Fall

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Fatigue Headache Nausea	Health	Zyban	PS	Glaxo Wellcome	ORAL
		Pain In Extremity	Professional	Tranxene	SS		ORAL
		Rib Fracture Road Traffic Accident Syncope Tendonitis		Unknown Selective Serotonin Reuptake Inhibitor	SS		ORAL

Date:05/01/02ISR Number: 3909203-3Report Type:Expedited (15-DaCompany Report #B0266558A  
Age:52 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1TAB Per day	3 DAY	Death		Zyban	PS	Glaxo Wellcome	ORAL
UNKNOWN					Fenoterol	C		
UNKNOWN					Formoterol	C		
UNKNOWN					Fluticasone	C	Glaxo Wellcome	

Date:05/01/02ISR Number: 3911535-XReport Type:Direct Company Report #CTU 167177  
Age: Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	100MG BID		Agitation		Wellbutrin Sr 100mg	PS		ORAL
ORAL					Depakote	C		
					Augmentin	C		
					Paxil	C		

Date:05/02/02ISR Number: 3909769-3Report Type:Expedited (15-DaCompany Report #A0367102A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged dosage text		Abdominal Distension		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Cholelithiasis					
		Constipation		Birth Control Pill Valtrex	C C	Glaxo Wellcome	ORAL ORAL
500MG As  required							

Date:05/02/02ISR Number: 3909780-2Report Type:Expedited (15-DaCompany Report #B0260822A  
Age:66 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN		International Normalised 75 DAY Ratio Increased		Zyban	PS	Glaxo Wellcome	
				Warfarin	SS	Glaxo Wellcome	
				Spirolactone	C		ORAL
25MG Per day				Enalapril	C		ORAL
5MG Per day				Unspecified Medication	C		
UNKNOWN				Furosemide	C	Glaxo Wellcome	ORAL
40MG Per day							

Date:05/02/02ISR Number: 3911428-8Report Type:Direct Company Report #CTU 167198  
Age:38 YR Gender:Female I/FU:I

Outcome  
Hospitalization -  
Initial or Prolonged  
  
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Required Intervention to Prevent Permanent Dose Duration Impairment/Damage	PT	Report Source	Product	Role	Manufacturer	Route
150MG PO QD	Confusional State Fall		Wellbutrin Sr 150mg (Glaxo Wellcome)	PS	Glaxo Wellcome	ORAL
50MG PO Q 1HR	Gaze Palsy Grand Mal Convulsion		Seroquel 25mg (Astra Zeneca)	SS	Astra Zeneca	ORAL
PRN	Lethargy		Birth Control Pills (Necon 0.5/35) Lorazepam	C C		

Date:05/03/02ISR Number: 3910249-XReport Type:Expedited (15-DaCompany Report #B0102171A  
Age:25 YR Gender:Female I/FU:F

Outcome Dose Duration Hospitalization - UNKNOW	PT	Report Source	Product	Role	Manufacturer	Route
Initial or Prolonged	Alanine Aminotransferase 2 DAY Increased Burning Sensation Decreased Appetite Dermatitis Exfoliative Dry Skin Fatigue Hyperhidrosis Neutrophilia Pain Pain Of Skin Pruritus Rash Maculo-Papular Skin Discolouration Vision Blurred Weight Decreased	Health Professional	Zyban No Concurrent Medications	PS C	Glaxo Wellcome	

Date:05/03/02ISR Number: 3910256-7Report Type:Expedited (15-DaCompany Report #B0265412A  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Aggression	Consumer	Zyban	PS	Glaxo Wellcome	
UNKNOWN	300MG per day	6 DAY					
Initial or Prolonged		Agitation					

Date:05/03/02ISR Number: 3910257-9Report Type:Expedited (15-DaCompany Report #B0265413A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Aggression	Consumer	Zyban	PS	Glaxo Wellcome	
UNKNOWN	2TABS per day	8 DAY					
Initial or Prolonged		Agitation		Celecoxib	C		
UNKNOWN	200MG per day	Emotional Disorder Headache Mood Swings					

Date:05/03/02ISR Number: 3910259-2Report Type:Expedited (15-DaCompany Report #B0266443A  
Age: Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Abnormal Behaviour
Initial or Prolonged	Atrial Fibrillation Brain Neoplasm

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Irritability Speech Disorder	Report Source	Product	Role	Manufacturer	Route
5	WK			Zyban	PS	Glaxo Wellcome	ORAL

Date:05/03/02ISR Number: 3910262-2Report Type:Expedited (15-DaCompany Report #B0266566A  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyntabac	PS	Glaxo Wellcome	
Other		Bundle Branch Block Left					
UNKNOWN		Myalgia		Contraceptives	C		
UNKNOWN							

Date:05/03/02ISR Number: 3910274-9Report Type:Expedited (15-DaCompany Report #NSADSS2002014533  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Risperidone	PS		ORAL
Hospitalization - Initial or Prolonged		Blood Ph Decreased		Divalproex	SS		ORAL
Other		Depressed Level Of		Bupropion	SS		
UNKNOWN		Consciousness					
		Drug Level Increased					
		Haemodialysis					
		Intentional Misuse					
		Respiratory Depression					
		Restlessness					

Date:05/03/02ISR Number: 3912130-9Report Type:Direct Company Report #CTU 167331  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS		
Other		Psychotic Disorder					

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Required ORAL	Blood Ph Decreased Confusional State Depressed Level Of Consciousness Drug Level Above Therapeutic Haemodialysis	Literature Health Professional	Risperidone (Unspecified) (Risperidone)	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage 180 TABLE, TOTAL, ORAL	Overdose Respiratory Depression Restlessness		Divalproex (Valproate Semisodium) Bupropion (Amfebutamone)	SS		ORAL

Outcome	PT
Other	Convulsion Crying Delusion Hallucinations, Mixed Headache

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	14 DAY	Memory Impairment Mood Swings Panic Attack Paranoia Schizoaffective Disorder	Zyban	PS	Glaxo Wellcome	ORAL
			Norplant	C		
			Synthroid	C	Glaxo Wellcome	
			Xanax	C		

Date:05/07/02ISR Number: 3911817-1Report Type:Expedited (15-DaCompany Report #A0365391A  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG See dosage text	11 DAY	PT Dysphagia Pulmonary Hypertension Throat Tightness	Zyban	PS	Glaxo Wellcome	ORAL

Date:05/07/02ISR Number: 3911837-7Report Type:Expedited (15-DaCompany Report #B0266864A  
Age:67 YR Gender:Male I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN Initial or Prolonged UNKNOWN UNKNOWN	32 DAY	PT Abdominal Pain Upper Blood Creatinine Increased Blood Urea Increased	Zyntabac Furosemide Doxazosine Eprosartan	PS C C C	Glaxo Wellcome Glaxo Wellcome Glaxo Wellcome	
		Hyperhidrosis Pyrexia				

Date:05/07/02ISR Number: 3911838-9Report Type:Expedited (15-DaCompany Report #B0266914A  
Age:55 YR Gender:Male I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Asthenia		Bupropion Hcl	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	7 DAY	Convulsion Dry Mouth Dry Throat Hyperhidrosis Insomnia Palpitations					

Date:05/07/02ISR Number: 3911839-0Report Type:Expedited (15-DaCompany Report #B0266928A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN	150MG	Convulsion See		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged dosage text		Intentional Misuse Suicide Attempt					

Date:05/07/02ISR Number: 3911840-7Report Type:Expedited (15-DaCompany Report #B0266930A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN		Psoriasis		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/07/02ISR Number: 3911841-9Report Type:Expedited (15-DaCompany Report #D0038216A  
 Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Unknown 27 DAY	Abortion Spontaneous		Zyban	PS	Glaxo Wellcome	ORAL
		Complications Of Maternal Exposure To Therapeutic Drugs					
		Maternal Drugs Affecting Foetus Pregnancy					

Date:05/07/02ISR Number: 3912989-5Report Type:Direct Company Report #CTU 167418  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2 WK Initial or Prolonged		Stevens-Johnson Syndrome		Wellbutrin	PS		

Date:05/07/02ISR Number: 3913646-1Report Type:Expedited (15-DaCompany Report #A103444  
 Age:79 YR Gender:Female I/FU:F

Outcome	PT
Required Intervention to Prevent Permanent Impairment/Damage	Anger
	Apathy
	Autoimmune Thyroiditis
	Balance Disorder
	Blood Oestrogen Decreased
	Blood Pressure Increased
	Blood Thyroid Stimulating
	Hormone Decreased
	Cerebral Haemorrhage
	Contusion
	Convulsion
	Dementia
	Drug Ineffective
	Dysphonia
	Emotional Disorder
	Epistaxis

Fall  
Head Injury  
Headache  
Heart Rate Decreased  
Heart Rate Increased  
Hostility  
Hyperhidrosis  
Laceration  
Meningioma  
Nervous System Disorder  
Nuclear Magnetic  
Resonance Imaging  
Abnormal  
Overdose  
Palpitations  
Parkinson'S Disease  
Personality Disorder  
Prothrombin Time  
Prolonged  
Syncope  
Thinking Abnormal

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
50.00 MG		Thyroid Function Test Abnormal	Consumer	Zoloft Tablets	PS		
TOTAL:DAILY		Transient Ischaemic Attack					
300.00 MG		Treatment Noncompliance		Wellbutrin	SS		ORAL
TOTAL:BID:ORA		Tremor					
L		Vaginal Haemorrhage		Coumadin	SS		
300.00 MG				Effexor Xr	SS		ORAL
TOTAL:BID:ORA							
L				Toprol	C		
				Zebeta	C		
				Lanoxin	C		
				Lasix	C		
				Synthroid	C		
				Vitamin E	C		
				Estrogen	C		
				Amantadine	C		
				Zocor	C		
				Estradiol Vaginal Ring	C		

Date:05/07/02ISR Number: 3913906-4Report Type:Expedited (15-DaCompany Report #A0367423A

Age: Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG/PER		Burning Sensation	Health	Wellbutrin Sr			
DAY/ORAL		Stevens-Johnson Syndrome	Professional	Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
		Stress					
		Vision Blurred					

Date:05/08/02ISR Number: 3913130-5Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 167539

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Delusion		Zyban 150mg			
Other		Disturbance In Attention		Glaxowellcome	PS	Glaxowellcome	ORAL
ONE TABLE							
FIRST 3 DAYS		Hallucinations, Mixed					
ORAL		Memory Impairment					
		Mood Swings					
		Panic Attack					
		Tinnitus					

Date:05/09/02ISR Number: 3913387-0Report Type:Expedited (15-DaCompany Report #A0367475A  
Age:47 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Angina Pectoris
Initial or Prolonged	Anxiety
	Blindness
	Cardiac Murmur
	Cardiac Valve Disease
	Chest Pain
	Dizziness
	Dyspnoea

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Hypoaesthesia Myocardial Infarction Thrombosis	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	1 DAY						

Date:05/09/02ISR Number: 3913400-0Report Type:Expedited (15-DaCompany Report #A0368187A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
Other		Hepatitis C					
150MG Twice							
per day							

Date:05/09/02ISR Number: 3913409-7Report Type:Expedited (15-DaCompany Report #B0267211A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxo Wellcome	ORAL
Other		Depression					
300MG per day	17 DAY			Norgestrel + Oestrone	C		ORAL
		Emotional Disorder Psychotic Disorder		Paracetamol	C	Glaxo Wellcome	ORAL
2TAB As							
required							

Date:05/09/02ISR Number: 3913429-2Report Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-11850310  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Sustiva Caps 600 Mg	PS	Bristol-Myers Squibb Company	ORAL
Hospitalization - Initial or Prolonged		Pancreatitis Acute	Health Professional	Stavudine	SS	Bristol-Myers Squibb Company	
				Didanosine	SS	Bristol-Myers Squibb	

Wellbutrin

SS

Company

Date:05/10/02ISR Number: 3913824-1Report Type:Expedited (15-DaCompany Report #A0360688A  
Age:9 DY Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly	9 MON	Dacryostenosis Congenital	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
		Drug Exposure Via Breast Milk Lethargy Maternal Drugs Affecting Foetus Somnolence	Professional	Vitamins	C		ORAL

Date:05/10/02ISR Number: 3913841-1Report Type:Expedited (15-DaCompany Report #B0264074A  
Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day 1 MON	Angina Unstable	Health	Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -	1MG Per day	Electrocardiogram St	Professional	Hyperium	C		ORAL
Initial or Prolonged	200MG Per day	Segment Abnormal		Fenofibrate	C		ORAL
	300MG Per day	Myocardial Infarction		Piascledine	C		ORAL
	10MG Per day			Stilnox	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/10/02ISR Number: 3913843-5Report Type:Expedited (15-DaCompany Report #B0266376A  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN	150MG Per day	14 DAY	Health	Zyban	PS	Glaxo Wellcome	
Initial or Prolonged Other		Condition Aggravated Fatigue Malaise Sleep Disorder	Professional	Proton Pump Inhibitor	C		

Date:05/10/02ISR Number: 3913844-7Report Type:Expedited (15-DaCompany Report #B0266756A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Zyban	PS	Glaxo Wellcome	ORAL
60TAB Single dose		Completed Suicide Epilepsy Intentional Misuse Urinary Incontinence					

Date:05/10/02ISR Number: 3913850-2Report Type:Expedited (15-DaCompany Report #D0038493A  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Intentional Misuse	Health	Zyban	PS	Glaxo Wellcome	ORAL
60TAB Single dose	1 DAY	Suicide Attempt	Professional				
40TAB Single dose				Diclofenac	SS		ORAL
10TAB Single dose	1 DAY			Diphenhydramine Hydrochloride	SS	Glaxo Wellcome	ORAL



Date:05/13/02ISR Number: 3914784-XReport Type:Expedited (15-DaCompany Report #A0366092A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
42 DAY		Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus	Professional				

Date:05/13/02ISR Number: 3914793-0Report Type:Expedited (15-DaCompany Report #B0130322A  
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Fall	Health	Zyban	PS	Glaxo Wellcome	ORAL
11 DAY							
Initial or Prolonged		Food Craving	Professional	Aspirin	C		
500MG Per day		Hypoglycaemia Loss Of Consciousness Malaise Muscle Rigidity Scratch Trismus Visual Disturbance		Nicotine	C	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/13/02ISR Number: 3914795-4Report Type:Expedited (15-DaCompany Report #B0259252A  
Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day	12 DAY	C-Reactive Protein	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged 40MG per day	276 DAY	Increased		Prazepam	SS		ORAL
1UNIT per day	1 DAY	Cheilitis		Propofan	C		ORAL
1UNIT per day	276 DAY	Corneal Epithelium Disorder		Zolpidem	C		ORAL
.25MG As required	276 DAY	Dermatitis Bullous		Alprazolam	C		ORAL
		Hypopyon Inflammation Intertrigo Leukocytosis Oedema Pyrexia Rash Pustular Skin Necrosis					

Date:05/14/02ISR Number: 3915446-5Report Type:Expedited (15-DaCompany Report #A0366642A  
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Chronic Obstructive		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	MON	Pulmonary Disease					
		Drug Ineffective Pneumonia		Nitroquick	C	Glaxo Wellcome	

Date:05/14/02ISR Number: 3915453-2Report Type:Expedited (15-DaCompany Report #A0368437A  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other  
Abortion Spontaneous  
Complications Of Maternal  
Exposure To Therapeutic  
Drugs  
Maternal Drugs Affecting  
Foetus  
Wellbutrin  
PS  
Glaxo Wellcome  
ORAL

Date:05/14/02ISR Number: 3915458-1Report Type:Expedited (15-DaCompany Report #B0267248A  
Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	7 WK	Diabetes Mellitus		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Insulin-Dependent					

Date:05/14/02ISR Number: 3915459-3Report Type:Expedited (15-DaCompany Report #B0267249A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Diabetes Mellitus		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Insulin-Dependent					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/14/02ISR Number: 3916520-XReport Type:Direct  
Age:53 YR Gender:Male I/FU:I

Company Report #CTU 168054

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Bupropion			
150MG BID PO		Dizziness		(Wellbutrin Sr)	PS		ORAL
		Fall		Diphenhyramine	C		
		Limb Injury		Hydrocodone	C		
		Loss Of Consciousness		Acetaminophen	C		
				Warfarin	C		
				Salsalate	C		
				Ranitidine	C		

Date:05/14/02ISR Number: 3917098-7Report Type:Expedited (15-DaCompany Report #200205-0247 (0)  
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Grand Mal Convulsion	Health	Halls Cough Drops			
Initial or Prolonged		Medication Error	Professional	(Menthol)	PS		ORAL
ORAL TOPICAL		Memory Impairment		Wellbutrin			
100 MG BID,				(Bupropion			
PER ORAL				Hydrochloride)	SS		ORAL
				Alcohol (Nos)	SS		
				Neurontin			
				(Gabapentin)	C		
				Trileptal			
				(Oxcarbazepine)	C		

Date:05/15/02ISR Number: 3916035-9Report Type:Expedited (15-DaCompany Report #B0102510A  
Age:73 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased	Health	Zyban	PS	Glaxo Wellcome	
38 DAY		Convulsion	Professional	Celecoxib	C		

Dizziness  
Dry Mouth  
Dysgeusia  
Fatigue  
Thought Blocking  
Thyroid Disorder  
Weight Decreased

Date:05/15/02ISR Number: 3916036-0Report Type:Expedited (15-DaCompany Report #B0115005A  
Age:73 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 38 DAY		Asthenia  Blood Pressure Increased Convulsion Dysgeusia Fatigue Loss Of Consciousness Weight Decreased	Consumer	Zyban	PS	Glaxo Wellcome	

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Freedom Of Information (FOI) Report

Date:05/15/02ISR Number: 3916042-6Report Type:Expedited (15-DaCompany Report #B0266928A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Convulsion	Health	Zyban	PS	Glaxo Wellcome	
UNKNOW	150MG	See					
Initial or Prolonged dosage text		Hallucination	Professional				
		Intentional Misuse		Benzodiazepine	SS		
UNKNOW		Suicide Attempt					

Date:05/15/02ISR Number: 3916043-8Report Type:Expedited (15-DaCompany Report #B0266929A

Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Aphasia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
Hospitalization -		Clonic Convulsion					
per day	21 DAY						
Initial or Prolonged		Fall					
		Head Injury					
		Hyperglycaemia					
		Hypotension					
		Salivary Hypersecretion					
		Syncope					
		Tremor					

Date:05/15/02ISR Number: 3916046-3Report Type:Expedited (15-DaCompany Report #B0267268A

Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Drug Withdrawal Syndrome		Quomem	PS	Glaxo Wellcome	ORAL
150MG							
Variable dose	15 DAY	Exophthalmos					
		Somnolence					
		Swelling					

Date:05/16/02ISR Number: 3916659-9Report Type:Expedited (15-DaCompany Report #A0368118A  
 Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG	Per day		Abortion Spontaneous	Wellbutrin	PS	Glaxo Wellcome	ORAL
			Complications Of Maternal Exposure To Therapeutic Drugs Road Traffic Accident				

Date:05/16/02ISR Number: 3916664-2Report Type:Expedited (15-DaCompany Report #A0368606A  
 Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
150MG	Twice		Abdominal Distension	Wellbutrin	PS	Glaxo Wellcome	
Initial or Prolonged	per day		Blindness				
20MG	Per day		Chromatopsia	Prozac	C		
2MG	As		Headache	Ambien	C		
required	YR		Intestinal Obstruction	Xanax	C		
325MG	Per day		Retinal Artery Thrombosis				
				Aspirin	C		
				Estradiol	C		
				Flomax	C		
				Percocet	C		
				Flexeril	C		
				Proventil	C	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/16/02ISR Number: 3916669-1Report Type:Expedited (15-DaCompany Report #B0264464A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day 8 DAY	Completed Suicide	Health	Zyban	PS	Glaxo Wellcome	ORAL
1.5MG Per day 8 DAY		Insomnia	Professional	Lexomil	C		ORAL

Date:05/16/02ISR Number: 3916673-3Report Type:Expedited (15-DaCompany Report #B0267386A  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Twice	Aggression		Zyban	PS	Glaxo Wellcome	ORAL
per day	32 DAY	Depressed Mood					
		Somnolence					

Date:05/16/02ISR Number: 3916675-7Report Type:Expedited (15-DaCompany Report #B0267494A  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG UNKNOWN	Hypomania		Zyban	PS	Glaxo Wellcome	
UNKNOWN		Unknown		Blood Pressure Medication	C		
		Psychotic Disorder					

Date:05/16/02ISR Number: 3916676-9Report Type:Expedited (15-DaCompany Report #B0267624A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	UNKNOWN	Death		Zyban	PS	Glaxo Wellcome	



Date:05/17/02ISR Number: 3917161-0Report Type:Expedited (15-DaCompany Report #B0266930A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN	1 DAY	Hypersensitivity		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged	1 DAY	Malaise Psoriasis Rash					

Date:05/17/02ISR Number: 3917165-8Report Type:Expedited (15-DaCompany Report #B0268075A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1 DAY	1 DAY	Drug Level Increased		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	1 DAY	Intentional Misuse		Prozac	SS		ORAL
1 DAY		Logorrhoea		Previscan	SS		ORAL

Date:05/20/02ISR Number: 3918595-0Report Type:Expedited (15-DaCompany Report #B0265714A  
Age:37 YR Gender:Female I/FU:F

Outcome	PT
Other	Anxiety Blood Creatine Phosphokinase Increased Chest Pain Dry Mouth Flushing

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Tachycardia Tremor				
Dose	Duration		Report Source	Product	Role	Manufacturer
150MG See				Zyban	PS	Glaxo Wellcome
dosage text	10 DAY					ORAL

Date:05/21/02ISR Number: 3919044-9Report Type:Expedited (15-DaCompany Report #A0149265B  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Congenital Anomaly		Accidental Exposure	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
7.5MG Twice		Breech Presentation	Professional	Buspar	C		ORAL
per day		Caesarean Section					
100MG Per day		Congenital Eyelid		Zoloft	C		ORAL
		Malformation					
		Finger Deformity					
		Plagiocephaly					
		Pregnancy					
		Premature Labour					
		Transmission Of Drug Via					
		Semen					

Date:05/21/02ISR Number: 3920491-XReport Type:Direct Company Report #CTU 168514  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Arthritis		Zyban	PS		
2 WK							
Initial or Prolonged		Pain					
Disability		Urticaria					

Date:05/22/02ISR Number: 3919534-9Report Type:Expedited (15-DaCompany Report #A0175184A  
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Facial Palsy		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Per day	3 DAY	Muscle Contractions		Celexa	C		ORAL
40MG Per day		Involuntary		Ziac	C		ORAL
2.5MG Per day		Paraesthesia		Spironolactone	C		
5MG As				Reglan	C	Glaxo Wellcome	
required							
10MG Per day				Zyrtec	C	Glaxo Wellcome	ORAL

Date:05/22/02ISR Number: 3919537-4Report Type:Expedited (15-DaCompany Report #A0364704A  
Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Alopecia		Zyban	PS	Glaxo Wellcome	ORAL
per day	47 DAY	Hair Disorder					
		Hair Growth Abnormal					
		Thyroid Disorder					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/22/02ISR Number: 3919542-8Report Type:Expedited (15-DaCompany Report #A0369185A  
Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Per day		Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus					

Date:05/22/02ISR Number: 3919870-6Report Type:Direct Company Report #CTU 168634  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Angioneurotic Oedema Nasopharyngeal Disorder		Wellbutrin 150 Sr Glaxo	PS	Glaxo	ORAL
1 BID ORAL		Pharmaceutical Product Complaint		Prozac	C		

Date:05/22/02ISR Number: 3921343-1Report Type:Expedited (15-DaCompany Report #MK200205-0227-1  
Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Cerebrovascular Accident Communication Disorder Drug Interaction		Pamelor 25mg Capsules Wellbutrin-Slow Release	PS SS		
100 MG BID				Celexa (Citalopram)	SS		

Date:05/22/02ISR Number: 3922173-7Report Type:Expedited (15-DaCompany Report #A103444  
Age:79 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Required	Blood Pressure Increased	Consumer	Zoloft Tablets	PS	
50.00 MG					
Intervention to	Cerebral Haemorrhage				
TOTAL: DAILY					
Prevent Permanent	Contusion		Wellbutrin	SS	ORAL
300.00 MG					
Impairment/Damage	Dementia				
TOTAL: BID:					
	Drug Ineffective				
ORAL					
	Epistaxis		Coumadin	SS	
	Fall		Effexor Xr	SS	ORAL
300.00 MG					
TOTAL:	Head Injury				
BID:ORAL	Headache				
	Heart Rate Decreased		Toprol	C	
	Heart Rate Increased		Zebeta	C	
	Hyperhidrosis		Lanoxin	C	
	Overdose		Lasix	C	
	Parkinson'S Disease		Synthroid	C	
	Syncope		Vitamin E	C	
			Estrogen	C	
			Amantadine	C	
			Zocor	C	
			Estradiol Vaginal		
			Ring	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/22/02ISR Number: 3922257-3Report Type:Expedited (15-DaCompany Report #A210802

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Depression	Consumer	Zoloft Tablets	PS		
50.000 MG							
Intervention to		Grand Mal Convulsion					
TOTAL							
Prevent Permanent				Wellbutrin Sr	SS		ORAL
150.00 MG							
Impairment/Damage							
TOTAL:BID:ORA							

L

Date:05/22/02ISR Number: 3922545-0Report Type:Expedited (15-DaCompany Report #B0267143A

Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Acidosis	Foreign	Bupropion			
Initial or Prolonged		Agitation	Literature	Hydrochloride Tablet			
		Confusional State	Health	- Zyban (Bupropion			
		Delusion Of Grandeur	Professional	Hydrochloride)	PS		
		Grand Mal Convulsion					
		Hallucination, Visual					
		Overdose					
		Respiratory Rate					
		Increased					
		Sinus Tachycardia					
		Vomiting					

Date:05/23/02ISR Number: 3920481-7Report Type:Expedited (15-DaCompany Report #B0266864A

Age:67 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abdominal Pain Upper		Zyntabac	PS	Glaxo Wellcome	
UNKNOWN		32 DAY					
Initial or Prolonged		Blood Creatinine		Furosemide	C	Glaxo Wellcome	
UNKNOWN							

UNKNOWN	Increased	Doxazosine	C	
UNKNOWN	Blood Urea Increased	Eprosartan	C	Glaxo Wellcome
	Hyperhidrosis			
	Pyrexia			

Date:05/23/02ISR Number: 3920964-XReport Type:Direct Company Report #CTU 168735  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour		Wellbutrin	PS		
200 MG	200 MG	Anger					
		Depression					
		Divorced					
		Drug Dependence					
		Feeling Abnormal					
		Job Dissatisfaction					
		Paranoia					
		Parent-Child Problem					
		Sleep Disorder					
		Weight Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/24/02ISR Number: 3921498-9Report Type:Expedited (15-DaCompany Report #A0368754A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:05/24/02ISR Number: 3921506-5Report Type:Expedited (15-DaCompany Report #B0268007A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Abdominal Distension Blood Glucose Abnormal Chest Pain		Zyban	PS	Glaxo Wellcome	ORAL

Date:05/24/02ISR Number: 3921507-7Report Type:Expedited (15-DaCompany Report #B0268089A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN		Facial Palsy		Zyban	PS	Glaxo Wellcome	

Date:05/24/02ISR Number: 3921511-9Report Type:Expedited (15-DaCompany Report #B0268467A  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG	Variable dose 9 DAY	Depressed Mood Feeling Drunk Suicidal Ideation		Zyban	PS	Glaxo Wellcome	ORAL

Date:05/28/02ISR Number: 3923289-1Report Type:Expedited (15-DaCompany Report #A0369795A  
Age:25 YR Gender:Male I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Conjunctival Hyperaemia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Vision Blurred					
per day							

Date:05/28/02ISR Number: 3923292-1Report Type:Expedited (15-DaCompany Report #B0260822A  
Age:66 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Health	Zyban	PS	Glaxo Wellcome	
UNKNOWN		75 DAY					
		International Normalised	Professional	Warfarin	SS	Glaxo Wellcome	
UNKNOWN		Ratio Increased		Spiroinolactone	C		ORAL
25MG Per day				Enalapril	C		ORAL
5MG Per day				Unspecified Medication	C		
UNKNOWN				Furosemide	C	Glaxo Wellcome	ORAL
40MG Per day				Ethanol	C		
UNKNOWN							

Date:05/29/02ISR Number: 3924107-8Report Type:Expedited (15-DaCompany Report #B0263173A  
Age:42 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Areflexia
Initial or Prolonged	Depressed Level Of Consciousness
	Dysphasia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	5 DAY	Epilepsy Hemiparesis Hemiplegia Paraesthesia	Zyban	PS	Glaxo Wellcome	ORAL

Date:05/29/02ISR Number: 3924108-XReport Type:Expedited (15-DaCompany Report #B0263835A  
Age:33 YR Gender:Male I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 14 DAY Initial or Prolonged		Difficulty In Walking Dysphagia Hypovolaemia Lymphocyte Percentage Decreased Neutrophil Percentage Decreased Oedema Peripheral Pain Painful Respiration Paralysis Peak Expiratory Flow Rate Decreased Urticaria Generalised White Blood Cell Count Increased	Zyban	PS	Glaxo Wellcome	ORAL

Date:05/29/02ISR Number: 3924111-XReport Type:Expedited (15-DaCompany Report #B0268361A  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN per day .25MG per day	150MG Twice	Confusional State Dizziness Insomnia	Zyban Alprazolam	PS C	Glaxo Wellcome	

Sensory Disturbance

Date:05/29/02ISR Number: 3925257-2Report Type:Direct  
 Age:43 YR Gender:Female I/FU:I

Company Report #CTU 169081

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash Pruritic		Zyban 150 Mg Catalytica	PS	Catalytica	ORAL
150 MG QD							
ORAL							
				Albuterol	C		
				Glucophage	C		
				Cetaphil	C		
				Calcium	C		
				Xanax	C		
				Nicoderm	C		

Date:05/29/02ISR Number: 3925280-8Report Type:Direct  
 Age:46 YR Gender:Female I/FU:I

Company Report #CTU 169088

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash Generalised Rash Pruritic		Zyban 150 Mg Catalytica	PS	Catalytica	ORAL
150 MG BID							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Synthroid C

Date:05/29/02ISR Number: 3925893-3Report Type:Direct  
 Age:35 YR Gender:Female I/FU:I

Company Report #CTU 169143

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pharmaceutical Product		Wellbutrin Sr 100 Mg	PS		
2 TABS BID		Complaint					
		Rash					

Date:05/30/02ISR Number: 3925597-7Report Type:Expedited (15-DaCompany Report #B0267897A  
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Zyban	PS	Glaxo Wellcome	ORAL
49 DAY		Panic Attack		Nicorette	C	Glaxo Wellcome	
SUBLINGUAL	2MG Five	Suicidal Ideation					
times per day				Ethanol	C		ORAL
4UNIT per day							

Date:05/30/02ISR Number: 3925600-4Report Type:Expedited (15-DaCompany Report #B0268687A  
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Fall					
per day		Malaise		Contraceptive (Unspecified)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600MG/DAY		Aggression		Zyban ( Bupropion )	PS		
		Agitation		Wellbutran	SS		
		Confusional State		Bupropion	SS		
		Convulsion					
		Gastroenteritis Viral					
		Medication Error					
		Vomiting					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Foreign	Depo-Medrone			
		Drug Interaction	Health	(Methylprednisolone)			
		Lumbar Vertebral Fracture	Professional	Suspension, Sterile	PS		
INTRAVENOUS	30 MG,		Other				
SINGLE, IV				Bupropion			
				(Amfebutamone)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/02ISR Number: 3926325-1Report Type:Expedited (15-DaCompany Report #A0370078A

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Drug Ineffective Drug Interaction Drug Level Decreased Suicide Attempt		Wellbutrin Protease Inhibitor	PS SS	Glaxo Wellcome	ORAL

Date:05/31/02ISR Number: 3926421-9Report Type:Direct

Company Report #CTU 169263

Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 150 MG BID PO Prevent Permanent Impairment/Damage		Rash		Bupropion 75 Mg Mylan Lot 1k1074	PS	Mylan	ORAL

Date:05/31/02ISR Number: 3927112-0Report Type:Expedited (15-DaCompany Report #A0366054A

Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coma Completed Suicide Drug Ineffective Drug Toxicity	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
SEE IMAGE/ORAL		Libido Decreased					
		Overdose Pulmonary Congestion Toxicologic Test Abnormal Urinary Retention		Venlafaxine Hydrochloride Gabapentin Ethanol Fexofenadine Hydrochlorid Alprazolam Hyoscyamine	C C C C C C C		

Date:06/03/02ISR Number: 3926797-2Report Type:Expedited (15-DaCompany Report #B0266443A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 6 WK Initial or Prolonged		Abnormal Behaviour Aggression Aphasia Ganglioneuroma Heart Rate Irregular Irritability Sleep Disorder Speech Disorder		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/03/02ISR Number: 3926808-4Report Type:Expedited (15-DaCompany Report #B0269491A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN	150MG Twice	Facial Palsy		Zyban	PS	Glaxo Wellcome	
per day UNKNOWN				Atorvastatin	C		
UNKNOWN				Amlodipine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/03/02ISR Number: 3926811-4Report Type:Expedited (15-DaCompany Report #D0038591A

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Bowel Sounds Abnormal	Zyban	PS	Glaxo Wellcome	ORAL
16TAB Single							
dose	1	DAY	Coma				
			Intentional Misuse	Atosil	SS	Glaxo Wellcome	ORAL
10TAB Single							
dose	1	DAY	Suicide Attempt				
			Toxicologic Test Abnormal	Normoc	SS		ORAL
10TAB Single							
dose	1	DAY		Extr. Rad. Valerianae	SS		ORAL
10TAB Single							
dose	1	DAY		Hop Extract	SS		ORAL
10TAB Single							
dose	1	DAY		Alcohol	SS		ORAL
1	DAY						

Date:06/03/02ISR Number: 3928596-4Report Type:Expedited (15-DaCompany Report #A211508

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required			Consumer	Zoloft Tablets	PS		
Intervention to				Wellbutrin Sr	SS		ORAL
150.00 MG							
Prevent Permanent			Grand Mal Convulsion				
TOTAL: BID:							
Impairment/Damage							
ORAL							



Date:06/04/02ISR Number: 3927256-3Report Type:Expedited (15-DaCompany Report #A0370262A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Abnormal Dreams Anorexia Back Injury Crying Epistaxis Insomnia Mood Altered Nasal Odour Rhinorrhoea		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/05/02ISR Number: 3927994-2Report Type:Expedited (15-DaCompany Report #A0369011A  
Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Single dose		Dysphagia Foreign Body Trauma Oesophageal Spasm Retching		Wellbutrin Elavil	PS C	Glaxo Wellcome Glaxo Wellcome	ORAL

Date:06/05/02ISR Number: 3932576-2Report Type:Periodic Company Report #A0160250A  
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG / TWICE PER DAY / ORAL		Eye Swelling Pruritus Rash	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/05/02ISR Number: 3932578-6Report Type:Periodic  
Age:29 YR Gender:Male I/FU:I

Company Report #A0161127A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Aggression Anger Anxiety	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
100 MG / TWICE PER DAY / ORAL		Asthenia Coma Depression Fear Grand Mal Convulsion Impatience Limb Injury Loss Of Consciousness Marital Problem Memory Impairment Mood Altered Movement Disorder Pain In Extremity Personality Change Salivary Hypersecretion Snoring Somnolence Staring Tremor Vomiting		Amitriptyline	C		

Date:06/05/02ISR Number: 3932579-8Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #A0161473A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Eye Swelling Joint Swelling Nasopharyngitis	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE PER DAY/ORAL		Oedema Peripheral Urticaria		Tylenol Cold Medication (Tylenol			

ORAL Cold Medication) SS ORAL  
Neocon C

Date:06/05/02ISR Number: 3932580-4Report Type:Periodic Company Report #A0164564A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Fall Loss Of Consciousness	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE PER DAY/ORAL							

Date:06/05/02ISR Number: 3932582-8Report Type:Periodic Company Report #A0168570A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ PER DAY/ ORAL 1 WK							
Thyroxine Sodium C							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/05/02ISR Number: 3932584-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0168990A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Chest Pain	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE						
PER DAY/ORAL	1 MON					

Date:06/05/02ISR Number: 3932586-5Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0169235A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Chest Pain	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /						
TWICE PER DAY						
/ ORAL	1 MON					

Date:06/05/02ISR Number: 3932587-7Report Type:Periodic  
Age: Gender:Unknown I/FU:I

Company Report #A0169236A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Chest Pain	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE						
PER DAY/ORAL	1 MON					

Date:06/05/02ISR Number: 3932588-9Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #A0170001A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion	Health Professional Company	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE			Representative				
PER DAY/ORAL							

Date:06/05/02ISR Number: 3932589-0Report Type:Periodic Company Report #A0170679A  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent 150 MG/TWICE Impairment/Damage PER DAY/ORAL		Angioneurotic Oedema Arthralgia Asthenia Eyelid Oedema Face Oedema Insomnia Joint Stiffness Lip Pain Musculoskeletal Stiffness Pain Rash Erythematous Skin Lesion Swelling Tenderness Throat Irritation Urticaria	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)  Esterified Estrogens Medroxyprogesterone Ace.	PS  C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/05/02ISR Number: 3932591-9Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #A0106207A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent 150 MG/ TWICE Impairment/Damage PER DAY/ ORAL	5	DAY	Foreign	Zyban Tablet - Zylan (Bupropion Hydrochloride)	PS		ORAL
				Diltiazem Hydrochloride	C		
				Conjugated Estrogens	C		
				Hydrochlorothiazide	C		
				Atorvastatin Calcium	C		
				Nitroglycerine	C		
				Lorazepam	C		
				Pantoprazole	C		
				Lorazepam	C		

Date:06/05/02ISR Number: 3932592-0Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0106346A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent 300 MG/ PER Impairment/Damage DAY; ORAL	2	WK	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
				Liver Function Test Abnormal Musculoskeletal Disorder			
				Vomiting			

Date:06/05/02ISR Number: 3932593-2Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0106354A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent 300 MG / PER			Foreign	Zyban Tablet- Zyban ( Bupropion Hydrochloride)	PS		ORAL
				Chest Pain Extrasystoles			

Impairment/Damage  
DAY / ORAL 10 DAY

Date:06/05/02ISR Number: 3932594-4Report Type:Periodic Company Report #A0143678A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Anxiety	Health	Zyban Tablet- Zyban			
Intervention to		Hypersensitivity	Professional	( Bupropion			
Prevent Permanent		Pruritus	Company	Hydrochloride)	PS		ORAL
ORAL							
Impairment/Damage		Rash	Representative				
		Throat Tightness					
		Urticaria					

Date:06/05/02ISR Number: 3932595-6Report Type:Periodic Company Report #A0148626A  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Convulsion	Health	Zyban Tablet- Zyban			
Disability		Laceration	Professional	( Bupropion			
Other			Company	Hydrochloride)	PS		ORAL
150 MG/ TWICE							
			Representative				
PER DAY/ ORAL 2	YR						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/05/02ISR Number: 3932596-8Report Type:Periodic  
Age:35 YR Gender:Male I/FU:I

Company Report #A0149866A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Drug Level Changed Hallucination, Auditory	Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL		Loss Of Consciousness  Photopsia Road Traffic Accident Tremor		Meclozine (Formulation Unknown) (Meclozine)	SS		

Date:06/05/02ISR Number: 3932597-XReport Type:Periodic  
Age:26 YR Gender:Male I/FU:I

Company Report #A0151283A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to Prevent Permanent Impairment/Damage PER DAY/ ORAL	19 DAY	Face Oedema Joint Swelling Malaise  Pain Pruritus Pyrexia Rash Maculo-Papular Serum Sickness	Foreign	Zyban Tablet- Zyban ( Bupropion Hydrochloride)	PS		ORAL

Date:06/05/02ISR Number: 3932598-1Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #A0151863A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged  150 MG/ TWICE PER DAY/ ORAL		Dyspnoea Hypersensitivity Oropharyngeal Swelling  Urticaria	Health Professional	Zyban Tablet- Zyban ( Bupropion Hydrochloride)	PS		ORAL



Date:06/05/02ISR Number: 3932599-3Report Type:Periodic Company Report #A0156006A  
Age:50 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Back Pain	Consumer	Zyban Tablet - Zyban			
Initial or Prolonged	Chest Discomfort		(Bupropion			
	Chest Pain		Hydrochloride)	PS		ORAL
150 MG/ PER						
DAY/ ORAL	Pain In Extremity					

Date:06/05/02ISR Number: 3932600-7Report Type:Periodic Company Report #A0159085A  
Age:40 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Anaphylactic Reaction	Health	Zyban Tablet- Zyban			
	Dyspnoea	Professional	( Bupropion			
	Fatigue	Company	Hydrochloride)	PS		ORAL
150 MG/ PER						
DAY/ ORAL	Laryngeal Dyspnoea	Representative				
	Rash					
	Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/05/02ISR Number: 3944093-4Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #A0172431A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Duration Hypersensitivity	Consumer	Zyban Tablet- Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:06/05/02ISR Number: 3944096-XReport Type:Periodic  
Age:34 YR Gender:Male I/FU:I

Company Report #A0173889A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/ TWICE PER DAY/ ORAL	Duration Arthralgia Difficulty In Walking Eye Swelling Joint Swelling Rash Urticaria	Foreign Consumer	Zyban Tablet- Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:06/05/02ISR Number: 3944097-1Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #A0359127A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG/ TWICE PER DAY/ORAL	Duration Angioneurotic Oedema Face Oedema Rash Urticaria	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:06/05/02ISR Number: 3944098-3Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #A0360041A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Life-Threatening Required Intervention to 150 MG/THREE Prevent Permanent TIMES PER Impairment/Damage DAY/ORAL	Angioneurotic Oedema Asthenia Dizziness  Eye Swelling  Face Oedema  Hypotension Oedema Mouth Palpitations Pruritus Rash Rash Vesicular Tremor Urticaria Urticaria Generalised	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS	ORAL
			Ziac Gemfibrozil	C C	

Date:06/05/02ISR Number: 3944099-5Report Type:Periodic Company Report #A0361077A  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Blood Pressure Fluctuation Blood Pressure Increased	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/  SINGLE DOSE/  ORAL				Atorvastatin Calcium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/05/02ISR Number: 3944100-9Report Type:Periodic  
Age:35 YR Gender:Female I/FU:F

Company Report #A0137115A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Pressure Increased Confusional State Convulsion	Health Professional	Zyban Tablet - Zyban (Bupropion Hydrchloride)	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL		Disorientation Muscle Rigidity Pain Panic Reaction Rhinorrhoea Salivary Hypersecretion Tremor Urinary Incontinence Visual Acuity Reduced					

Date:06/05/02ISR Number: 3944101-0Report Type:Periodic  
Age:38 YR Gender:Female I/FU:F

Company Report #A0144963A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Chest Pain Dyspnoea Hypersensitivity	Consumer	Zyban Tablet - Zyban (Bupropion Hydrchloride)	PS		ORAL
150 MG/ TWICE PER DAY/ ORAL		Pruritus Rash Urticaria Vomiting					

Date:06/05/02ISR Number: 3944102-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A0145155A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Loss Of Consciousness	Consumer	Zyban Tablet - Zyban (Bupropion			

ORAL

Hydrchloride)

PS

ORAL

Ethanol (Alcohol)

SS

Date:06/06/02ISR Number: 3930360-7Report Type:Expedited (15-DaCompany Report #A103444

Age:79 YR Gender:Female I/FU:F

Outcome	PT
Required	Abnormal Behaviour
Intervention to	Anger
Prevent Permanent	Balance Disorder
Impairment/Damage	Blood Oestrogen Decreased
	Blood Pressure Increased
	Blood Thyroid Stimulating
	Hormone Decreased
	Cerebral Haemorrhage
	Cognitive Disorder
	Contusion
	Convulsion
	Drug Ineffective
	Dysphonia
	Ear Disorder
	Epistaxis
	Fall
	Head Injury

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
50.00 MG	TOTAL:DAILY	Headache Heart Rate Decreased Heart Rate Increased	Consumer	Zoloft Tablets	PS		
300.00 MG	TOTAL:BID:ORA	Hostility Hyperhidrosis		Wellbutrin	SS		ORAL
L		Injury Masked Facies					
300.00 MG	TOTAL:BID:ORA	Medication Error Meningioma Mood Swings		Coumadin Effexor Xr	SS SS		ORAL
L		Nuclear Magnetic Resonance Imaging					
		Abnormal Palpitations Parkinson'S Disease Prothrombin Time Prolonged Sluggishness Syncope Thyroxine Abnormal Transient Ischaemic Attack Treatment Noncompliance Tremor Vaginal Haemorrhage		Toprol Zebeta Lanoxin Lasix Synthroid Vitamin E Estrogen Amantadine Zocor Estradiol Vaginal Ring	C C C C C C C C C C		

Date:06/06/02ISR Number: 3933557-5Report Type:Expedited (15-DaCompany Report #B0264771A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to ORAL		Anxiety Depression	Foreign Health	Paxil (Paroxetine Hydrochloride)	PS		ORAL
Prevent Permanent Impairment/Damage		Malaise Nausea	Professional	Zyban Tablet (Bupropion			

1 TABLET/ Hydrochloride) SS ORAL  
 TWICE PER  
 DAY/ ORAL Nicotine C

Date:06/07/02ISR Number: 3929339-0Report Type:Expedited (15-DaCompany Report #B0268848A  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged RESPIRATORY		Angina Pectoris Blood Creatine		Zyban Becotide	PS C	Glaxo Wellcome Glaxo Wellcome	ORAL
(INHALATION)		Phosphokinase Increased					
RESPIRATORY		Chest Discomfort		Serevent	C	Glaxo Wellcome	
(INHALATION)		Chest Pain					
UNKNOWN		Dyspnoea		Ventoline	C	Glaxo Wellcome	
		Electrocardiogram St Segment Abnormal					

Date:06/07/02ISR Number: 3929340-7Report Type:Expedited (15-DaCompany Report #B0269619A  
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN	150MG	Impaired Driving Ability		Zyban	PS	Glaxo Wellcome	
		Per day 1 MON Labyrinthitis		Atenolol	C		
	.25MG per day						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

.5MG per day Alprazolam C

Date:06/07/02ISR Number: 3929341-9Report Type:Expedited (15-DaCompany Report #B0269621A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Zyban	PS	Glaxo Wellcome	ORAL
150MG per day							

Date:06/10/02ISR Number: 3930009-3Report Type:Expedited (15-DaCompany Report #A0169595A  
Age:24 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Antisocial Behaviour Apathy	Consumer	Wellbutrin Synthroid	PS C	Glaxo Wellcome Glaxo Wellcome	ORAL ORAL
.075MCG Per day		Hallucination, Auditory					
25MG Twice per day		Memory Impairment Nervousness		Ritalin	C		ORAL
15 DAY		Paranoia Pruritus		Vitamin Depakote	C C		ORAL ORAL
		Rash Erythematous Rash Vesicular Scar Suicidal Ideation					

Date:06/10/02ISR Number: 3930012-3Report Type:Expedited (15-DaCompany Report #A0367475A  
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG Per day 1 DAY Initial or Prolonged		Angina Pectoris Anxiety	Consumer	Zyban	PS	Glaxo Wellcome	ORAL



Blindness  
 Cardiac Murmur  
 Cardiac Valve Disease  
 Chest Pain  
 Dyspnoea  
 Hypoaesthesia  
 Myocardial Infarction  
 Thrombosis

Date:06/10/02ISR Number: 3930013-5Report Type:Expedited (15-DaCompany Report #A0370078A

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - YR Initial or Prolonged	Drug Ineffective Drug Interaction Drug Level Decreased Intentional Misuse Suicide Attempt	Consumer	Wellbutrin Protease Inhibitor Ethanol Ambien	PS SS SS	Glaxo Wellcome	ORAL  ORAL ORAL

Date:06/10/02ISR Number: 3930014-7Report Type:Expedited (15-DaCompany Report #A0370671A

Age:36 YR Gender:Female I/FU:F

Outcome	PT
Other	Fatigue Grand Mal Convulsion Hypercalcaemia

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Expectorant  
(Expectorant) SS

Date:06/11/02ISR Number: 3934027-0Report Type:Expedited (15-DaCompany Report #B0269609A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complex Partial Seizures Malaise	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		
UNKNOWN	3	WK					

Date:06/11/02ISR Number: 3934217-7Report Type:Expedited (15-DaCompany Report #B0269682A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ PER DAY/ ORAL		Drugs Maternal Drugs Affecting Foetus Pregnancy					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/11/02ISR Number: 3934219-0Report Type:Expedited (15-DaCompany Report #D0038599A

Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Foreign	Zyban Tablet - Zyban			
Life-Threatening		Loss Of Consciousness	Health	(Bupropion			
			Professional	Hydrochloride)	PS		ORAL
300 MG/ PER							
DAY/ ORAL				Antihypertensive	C		

Date:06/12/02ISR Number: 3931860-6Report Type:Expedited (15-DaCompany Report #A0364976A

Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cough		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	5 MON	Diarrhoea					
		Epistaxis					
		Gastrointestinal Disorder					
		Vomiting					

Date:06/12/02ISR Number: 3931863-1Report Type:Expedited (15-DaCompany Report #A0368754A

Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arteriosclerosis		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Completed Suicide					
		Convulsion					
		Drug Level Increased					
		Intentional Misuse					
		Pulmonary Congestion					
		Pulmonary Oedema					
		Somnolence					
		Urinary Incontinence					

Date:06/12/02ISR Number: 3931870-9Report Type:Expedited (15-DaCompany Report #B0107035A  
Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anorexia		Zyban	PS	Glaxo Wellcome	ORAL
2 WK		Anxiety		Chloroquine	C		
		Completed Suicide					
		Depression					
		Dissociation					
		Disturbance In Attention					
		Feeling Abnormal					
		Headache					
		Injury Asphyxiation					
		Irritability					
		Mood Swings					
		Palpitations					
		Vision Blurred					

Date:06/12/02ISR Number: 3931875-8Report Type:Expedited (15-DaCompany Report #B0269481A  
Age:56 YR Gender:Female I/FU:I

Outcome  
Life-Threatening  
Hospitalization -

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Initial or Prolonged

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	32 DAY	Ventricular Tachycardia		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/12/02ISR Number: 3931879-5Report Type:Expedited (15-DaCompany Report #B0270232A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 15 DAY Initial or Prolonged		Major Depression		Zyban	PS	Glaxo Wellcome	ORAL

Date:06/13/02ISR Number: 3932412-4Report Type:Expedited (15-DaCompany Report #A0173931A  
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Cholelithiasis Grand Mal Convulsion		Wellbutrin Singulair Levoxyl Prevacid Luvox	PS C C C C	Glaxo Wellcome	ORAL

Date:06/13/02ISR Number: 3932418-5Report Type:Expedited (15-DaCompany Report #A0370680A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Blood Pressure Increased Brain Oedema		Wellbutrin	PS	Glaxo Wellcome	ORAL

Glaucoma	Lipitor	C
Haematemesis	Estrace	C
Headache	Timoptic	C
Intracranial Pressure	Carbachol	C
Increased	Vagifem	C
Intraocular Pressure	Baby Aspirin	C
Increased	Protonix	C

Date:06/13/02ISR Number: 3932423-9Report Type:Expedited (15-DaCompany Report #A0371413A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
38 DAY		Gastrooesophageal Reflux		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
		Disease		Prozac	C		ORAL
		Retching		Nicoderm	C	Glaxo Wellcome	
TOPICAL							
.075MG		Thyroid Neoplasm		Levoxyl	C	Glaxo Wellcome	ORAL
Unknown							

Date:06/13/02ISR Number: 3932449-5Report Type:Expedited (15-DaCompany Report #B0270183A  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability							
300MG per day 41 DAY		Colitis Ulcerative		Zyntabac	PS	Glaxo Wellcome	ORAL
		Rectal Haemorrhage					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/13/02ISR Number: 3932452-5Report Type:Expedited (15-DaCompany Report #B0270350A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Twice				Zyban	PS	Glaxo Wellcome	ORAL
per day	35 DAY	Jaundice					

Date:06/14/02ISR Number: 3933152-8Report Type:Expedited (15-DaCompany Report #B0269764A

Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
20TAB Single				Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged							
dose	1 DAY	Clonic Convulsion					
14TAB Single		Convulsion		Deroxat	SS	Glaxo Wellcome	ORAL
dose	1 DAY	Depressed Level Of					
16TAB Single		Consciousness		Arestal	C		ORAL
dose	1 DAY	Intentional Misuse					
1 DAY		Somnolence		Nifuroxazide	C	Glaxo Wellcome	ORAL
30TAB Single		Suicide Attempt		Lexomil	C		ORAL
dose	1 DAY	Tracheal Obstruction					

Date:06/14/02ISR Number: 3933157-7Report Type:Expedited (15-DaCompany Report #B0269840A

Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death							
Hospitalization -							
1 DAY				Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged				Ecstasy	SS		ORAL
		Blood Creatinine Increased					
		Blood Ph Decreased					



Cardiac Arrest  
 Diarrhoea  
 Disseminated  
 Intravascular Coagulation  
 Drug Interaction  
 General Physical Health  
 Deterioration  
 Hyperhidrosis  
 Hyperpyrexia  
 Hypotension  
 Metabolic Acidosis  
 Mydriasis  
 Renal Failure  
 Rhabdomyolysis  
 Serotonin Syndrome  
 Shock

Date:06/18/02ISR Number: 3934255-4Report Type:Expedited (15-DaCompany Report #A0371095A  
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Per day	Cystitis	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged 50MG Per day	Drug Interaction		Atenolol	SS		ORAL
360MG Unknown	Hypotension		Cardizem Cd	SS	Glaxo Wellcome	ORAL
	Medication Error Pharmaceutical Product Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/18/02ISR Number: 3934256-6Report Type:Expedited (15-DaCompany Report #A0371180A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Clonic Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day							
				Marijuana	C		
				Naprosyn	C		
				Flexeril	C		

Date:06/18/02ISR Number: 3934265-7Report Type:Expedited (15-DaCompany Report #B0270362A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Discomfort		Zyban	PS	Glaxo Wellcome	ORAL
		Dyspnoea		Amarel	C		
		Headache					
		Hypertension					
		Pulmonary Oedema					

Date:06/18/02ISR Number: 3934267-0Report Type:Expedited (15-DaCompany Report #B0270415A  
Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Zyban	PS	Glaxo Wellcome	
UNKNOWN	150MG	Unknown					
Other		Overdose					
		Psychotic Disorder					

Date:06/18/02ISR Number: 3934269-4Report Type:Expedited (15-DaCompany Report #B0270422A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Carotid Artery Thrombosis		Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown	23 DAY						

Hospitalization - 1.25MG Initial or Prolonged Unknown	Cerebrovascular Accident  Transient Ischaemic Attack	Oestrone	C		ORAL
RESPIRATORY  (INHALATION)	50MCG Twice  per day	Salmeterol	C	Glaxo Wellcome	
RESPIRATORY  (INHALATION)	2PUFF Four  times per day	Salbutamol	C	Glaxo Wellcome	
RESPIRATORY  (INHALATION)	500MCG Four  times per day	Beclomethasone	C	Glaxo Wellcome	
RESPIRATORY  (INHALATION)	40MCG Four  times per day	Ipratropium	C	Glaxo Wellcome	

Date:06/18/02ISR Number: 3934270-0Report Type:Expedited (15-DaCompany Report #B0270659A  
Age:43 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - UNKNOWN	Chest Pain		Zyntabac	PS	Glaxo Wellcome	
Initial or Prolonged UNKNOWN	Hypothermia		Alendronate Sodium	C		
	Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/18/02ISR Number: 3934271-2Report Type:Expedited (15-DaCompany Report #D0038028A

Age:32 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 7.5G per day 1 DAY	Diplopia	Health	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged Other	Fear Hallucination Headache Intentional Misuse Suicide Attempt Tachycardia Tremor Vomiting	Professional				

Date:06/18/02ISR Number: 3935704-8Report Type:Expedited (15-DaCompany Report #A0371291A

Age:36 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Confusional State Convulsion Crying Dysgeusia	Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
SEE DOSAGE TEXT ORAL 2 YEARS	Emotional Disorder Thinking Abnormal					
25 MG ORAL	Vaginal Infection		Lamictal Tablet (Lamotrigine)	SS		ORAL
ORAL			Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	SS		ORAL
			Clonazepam Lorazepam Ortho-Tri-Cyclen	C C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Accident Eye Pain Thrombosis	Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
1							
TABLET/TWICE							
DALY/ORAL							
				Cetirizine Hydrochloride	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Arrest Epilepsy Respiratory Arrest		Zyban Miranova	PS C	Glaxo Wellcome	ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/20/02ISR Number: 3935966-7Report Type:Expedited (15-DaCompany Report #A0170310A  
Age:69 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day Initial or Prolonged Disability		Cardiac Failure Congestive Coronary Artery Disease Dystonia	Health Professional	Wellbutrin Sr	PS	Glaxo Wellcome	ORAL

Date:06/20/02ISR Number: 3935970-9Report Type:Expedited (15-DaCompany Report #A0371430A  
Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Withdrawal Convulsions		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:06/20/02ISR Number: 3935973-4Report Type:Expedited (15-DaCompany Report #A0372010A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 75MG Per day 12 DAY Initial or Prolonged		Drug Interaction International Normalised Ratio Decreased	Health Professional	Wellbutrin Trazodone Warfarin Lovenox Zyprexa Clonazepam Colace Zoloft Tylenol Motrin Diphenhydramine Reglan	PS SS SS C C C C C C C C C C	Glaxo Wellcome Glaxo Wellcome	ORAL

Date:06/20/02ISR Number: 3935978-3Report Type:Expedited (15-DaCompany Report #B0122442A  
Age:34 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day	12 DAY	C-Reactive Protein	Health	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	1UNIT per day	1 DAY	Increased	Professional	Propofan	SS		ORAL
Disability	40MG per day	277 DAY	Cheilitis		Prazepam	SS		ORAL
	.25MG per day	277 DAY	Dermatitis Bullous		Alprazolam	C		ORAL
	1UNIT per day	277 DAY	Erythema Multiforme		Zolpidem	C		ORAL
			Gamma-Glutamyltransferase					
			Increased					
			Hypopyon					
			Intertrigo					
			Leukocytosis					
			Oedema					
			Oral Soft Tissue Disorder					
			Pyrexia					
			Rash Pustular					
			Skin Exfoliation					
			Skin Necrosis					
			White Blood Cell Count					
			Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/20/02ISR Number: 3935979-5Report Type:Expedited (15-DaCompany Report #B0259252A  
 Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day 12 DAY	C-Reactive Protein	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	40MG per day 276 DAY	Increased		Prazepam	SS		ORAL
	1UNIT per day 1 DAY	Cheilitis		Propofan	C		ORAL
	1UNIT per day 276 DAY	Dermatitis Bullous		Zolpidem	C		ORAL
.25MG As		Hypopyon		Alprazolam	C		ORAL
required	276 DAY	Inflammation					
		Intertrigo					
		Leukocytosis					
		Oedema					
		Pyrexia					
		Rash Pustular					
		Skin Exfoliation					
		Skin Necrosis					

Date:06/20/02ISR Number: 3935984-9Report Type:Expedited (15-DaCompany Report #B0270679A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG	Overdose		Zyban	PS	Glaxo Wellcome	
UNKNOWN		Unknown					
		Serotonin Syndrome					

Date:06/20/02ISR Number: 3935986-2Report Type:Expedited (15-DaCompany Report #B0270906A  
 Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG As	Psychotic Disorder		Zyban	PS	Glaxo Wellcome	
UNKNOWN							
directed	52 DAY						



UNKNOWN Paracetamol + Codeine C

RESPIRATORY Fenoterol + Ipratropium C

( INHALATION)

UNKNOWN Cocaine C

Date:06/20/02ISR Number: 3936777-9Report Type:Direct Company Report #CTU 170637  
 Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Wellbutrin	PS		ORAL
MINIMUM DAILY		Hostility					
ORAL							

Date:06/21/02ISR Number: 3936682-8Report Type:Expedited (15-DaCompany Report #A0372195A  
 Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Metabolic Acidosis	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
		Overdose	Professional	Topamax	SS		ORAL
				Amaryl	SS		
				Prozac	SS		
				Vasotec	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/21/02ISR Number: 3936684-1Report Type:Expedited (15-DaCompany Report #B0122944A

Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day	41 DAY	Depression	Zyban	PS	Glaxo Wellcome	ORAL
Drug Withdrawal Syndrome							

Date:06/21/02ISR Number: 3936688-9Report Type:Expedited (15-DaCompany Report #B0269955A

Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day	3 WK	Alpha 2 Globulin	Zyban	PS	Glaxo Wellcome	ORAL
Increased Arthralgia C-Reactive Protein Increased Myalgia Red Blood Cell Sedimentation Rate Increased							

Date:06/21/02ISR Number: 3936696-8Report Type:Expedited (15-DaCompany Report #B0271139A

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Enterocolitis Haemorrhagic	Zyban	PS	Glaxo Wellcome	ORAL

Date:06/21/02ISR Number: 3937138-9Report Type:Direct Company Report #CTU 170729

Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required	75MG BID ORAL		Aggression	Bupropion	PS		ORAL
Intervention to			Agitation	Diltiazem Sa	C		

Prevent Permanent Anxiety  
Impairment/Damage Homicidal Ideation

Hctz  
25mg/Triamterene C  
Potassium Chloride C  
Oxycodone Hcl C  
Baclofen C  
Pseudoephedrine C  
Promethazine C

Date:06/21/02ISR Number: 3937144-4Report Type:Direct  
Age:41 YR Gender:Male I/FU:I

Company Report #CTU 170717

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tremor		Zyban 150mg	PS		ORAL
1T TAB PO QD							
X 3 D THEN PO							
BID , NOW QD							
				Habitrol Patch	C		

Date:06/24/02ISR Number: 3937901-4Report Type:Direct  
Age:25 YR Gender:Female I/FU:I

Company Report #CTU 170796

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion		Bupropriion-Wellbutri n- 100mg Tabs			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

100MG 2X			Glaxosmithkline	PS	Glaxosmithkline	ORAL
DAILY ORAL			Prozac	C		
			Atarax	C		

Date:06/24/02ISR Number: 3939099-5Report Type:Expedited (15-DaCompany Report #2002-BP-02683AU (0)  
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional	Atrovent (Ipratropium Bromide) (Nr)			
Other			Other	(Ipratropium-Br)	PS		
				Zyban Sr (Nr)	SS		
				Temazepam (Temazepam) (Nr)	SS		
				Antenex (Diazepam) (Nr)	SS		
				Dichlotride (Hydrochlorothiazide) (Nr)	SS		
				Epilim (Valproate Sodium) (Nr)	SS		
				Ventolin (Salbutamol) (Nr)	SS		
				Flixotide (Fluticasone Propionate) (Nr)	C		

Date:06/27/02ISR Number: 3940684-5Report Type:Direct Company Report #CTU 171129  
 Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG SR PO		Dyspnoea		Wellbutrin Sr 150mg	PS		ORAL
Initial or Prolonged BID	11 DAY	Face Oedema					
		Medication Error		Prenatal Vitamin	C		
		Pruritus		Acetaminophen	C		
				Ibuprofen	C		

Date:06/27/02ISR Number: 3940929-1Report Type:Expedited (15-DaCompany Report #A0372287A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Foetal Growth Retardation Maternal Drugs Affecting Foetus	Study Health Professional	Bupropion Unspecified Tablet (Bupropion Hydrochloride)			
ORAL					PS		ORAL

Date:06/27/02ISR Number: 3940963-1Report Type:Expedited (15-DaCompany Report #A0372286A

Age: Gender: I/FU:I

Outcome	PT
Other	Angiopathy Complications Of Maternal Exposure To Therapeutic Drugs Drug Exposure During

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pregnancy  
Placental Disorder

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
TRANSPLACENTAL	ORAL	Study Health Professional	Bupropion Unspecified (Bupropion Hydrochloride)	PS		

Date:06/27/02ISR Number: 3941582-3Report Type:Expedited (15-DaCompany Report #B0271009A  
Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Anxiety Nausea Panic Attack	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG PER DAY	ORAL	Tremor					

Date:06/28/02ISR Number: 3942017-7Report Type:Expedited (15-DaCompany Report #B0261044A  
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Asthenia Blindness Transient Conjunctival Hyperaemia	Foreign Health Professional	Amoxil (Formulation Unknown) (Amoxicillin)	PS		ORAL
ORAL		Depressed Level Of Consciousness Diarrhoea		Zyban Tablet - Zyban (Bupropion Hydrochloride)	SS		ORAL
ORAL		Dysstasia Hypertension Syncope Urticaria		Acetaminophen + Codeine	C		

Date:07/01/02ISR Number: 3942081-5Report Type:Direct  
Age:44 YR Gender:Male I/FU:I

Company Report #CTU 171325

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Zyban 150mg Glaxco	PS	Glaxco	ORAL
1 PILL 2		Feeling Abnormal					
TIMES ORAL		Hypoaesthesia					
		Hypoaesthesia Oral					
		Nervousness					
		Panic Attack					
		Paraesthesia					

Date:07/01/02ISR Number: 3942537-5Report Type:Expedited (15-DaCompany Report #A0364730A  
Age:43 YR Gender:Female I/FU:F

Outcome	PT
Other	Convulsion
	Crying
	Delusion
	Hallucination, Auditory
	Hallucination, Visual
	Headache
	Memory Impairment
	Mood Swings
	Panic Attack

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Paranoia Schizoaffective Disorder	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE PER DAY/ORAL			Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
				Levonorgestrel	C		
				Thyroxine Sodium	C		
				Alprazolam	C		

Date:07/01/02ISR Number: 3942987-7Report Type:Expedited (15-DaCompany Report #B0271436A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other SEE DOSAGE TEXT / ORAL		Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
		Drugs	Company				
		Maternal Drugs Affecting Foetus	Representative				

Date:07/01/02ISR Number: 3942996-8Report Type:Expedited (15-DaCompany Report #D0038746A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Acidosis Circulatory Collapse Convulsion	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
		Overdose		Metformin Hydrochloride	C		



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage	150 MG/TWICE PER DAY	Tongue Ulceration	Foreign Health Professional	Nicoderm Cq Patch (Nicotine)	PS		
				Zyban Tablet - Zyban (Bupropion Hydrochloride)	SS		
				Ranitidine Hydrochloride	C		
				Labetalol Hydrochloride	C		
				Napratec	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG/ TWICE PER DAY/ ORAL	Bundle Branch Block Left Chest Discomfort Myalgia	Foreign Health Professional	Zyban Tablet- Zyban (Bupropion Hydrochloride)	PS		ORAL
		Myocardial Ischaemia	Company Representative	Birth Control	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/02/02ISR Number: 3942578-8Report Type:Direct  
Age:74 YR Gender:Male I/FU:I

Company Report #CTU 171445

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG BID		Atrioventricular Block		Bupropion 150mg	PS		ORAL
Initial or Prolonged ORAL		Bundle Branch Block Right					
		Dizziness		Verapamil	C		
		Fall		Lisinopril	C		
				Glipizide	C		
				Hctz	C		

Date:07/02/02ISR Number: 3943700-XReport Type:Expedited (15-DaCompany Report #A0372639A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous	Study	Wellbutrin			
		Blighted Ovum	Health	Unspecified Tablet			
		Complications Of Maternal	Professional	(Bupropion			
300 MG PER		Exposure To Therapeutic		Hydrochloride)	PS		ORAL
DAY ORAL		Drugs					
		Maternal Drugs Affecting					
		Foetus					
		Pregnancy					

Date:07/02/02ISR Number: 3944131-9Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #260829

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Euphoric Mood	Consumer	Klonopin Tablets			
1 PER PRN		Fall	Health	(Clonazepam) 0.5 Mg	PS		ORAL
ORAL		Fatigue	Professional				
		Grand Mal Convulsion	Other	Wellbutrin			
		Libido Decreased		(Ibuproprion			

150 MG 2 PER	Memory Impairment	Hydrochloride)	SS	ORAL
DAY ORAL	7 DAY	Tooth Injury		
20 MG DAILY		Paxil (Paroxetine)	SS	ORAL
ORAL				

Date:07/03/02ISR Number: 3944208-8Report Type:Expedited (15-DaCompany Report #B0125399A  
Age:42 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Disability 150 MG ORAL	Duration Cerebrovascular Accident Confusional State Coordination Abnormal	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
	Dizziness Drug Toxicity Dysarthria Fall Feeling Abnormal Hallucination, Visual Metabolic Disorder Motion Sickness Swelling Vision Blurred		Ferroul Sulfate Promethazine Propiomazine	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/03/02ISR Number: 3944209-XReport Type:Expedited (15-DaCompany Report #B0130344A

Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150 MG/ PER DAY/ORAL	Alcohol Poisoning Atherosclerosis Cardiac Arrest	Foreign Health Professional	Zyban Tablet -Zyban (Bupropion Hydrochloride)	PS		ORAL
		Drug Toxicity					
	50 MG SEE	Overdose Pulmonary Congestion Pulmonary Oedema		Dothiepin (Formulation Unknown) (Dothiepin)	SS		ORAL
	DOSAGE TEXT / ORAL	Toxicologic Test Abnormal					
		Ventricular Hypertrophy					
				Ethanol Liquid (Alcohol)	SS		ORAL
				Prempak-C	C		
				Salbutamol Sulphate	C		
				Diazepam	C		
				Fluoxetine	C		
				Zopiclone	C		
				Beclomethasone			
				Dipropion	C		

Date:07/03/02ISR Number: 3944302-1Report Type:Expedited (15-DaCompany Report #B0270659A

Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG / ORAL	Chest Pain Hypothermia Urticaria	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
				Alendronate Sodium	C		

Date:07/03/02ISR Number: 3944304-5Report Type:Expedited (15-DaCompany Report #B0270415A

Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Required Intervention to 150 MG/ AS Prevent Permanent DIRECTED Impairment/Damage		Completed Suicide Overdose Psychotic Disorder	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		
				Oxazepam	C		
				Mirtazapine	C		

Date:07/03/02ISR Number: 3944909-1Report Type:Expedited (15-DaCompany Report #US008746  
Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 200 MG QD Initial or Prolonged ORAL		Atelectasis Bipolar Disorder	Health Professional	Provigil	PS		ORAL
6 TAB ONCE ORAL		Coma Delusion Depression		Provigil Wellbutrin	SS SS		ORAL
200 MG QD ORAL		Electrocardiogram T Wave Abnormal		Wellbutrin	SS		ORAL
		Hypomania Mania Overdose Psychotic Disorder Suicidal Ideation		Zyprexa Lithium	SS C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/05/02ISR Number: 3944959-5Report Type:Expedited (15-DaCompany Report #B0271767A  
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Drug Interaction Hypomania	Literature Health Professional	Paxil (Paroxetine Hydrochloride) Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS  SS		
1	YR			Carbamazepine	C		

Date:07/05/02ISR Number: 3945789-0Report Type:Expedited (15-DaCompany Report #B0272251A  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Asthenia Fall Hyperthyroidism	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG /							
VARIABLE DOSE							
/ ORAL				Propranolol Hydrochloride Ethanol Silymarin	C C C		

Date:07/05/02ISR Number: 3945968-2Report Type:Expedited (15-DaCompany Report #A0372635A  
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Dizziness Feeling Abnormal Heart Rate Decreased	Foreign Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER							

DAY / ORAL

Date:07/05/02ISR Number: 3945981-5Report Type:Expedited (15-DaCompany Report #B0271148A

Age:33 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Pruritus	Foreign	Zyban Tablet - Zyban			
Initial or Prolonged	Tenosynovitis		(Bupropion Hydrochloride)	PS		ORAL
150 MG/ PER						

DAY / ORAL

Date:07/05/02ISR Number: 3945992-XReport Type:Expedited (15-DaCompany Report #B0114445A

Age:55 YR Gender:Female I/FU:F

Outcome	PT
Other	Anxiety
	Arthralgia
	Depression
	Difficulty In Walking
	Fatigue
	Headache
	Hypersomnia
	Insomnia
	Malaise

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Movement Disorder				
		Myalgia				
Dose	Duration	Nausea	Report Source	Product	Role	Manufacturer
		Pain In Extremity	Foreign	Zyban Tablet - Zyban		
		Polymyalgia Rheumatica	Health	(Bupropion		
150 MG/ TWICE		Red Blood Cell	Professional	Hydrochloride)	PS	
		Sedimentation Rate				
PER DAY/		Increased				
		Somnolence				
		Tremor				

Date:07/05/02ISR Number: 3945999-2Report Type:Expedited (15-DaCompany Report #B0269491A  
 Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Facial Palsy	Foreign	Zyban Tablet - Zyban			
Initial or Prolonged			Health	(Bupropion			
			Professional	Hydrochloride)	PS		ORAL
150 MG/							
VARIABLE							
DOSE/ ORAL							

Atorvastatin Calcium C  
 Amlodipine C  
 Enalapril C

Date:07/05/02ISR Number: 3946022-6Report Type:Expedited (15-DaCompany Report #B0271161A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Mean Cell Volume Abnormal	Foreign	Zyban Tablet - Zyban			
		Neuropathy Peripheral		(Bupropion			
150 MG/PER				Hydrochloride)	PS		ORAL
DAY/ORAL				Miniphase Tablet			



1	TABLET/PER	(Miniphase)	SS	ORAL
	DAY/ORAL			
.5		Bromazepam Tablet (Bromazepam)	SS	ORAL
	TABLET/TWICE			
	PER DAY/ORAL			
2		Di-Antalvic (Di-Antalvic)	SS	ORAL
	UNIT/MONTHLY/			
	ORAL			

Date:07/05/02ISR Number: 3946044-5Report Type:Expedited (15-DaCompany Report #B0271860A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation Anxiety Depressed Mood	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
SEE DOSAGE		Depression					
TEXT/ORAL		Dry Mouth Insomnia Malaise Medication Error Panic Attack Suicidal Ideation		Temazepam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/05/02ISR Number: 3946237-7Report Type:Expedited (15-DaCompany Report #A0367126A  
 Age:20 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Required Intervention to 1500 MG ORAL Prevent Permanent Impairment/Damage	Convulsion Intentional Misuse Intentional Self-Injury Vomiting	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)  Methylphenidate Hci	PS  C		ORAL

Date:07/05/02ISR Number: 3946287-0Report Type:Expedited (15-DaCompany Report #B0271672A  
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other  ORAL	Pneumothorax	Foreign Health Professional  Company Representative	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:07/05/02ISR Number: 3946325-5Report Type:Expedited (15-DaCompany Report #B0272116A  
 Age:46 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged  150 MG TWICE  PER DAY ORAL	Disinhibition Elevated Mood Hallucination, Auditory  Mania  Psychomotor Hyperactivity	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:07/05/02ISR Number: 3946593-XReport Type:Expedited (15-DaCompany Report #A0373043A  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Drug Interaction Potentiation	Health Professional Company Representative	Wellburtin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL				Cetirizine Hydrochloride (Formulation Unknown) (Cetirizine Hydrochloride)	SS		ORAL
ORAL				Antibiotics (Formulation Unknown) (Antibiotics)	SS		ORAL
ORAL				Montelukast Sodium (Formulation Unknown) (Montelukast Sodium)	SS		ORAL
ORAL				Allergy Medication (Formulation Unknown) (Allergy Medication)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/05/02ISR Number: 3946610-7Report Type:Expedited (15-DaCompany Report #B0270679A  
 Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Overdose	Foreign	Zyban Tablet - Zyban			
		Serotonin Syndrome	Health	(Bupropion			
150 MG			Professional	Hydrochloride)	PS		
				Quetiapine	C		
				Clonazepam	C		

Date:07/08/02ISR Number: 3944409-9Report Type:Expedited (15-DaCompany Report #B0269955A  
 Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alpha 2 Globulin		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	3 WK	Increased					
		Arthralgia					
		C-Reactive Protein					
		Increased					
		Myalgia					
		Red Blood Cell					
		Sedimentation Rate					
		Increased					

Date:07/08/02ISR Number: 3944418-XReport Type:Expedited (15-DaCompany Report #D0038493A  
 Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Apathy		Zyban	PS	Glaxo Wellcome	ORAL
60TAB Single							
Initial or Prolonged		Disorientation					
dose	1 DAY						
		Disturbance In Attention		Diclofenac	SS		ORAL
40TAB Single							
		Hypertension					
dose	1 DAY						
		Intentional Misuse		Diphenhydramine			

10TAB Single      Suicide Attempt      Hydrochloride      SS      Glaxo Wellcome      ORAL  
 Tachycardia  
 dose      1      DAY  
 UNKNOWN      Amphetamine      SS

Date:07/09/02ISR Number: 3945042-5Report Type:Expedited (15-DaCompany Report #A0360169A  
 Age:52 YR      Gender:Female      I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice		Depressed Mood					
per day	8	MON		Paxil	C	Glaxo Wellcome	ORAL
20MG Per day		Fatigue					
750MG At		Intentional Misuse		Epival	C		
night				Lipitor	C		ORAL
40MG Per day				Remeron	C		
45MG At night							

Date:07/09/02ISR Number: 3945151-0Report Type:Direct      Company Report #CTU 171795  
 Age:34 YR      Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Alopecia		Sulfasalazine 500mg	PS	Watson Laboratories	ORAL
500MG TWICE				Watson Laboratories			
DAILY ORAL				Wellbutrin 150mg	SS		ORAL
150MG ONCE							
DAILY ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/09/02ISR Number: 3945742-7Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #CTU 171802

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Hypersensitivity		Wellbutrin Sr Tab 100 Mg Glaxo Smithkline	PS	Glaxo Smithkline	ORAL
100MG SR QAM							
ORAL				Dilantin	C		
				Norvasc	C		
				Cipro	C		
				Nicoderm Patch	C		
				Ativan	C		
				Ambien	C		
				Tylenol	C		

Date:07/09/02ISR Number: 3945771-3Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #CTU 171815

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Serum Sickness		Wellbutrin Sr 150mg	PS		
BID							
Intervention to				Lescol Xl	C		
Prevent Permanent				Adallat Cc	C		
Impairment/Damage							

Date:07/09/02ISR Number: 3946846-5Report Type:Expedited (15-DaCompany Report #A0373063A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Convulsion Drug Abuser Fall	Health Professional Company	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
100 MG /							
THREE TIMES		Laceration	Representative				
PER DAY /		Road Traffic Accident					

ORAL

Wellbutrin Sr  
Tablet-Controlled  
Release SS

ORAL

ORAL

Methadone  
Hydrochloride C

Date:07/09/02ISR Number: 3946987-2Report Type:Expedited (15-DaCompany Report #USA-2002-0001076

Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Aortic Atherosclerosis	Consumer	Oxycontin Tablets			
Other		Coronary Artery Atherosclerosis	Health Professional	(Oxycodone Hydrochloride) Cr			
		Drug Abuser	Other	Tablet	PS		
		Drug Dependence		Methadone			
		Drug Toxicity		(Methadone) Unknown	SS		
		Fibrosis		Hydrocodone			
		Mitral Valve Disease		Bitartrate (Similar			
		Pericardial Effusion		To Ind 59,175)			
		Pleural Effusion		(Hydrocodone			
		Pneumonia Bacterial		Bitartrate) Unknown	SS		
		Respiratory Tract		Dihydrocodeine /			
		Infection		Caffeine /			
				Acetaminophen			

MG

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Freedom Of Information (FOI) Report

MG	(Similar To Anda 88-584)	SS
MG	Bupropion (Amfebutamone) Unknown	SS
MG	Ibuprofen (Ibuprofen) Tablet	SS
MG	Dextromethorphan (Dextromethorphan)	SS
MG	Ephedrine (Ephedrine) Unknown	SS
MG	Pseudoephedrine (Pseudoephedrine) Unknown	SS
MG	Doxylamine (Doxylamine) Unknown	SS
MG	Levorphanol (Levorphanol) Unknown	SS
MG	Alprazolam (Alprazolam) Unknown	SS
MG	Codeine (Codeine) Unknown	SS
MG	Heroin (Diamorphine) Unknown	SS

Date:07/09/02ISR Number: 3947122-7Report Type:Expedited (15-DaCompany Report #A0371095A

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/PER	Blood Pressure Decreased Cystitis	Consumer	Zyban (Bupropion Hydrochloride)	PS		ORAL



Drug Interaction

DAY/ORAL

Myocardial Infarction

Atenolol  
(Formulation  
Unknown) (Generic)  
(Atenolol) SS ORAL

50 MG/PER

DAY/ORAL

Diltiazem  
Hydrochloride SS ORAL

360 MG/ORAL

Date:07/09/02ISR Number: 3947135-5Report Type:Expedited (15-DaCompany Report #A0362556A  
Age:16 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Conversion Disorder	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
200 MG/TWICE							
PER DAY/ORAL				Amitriptyline Hcl	C		

Date:07/09/02ISR Number: 3947185-9Report Type:Expedited (15-DaCompany Report #A0359984A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Ecchymosis Overdose Pulmonary Congestion	Foreign Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

150 MG PER				Hydrochloride)	PS		ORAL
DAY/ORAL				Citalopram			
				Hydrobromide	C		
				Clonazepam	C		

Date:07/09/02ISR Number: 3947262-2Report Type:Expedited (15-DaCompany Report #B0272342A  
 Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthralgia Insomnia Joint Effusion	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrchoride)	PS		ORAL
ORAL		Nephrolithiasis Renal Colic Tremor					

Date:07/09/02ISR Number: 3947273-7Report Type:Expedited (15-DaCompany Report #B0272177A  
 Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Sudden Death	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL				Zestoretic	C		

Date:07/10/02ISR Number: 3945938-4Report Type:Expedited (15-DaCompany Report #A0370262A  
 Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	61 DAY	Abnormal Dreams Anorexia Back Injury		Zyban	PS	Glaxo Wellcome	ORAL

Crying  
Epistaxis  
Insomnia  
Mood Altered  
Nasal Odour  
Rhinorrhoea  
Stress

Date:07/10/02ISR Number: 3945944-XReport Type:Expedited (15-DaCompany Report #B0260007A  
Age:49 YR Gender:Male I/FU:F

Outcome  
Death

PT  
Aphasia  
Brain Oedema  
Bronchospasm  
C-Reactive Protein  
Increased  
Cardio-Respiratory Arrest  
Chest X-Ray Abnormal  
Coma  
Computerised Tomogram  
Abnormal  
Cyanosis  
Hepatic Failure  
Injury

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Laryngeal Oedema Leukocytosis Malaise					
75MG Per day		Mydriasis Painful Respiration		Zyban Kardegic	PS SS	Glaxo Wellcome	ORAL ORAL
1 DAY		Pharyngolaryngeal Pain		Rulid	SS	Glaxo Wellcome	ORAL
400MG per day	1 DAY	Pyrexia		Surgam	C		ORAL
TRANSDERMAL		Renal Failure Syncope		Elisor Discotrine	C C	Glaxo Wellcome	ORAL
		Ventricular Fibrillation		Sectral Artotec	C C		ORAL ORAL

Date:07/11/02ISR Number: 3948193-4Report Type:Expedited (15-DaCompany Report #0714201A  
Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tongue Ulceration	Foreign Health Professional	Habitrol-Nicotine Transdermal 21mg-Hvch	PS	Nvch	
1			Other				
PATCH/QD/TTS				Zyban (Bupropion)	SS		ORAL
150 MG/BD/PO				Ranitidine Labetalol Naparatec (Naproxen, Misoprostol)	C C C		

Date:07/12/02ISR Number: 3948285-XReport Type:Expedited (15-DaCompany Report #2002114146GB  
Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tongue Ulceration	Foreign Health	Nicoderm (Nicotine) Patch	PS		
TRANSDERMAL	SEE IMAGE						

150 MG, BID,

ORAL

Professional  
Other

Zyban (Amfebutamone  
Hydrochloride) SS

ORAL

Ranitidine  
Hydrochloride C  
Labetalol  
Hydrochloride  
(Labetalol  
Hydrochloride) C

Date:07/12/02ISR Number: 3948713-XReport Type:Expedited (15-DaCompany Report #A103444

Age:79 YR Gender:Female I/FU:F

Outcome	PT
Required	Anger
Intervention to	Balance Disorder
Prevent Permanent	Blood Oestrogen Decreased
Impairment/Damage	Blood Pressure Increased
	Blood Thyroid Stimulating
	Hormone Decreased
	Cerebral Haemorrhage
	Contusion
	Convulsion
	Drug Ineffective
	Epistaxis
	Fall

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
50.00 MG	TOTAL : DAILY	Head Injury Headache	Consumer	Zoloft Tablets	PS		ORAL
	: ORAL	Heart Rate Decreased					
300.00 MG	TOTAL: BID:	Heart Rate Increased					
	ORAL	Hostility					
		Hyperhidrosis					
		Laceration		Wellbutrin Sr	SS		ORAL
		Medication Error					
		Meningioma					
300.00 MG	TOTAL : BID :	Nuclear Magnetic Resonance Imaging Brain		Coumadin Effexor Xr	SS SS		ORAL
	ORAL	Abnormal					
		Overdose					
		Palpitations		Toprol	C		
		Parkinsonism		Zebeta	C		
		Prothrombin Time Prolonged		Lanoxin Lasix	C C		
		Syncope		Synthroid	C		
		Transient Ischaemic Attack		Vitamin E Estrogen	C C		
		Tremor		Zocor	C		
		Vaginal Haemorrhage		Estradiol With Vaginal Ring	C C		
				Inderal	C		
				Monopril	C		
				Cardizem	C		
				Folic Acid	C		
				Amantadine (Subject)	C		

Date:07/15/02ISR Number: 3948156-9Report Type:Expedited (15-DaCompany Report #A0373618A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dyspnoea	Health	Zantac	PS	Glaxo Wellcome	

Urticaria

Professional

Wellbutrin

SS

Glaxo Wellcome

Paxil

SS

Glaxo Wellcome

Celebrex

SS

Neurontin

SS

Topamax

SS

Klonopin

SS

Progesterone

C

Salbutamol

C

Glaxo Wellcome

Pentosane

C

Polysulfate

C

Hydroxyzine

C

Amitriptyline

C

Oxycodone

C

Hydrochloride

C

Date:07/15/02ISR Number: 3948158-2Report Type:Expedited (15-DaCompany Report #A0373917A

Age:33 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice	Toxic Epidermal	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	Necrolysis					
2	MON					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/15/02ISR Number: 3948168-5Report Type:Expedited (15-DaCompany Report #B0271008A

Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Health	Zyban	PS	Glaxo Wellcome	ORAL
10 DAY		Erythema	Professional	Nicorette	C	Glaxo Wellcome	ORAL
		Joint Swelling					
		Malaise					
		Myalgia					
		Oedema Peripheral					
		Pyrexia					
		Serum Sickness					
		Urticaria					

Date:07/15/02ISR Number: 3948439-2Report Type:Direct

Company Report #CTU 172287

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Confusional State		Bupropion 150 Mg			
Initial or Prolonged		Hallucination		Glaxo-Wellcome	PS	Glaxo-Wellcome	ORAL
150 MG. BID							
ORAL		Hypoglycaemia					
				Amaryl	C		
				Actos	C		
				Glucophage	C		

Date:07/15/02ISR Number: 3948605-6Report Type:Direct

Company Report #CTU 172300

Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Chills		Zyban 150mg			
Hospitalization -		Confusional State		Glaxo-Wellcome	PS	Glaxo-Wellcome	
ONE IN AM							
Initial or Prolonged		Dizziness		Premarin 1.25mg	C		
Disability		Dysarthria		Aspirin	C		
		Euphoric Mood		Centr	C		
		Eye Rolling		Silver Vitiman	C		
		Fear					



Hyperhidrosis  
Movement Disorder  
Oedema  
Sensation Of Heaviness

Date:07/16/02ISR Number: 3948684-6Report Type:Expedited (15-DaCompany Report #A0369795A  
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	21 DAY	Conjunctival Hyperaemia Eye Irritation Vision Blurred Visual Disturbance	Health Professional	Zyban	PS	Glaxo Wellcome	ORAL

Date:07/16/02ISR Number: 3948687-1Report Type:Expedited (15-DaCompany Report #A0373866A  
Age:59 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Dyspepsia Fall Lower Limb Fracture Tinnitus

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Upper Limb Fracture

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice			Wellbutrin	PS	Glaxo Wellcome	ORAL
per day						
240MG per day			Diltiazem	C	Glaxo Wellcome	
5MG per day			Prinivil	C		
1.5MG per day			Clonazepam	C		
250MG Twice			Ticlid	C		
per day						
600MG per day			Neurontin	C		
40UNIT Twice			Insulin	C		
per day						
			Vitamin B12	C	Glaxo Wellcome	
			Tylenol	C	Glaxo Wellcome	

Date:07/16/02ISR Number: 3948707-4Report Type:Expedited (15-DaCompany Report #B0271860A  
 Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation	Health	Zyban	PS	Glaxo Wellcome	ORAL
18 DAY		Anxiety	Professional	Temazepam	C		
UNKNOWN		Depressed Mood					
		Depression					
		Drug Ineffective					
		Dry Mouth					
		Insomnia					
		Malaise					
		Medication Error					
		Panic Attack					
		Suicidal Ideation					

Date:07/16/02ISR Number: 3949386-2Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 172424

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MGM BID	Grand Mal Convulsion		Wellbutrin Sr	PS		
Required				Ritalin	C		
Intervention to				Ambien	C		
Prevent Permanent							
Impairment/Damage							

Date:07/16/02ISR Number: 3949721-5Report Type:Expedited (15-DaCompany Report #B0272924A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abnormal Behaviour	Foreign	Zyban Tablet - Zyban	PS		ORAL
ORAL							
Initial or Prolonged		Pyromania	Health Professional Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/16/02ISR Number: 3950017-6Report Type:Expedited (15-DaCompany Report #B0270415A  
Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Required		Completed Suicide	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		
Intervention to Prevent Permanent Impairment/Damage		Psychotic Disorder					
		Skin Injury		Oxazepam	C		
		Tendon Injury		Mirtazapine	C		

Date:07/16/02ISR Number: 3950020-6Report Type:Expedited (15-DaCompany Report #D0038838A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Aggression	Foreign Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
		Anxiety					
		Bipolar Disorder					
150 MG/ ORAL		Depression		Paxil (Formulation Unknown) (Paroxetine Hydrochloride)	SS		
		Diarrhoea					
		Disturbance In Attention					
		Hyperhidrosis					
		Hypomania					
		Migraine					
		Nausea					
		Nervousness					
		Sleep Disorder					
		Tachycardia					
		Vertigo					
		Weight Decreased					

Date:07/16/02ISR Number: 3950278-3Report Type:Expedited (15-DaCompany Report #A0072356A  
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required		Akathisia	Health Professional	Wellbutrin Tablet (Bupropion			
		Clonic Convulsion					

Intervention to 100 MG/THREE Prevent Permanent TIMES PER Impairment/Damage DAY/ORAL	Depersonalisation  Dystonia  Nervous System Disorder  Respiratory Alkalosis Tardive Dyskinesia Tremor	Company  Representative	Hydrochloride)    Triamcinolone Acetonide Melatonin	PS    C C	ORAL
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Date:07/16/02ISR Number: 3950292-8Report Type:Expedited (15-DaCompany Report #A0366635A  
Age:38 YR Gender:Male I/FU:F

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required	Atrial Flutter Blindness Transient Dyspnoea	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
Intervention to SEE DOSAGE Prevent Permanent TEXT/ORAL Impairment/Damage	Heart Rate Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/17/02ISR Number: 3949441-7Report Type:Expedited (15-DaCompany Report #A0373933A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	Death	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
			Professional	Tegretol	C		

Date:07/17/02ISR Number: 3949442-9Report Type:Expedited (15-DaCompany Report #A0374100A

Age:24 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day 4 DAY	Abortion Spontaneous	Consumer	Wellbutrin	PS	Glaxo Wellcome	ORAL
		Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus					

Date:07/17/02ISR Number: 3949555-1Report Type:Expedited (15-DaCompany Report #316974

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dyspnoea		Dyspnoea	Health	Klonopin	PS	Roche	
Urticaria		Urticaria	Professional	Celebrex	SS		
				Neurontin	SS		
				Topamax	SS		
				Paxil	SS		
				Zantac	SS		
				Wellbutrin	SS		
				Progesterone	C		
				Salbutamol	C		
				Pentosane			
				Polysulphate	C		
				Hydroxyzine	C		
				Amitriptyline	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Eczema	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER DAY / ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Feeling Hot Haemoglobin Decreased Headache	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ PER DAY/ ORAL							
		Malaise					
		Oral Intake Reduced		Nitrazepam Glibenclamide Fluoxetine Hydrochloride	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/17/02ISR Number: 3950886-XReport Type:Expedited (15-DaCompany Report #A0373158A

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Duration Muscle Spasms	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
200 MG/ TWICE PER DAY/ORAL			Citalopram Hydrobrome	C		

Date:07/18/02ISR Number: 3950201-1Report Type:Expedited (15-DaCompany Report #A0374178B

Age: Gender:Unknown I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Unknown	Duration Developmental Delay		Wellbutrin	PS	Glaxo Wellcome	
	Maternal Drugs Affecting Foetus Premature Baby					

Date:07/19/02ISR Number: 3950827-5Report Type:Expedited (15-DaCompany Report #A0373936A

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 450MG Unknown	Duration Convulsion		Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
			Unknown	C		

Date:07/19/02ISR Number: 3950830-5Report Type:Expedited (15-DaCompany Report #B0120942A

Age:20 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization -	Duration Arthritis		Bupropion			



Initial or Prolonged      Arthritis/Reactive      Hydrochloride      PS      Glaxo Wellcome      ORAL  
150MG Twice  
per day      26      DAY      Asthma  
Dyspnoea      Decortin      C  
Heart Rate Increased  
Hypersensitivity  
Osteoarthritis  
Pyrexia  
Rash  
Tobacco Abuse  
Urticaria  
Yersinia Infection

Date:07/19/02ISR Number: 3950834-2Report Type:Expedited (15-DaCompany Report #B0273496A  
Age:23 YR      Gender:Female      I/FU:I

Outcome      PT  
Hospitalization -      Arthralgia  
Initial or Prolonged      C-Reactive Protein  
Increased  
Coagulation Factor V  
Level Decreased  
Coagulation Factor Vii  
Level Decreased  
Fibrin D Dimer Increased  
Proteinuria

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Prothrombin Level Decreased					
		Urticaria Generalised	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Zyban	PS	Glaxo Wellcome	ORAL
21	DAY			Adepal	C		
UNKNOWN							

Date:07/19/02ISR Number: 3950837-8Report Type:Expedited (15-DaCompany Report #B0273786A  
Age: Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Depression		Zyban	PS	Glaxo Wellcome	
Dose							
Other	150MG	Mania					
UNKNOWN							
Variable dose 11 DAY							

Date:07/19/02ISR Number: 3950842-1Report Type:Expedited (15-DaCompany Report #D0038746A  
Age:55 YR Gender:Male I/FU:F

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Acidosis		Zyban	PS	Glaxo Wellcome	ORAL
Dose							
Hospitalization -		Blood Lactic Acid					
1800MG per		Increased		Metformin	C		ORAL
Initial or Prolonged	2 DAY	Circulatory Collapse					
day		Convulsion					
Other		Overdose					
850MG Per day		White Blood Cell Count Increased					

Date:07/22/02ISR Number: 3951379-6Report Type:Direct Company Report #CTU 172642  
Age:53 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose							

Other 150 MG PO BID	Condition Aggravated	Wellbutrin Sr	PS	ORAL
	Cough	Aspirin	C	
	Extrasystoles	Vitamins	C	
	Sinus Bradycardia			
	Supraventricular			
	Extrasystoles			
	Ventricular Arrhythmia			
	Ventricular Extrasystoles			

Date:07/22/02ISR Number: 3951863-5Report Type:Direct Company Report #CTU 172700  
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Bupropion	PS		ORAL
100MG PO		Crying					
		Decreased Appetite					
		Dyspnoea					
		Fatigue					

Date:07/23/02ISR Number: 3951888-XReport Type:Expedited (15-DaCompany Report #A0374687A  
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ventricular Tachycardia		Bupropion	PS	Glaxo Wellcome	ORAL
200MG Per day	3 DAY						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/23/02ISR Number: 3951892-1Report Type:Expedited (15-DaCompany Report #B0131464A

Age:25 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Unknown 29 DAY	Abdominal Pain		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged 2 DAY	Angina Pectoris Blood Creatine Increased Blood Creatine Phosphokinase Increased Cardiac Disorder Cardiac Failure Chest Pain Cutaneous Vasculitis Cytomegalovirus Infection Diarrhoea Ecchymosis Ejection Fraction Decreased Electrocardiogram St Segment Elevation Gangrene Hepatomegaly Hyperpyrexia Hypotension Leukocytoclastic Vasculitis Malaise Myocardial Infarction Skin Necrosis Somnolence Splenomegaly Troponin I Increased Vomiting		Methylenedioxymetham phetamine Alcohol Cannabis	SS C C		ORAL

Date:07/23/02ISR Number: 3951896-9Report Type:Expedited (15-DaCompany Report #B0267249A

Age:50 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 3 WK	Diabetes Mellitus	Health	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Insulin-Dependent	Professional				

Weight Decreased

Date:07/24/02ISR Number: 3952669-3Report Type:Expedited (15-DaCompany Report #A0372286A  
Age:15 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Maternal Drugs Affecting Foetus Placental Disorder Placental Insufficiency Stillbirth		Bupropion	PS	Glaxo Wellcome	ORAL

Date:07/24/02ISR Number: 3952690-5Report Type:Expedited (15-DaCompany Report #B0272783A  
Age:32 YR Gender:Female I/FU:F

Outcome	PT
Other	Alanine Aminotransferase Increased Aspartate

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
16 DAY		Aminotransferase Increased Blood Creatine Phosphokinase Increased	Zyban	PS	Glaxo Wellcome	ORAL
TRANSDERMAL		Blood Lactate	Nicotine Patch	SS	Glaxo Wellcome	
YR		Dehydrogenase Increased	Ikaran	C		
YR		Insomnia	Adepal	C		
		Muscle Spasms				

Date:07/24/02ISR Number: 3952692-9Report Type:Expedited (15-DaCompany Report #B0273874A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	31 DAY	Aspartate		Zyban	PS	Glaxo Wellcome	ORAL
Life-Threatening	26 DAY	Aminotransferase		Lectil	SS		ORAL
Hospitalization -	10 DAY	Increased		Tanganil	SS		ORAL
Initial or Prolonged	30 DAY	Blood Creatine		Stilnox	SS		ORAL
Other		Phosphokinase Increased Blood Creatinine Increased Brain Oedema Cardio-Respiratory Arrest Cervical Vertebral Fracture Coma Corneal Reflex Decreased Electrocardiogram T Wave Amplitude Increased Fall Hyperkalaemia Laboratory Test Abnormal Malaise Mydriasis Pallor Sinus Tachycardia Ventricular Fibrillation					

Date:07/24/02ISR Number: 3952698-XReport Type:Expedited (15-DaCompany Report #B0274253A  
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Acute Coronary Syndrome		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -		Blood Creatine		Tahor	C		ORAL
40MG Per day							
Initial or Prolonged		Phosphokinase Mb		Discotrine	C	Glaxo Wellcome	
TRANSDERMAL	10MGD	Per day					
		Increased		Kardegic	C		ORAL
160MG Per day							
		Chest Pain		Monotildiem	C	Glaxo Wellcome	ORAL
200MG Per day							
		Troponin I Increased					

Date:07/24/02ISR Number: 3952699-1Report Type:Expedited (15-DaCompany Report #B0274315A  
Age:35 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Angioneurotic Oedema
Initial or Prolonged	Dyspnoea
Disability	Face Oedema
	Gamma-Glutamyltransferase
	Increased
	Hepatic Enzyme Increased
	Myalgia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Oedema Peripheral

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1 WK		Zyban	PS	Glaxo Wellcome	ORAL

Date:07/24/02ISR Number: 3952836-9Report Type:Direct Company Report #CTU 172860  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anorexia Anxiety		Wellbutrin Sr 150mg Glaxo/Smit	PS	Glaxo/Smit	ORAL
150MG BID		Confusional State					
ORAL		Mydriasis Tachycardia		Fluoxetine 10mg Generic	SS		ORAL
10MG QD ORAL		Tremor Vomiting					

Date:07/25/02ISR Number: 3953122-3Report Type:Expedited (15-DaCompany Report #A0374100A  
 Age:24 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Per day	4 DAY	Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus					

Date:07/25/02ISR Number: 3954978-0Report Type:Expedited (15-DaCompany Report #B0274041A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Other	Psoriasis	Foreign Literature Health	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS	ORAL
150 MG /		Professional			
TWICE PER DAY					
/ ORAL					

Date:07/25/02ISR Number: 3954981-0Report Type:Expedited (15-DaCompany Report #B0274017A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG/ TWICE PER DAY /	Dehydration Dermatitis Exfoliative Ectropion	Foreign Literature Health	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
		Malaise	Professional				
		Mean Cell Volume					
		Increased Staphylococcal Infection		Emollient	C		

Date:07/25/02ISR Number: 3954983-4Report Type:Expedited (15-DaCompany Report #B0093634A  
Age:43 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Face Oedema Malaise

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Neutrophil Count Increased Psoriasis	Report Source	Product	Role	Manufacturer	Route
ORAL		Red Blood Cell Sedimentation Rate Increased	Foreign Literature Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
				Cyclosporin	C		

Date:07/26/02ISR Number: 3953754-2Report Type:Expedited (15-DaCompany Report #B0273542A  
 Age:35 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG	Unknown	1 DAY		Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY (INHALATION)			200MCG		Beclomethasone	C	Glaxo Wellcome	
per day			Twice					
RESPIRATORY (INHALATION)			2PUFF		Salbutamol	C	Glaxo Wellcome	
			Unknown					

Date:07/26/02ISR Number: 3956680-8Report Type:Expedited (15-DaCompany Report #B0089512A  
 Age:37 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG /			Foreign Literature Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
TWICE PER DAY				Company Representative				
/ ORAL					Co-Proxamol	C		
					Dothiepin	C		

Coal Tar C  
Corticosteroid C  
Dithranol C

Date:07/29/02ISR Number: 3954537-XReport Type:Expedited (15-DaCompany Report #B0274449A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Zyban	PS	Glaxo Wellcome	ORAL

Date:07/29/02ISR Number: 3954541-1Report Type:Expedited (15-DaCompany Report #B0274763A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 7 WK		Keratitis Sjogren'S Syndrome		Zyban	PS	Glaxo Wellcome	ORAL

Date:07/29/02ISR Number: 3954542-3Report Type:Expedited (15-DaCompany Report #B0275022A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1TAB Per day	8 DAY	Abortion Spontaneous		Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/29/02ISR Number: 3954606-4Report Type:Direct  
Age:16 YR Gender:Female I/FU:I

Company Report #CTU 173068

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 100 MG TWICE Initial or Prolonged DAILY ORAL		Abnormal Behaviour Crying Eye Disorder		Bupropion 100mg Teva	PS		ORAL
		Feeling Cold Hypersomnia Incoherent Insomnia Malaise Medication Error Mental Disorder Mood Altered Overdose Speech Disorder Suicide Attempt Thinking Abnormal Trance		Concerta	C		

Date:07/29/02ISR Number: 3954989-5Report Type:Direct  
Age:27 YR Gender:Female I/FU:I

Company Report #CTU 173122

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1TAB PO QD X 3 D , THEN BID		Muscle Spasms		Zyban 150mg Tbsr	PS	Tbsr	ORAL
				Seroquel	C		
				Serzone	C		
				Prozac	C		
				Imipramine	C		
				Buspar	C		

Date:07/30/02ISR Number: 3955018-XReport Type:Expedited (15-DaCompany Report #B0271436A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Health	Zyban	PS	Glaxo Wellcome	ORAL
1	MON	Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus	Professional				

Date:07/30/02ISR Number: 3955033-6Report Type:Expedited (15-DaCompany Report #B0275127A  
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Zyban	PS	Glaxo Wellcome	ORAL
17	DAY	Depression Dissociative Disorder Suicidal Ideation					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/30/02ISR Number: 3955607-2Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #CTU 173219

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly	100MG QD	Anotia		Bupropion Sr	PS		
STARTED		Maternal Drugs Affecting					
BEFORE		Foetus					
PREGNANCY		Microtia					
PRN VERY				Desonide Ointment	SS		
SPARINGLY							
EARLY							
PREGNANCY				Metronidazole Lotion	C		

Date:07/30/02ISR Number: 3955958-1Report Type:Expedited (15-DaCompany Report #A0072356A  
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required	100 MG / SEE	Acquired Porphyria	Health Professional Company	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
Intervention to Prevent Permanent	DOSAGE TEXT /	Anxiety	Representative				
Impairment/Damage	ORAL	Clonic Convulsion					
VARIABLE DOSE		Convulsion		Diazepam (Formulation			
/ ORAL		Cytomegalovirus Antibody Positive		Unknown) (Diazepam)	SS		ORAL
		Depersonalisation					
		Depressed Level Of Consciousness		Bismuth Subsalicylate (Formulation			
		Dissociation					

Drug Toxicity  
 Dystonia  
 Encephalopathy  
 Epstein-Barr Virus  
 Antibody Positive  
 Joint Stiffness  
 Mood Altered  
 Muscle Contracture  
 Nail Discolouration  
 Nail Disorder  
 Pressure Of Speech  
 Rash Erythematous  
 Skin Discolouration  
 Staring  
 Tardive Dyskinesia  
 Thyroid Neoplasm  
 Tremor  
 Visual Disturbance  
 Weight Decreased

Unknown) (Bismuth  
 Subsaliolate) SS  
 Triamcinolone  
 Acetonide C  
 Melatonin C  
 Herbal Medication C

Date:07/31/02ISR Number: 3955787-9Report Type:Direct  
 Age:17 YR Gender:Male I/FU:I

Company Report #CTU 173243

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin 150 Mg-Sr			
Other		Decreased Appetite		1 Po Bid	PS		ORAL
1 PO BID		Dehydration					
		Emotional Distress					
		Insomnia					
		Sunburn					

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Other	Alanine Aminotransferase	Zyban	PS	Glaxo Wellcome	ORAL
16 DAY	Increased	Nicotine Patch	C	Glaxo Wellcome	
TRANSDERMAL	Aspartate	Ikaran	C		
10MG per day	Aminotransferase	Adepal	C		
YR	Increased				
	Blood Creatine				
	Phosphokinase Increased				
	Blood Lactate				
	Dehydrogenase Increased				
	Insomnia				
	Muscle Spasms				
	Myalgia				

Date:08/01/02ISR Number: 3956398-1Report Type:Expedited (15-DaCompany Report #B0274315A  
Age:32 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Angioneurotic Oedema
Initial or Prolonged	Arthralgia
Disability	C-Reactive Protein
	Increased
	Dyspnoea
	Face Oedema
	Liver Function Test

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Abnormal  
Myalgia  
Oedema Peripheral

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	8 DAY		Zyban	PS	Glaxo Wellcome	ORAL

Date:08/01/02ISR Number: 3956413-5Report Type:Expedited (15-DaCompany Report #B0275303A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN		Physical Abuse		Zyban	PS	Glaxo Wellcome	

Date:08/01/02ISR Number: 3958181-XReport Type:Expedited (15-DaCompany Report #A202614  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Blood Sodium Abnormal Blood Testosterone Decreased	Health Professional	Zoloft Tablets Neurontin Paxil Wellbutrin Ambien Albuterol Inhaler Atenolol Folic Acid Zocor Hydrochlorothiazide	PS SS SS SS C C C C C C		

Date:08/01/02ISR Number: 3958565-XReport Type:Expedited (15-DaCompany Report #B0274494A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1500 MG / PER	Anorexia Bradycardia Bradyphrenia Depression	Literature Health Professional	Eskalith (Formulation Unknown) (Lithium Carbonate)	PS		

DAY

Disturbance In Attention

Drug Ineffective	Topiramate	
Drug Interaction	(Formulation	
Drug Level Increased	Unknown)	
Fatigue	(Topiramate)	SS
Lethargy	Wellbutrin	
Medication Error	(Formulation	
Memory Impairment	Unknown) (Bupropion	
Nausea	Hydrochloride)	SS
Neurotoxicity	Semisodium Valproate	
Nystagmus	(Formulation	
Tremor	Unknown) (Divalproex	
Weight Increased	Sodium)	SS
	Citalopram	C

Date:08/02/02ISR Number: 3956954-0Report Type:Expedited (15-DaCompany Report #A0372798A

Age:34 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Depressed Level Of
Initial or Prolonged	Consciousness
	Grand Mal Convulsion
	Intentional Misuse

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
9000MG	cumulative dose	Nystagmus Respiratory Depression	Wellbutrin Sr	PS	Glaxo Wellcome	ORAL
			Ethanol	C		ORAL
			Metabolite Herbal Supplement	C		
			Enbrel	C		ORAL

Date:08/02/02ISR Number: 3956957-6Report Type:Expedited (15-DaCompany Report #A0376136A  
Age: YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly	100MG	Unknown	2 YR	Anotia	Wellbutrin	PS	Glaxo Wellcome	
TOPICAL				Maternal Drugs Affecting	Metronidazole	C	Glaxo Wellcome	
TOPICAL				Foetus	Desonide	C		
				Microtia				

Date:08/02/02ISR Number: 3957744-5Report Type:Direct Company Report #CTU 173451  
Age:39 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	600MG/24HOURS	1 DAY		Amnesia	Wellbutrin	PS		
Initial or Prolonged				Blindness Transient				
				Productive Cough				
				Tremor				

Date:08/02/02ISR Number: 3960456-5Report Type:Periodic Company Report #2001AP05222  
Age:48 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Amitriptylline Bupropion	PS SS		

Date:08/05/02ISR Number: 3957571-9Report Type:Expedited (15-DaCompany Report #A0062462A  
 Age:61 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day 1.5MG per day 12.5MG per day 40MG per day		Angiopathy Diplopia Eye Disorder Vision Blurred		Zyban Xanax Triamterene Prinivil	PS C C C	Glaxo Wellcome Glaxo Wellcome	ORAL

Date:08/05/02ISR Number: 3957575-6Report Type:Expedited (15-DaCompany Report #B0120942A  
 Age:20 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Arthritis Reactive Asthma Fungal Infection Heart Rate Increased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Hypersensitivity Osteoarthritis Tobacco Abuse	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	26 DAY	Yersinia Infection		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
				Decortin	C		

Date:08/05/02ISR Number: 3957576-8Report Type:Expedited (15-DaCompany Report #B0275736A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	2TAB per day	Aggression		Zyban	PS	Glaxo Wellcome	
Hospitalization - UNKNOWN	Initial or Prolonged	Agitation Depression Palpitations Panic Attack					

Date:08/05/02ISR Number: 3957804-9Report Type:Direct Company Report #CTU 173480  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150 MG PO TD	Chest Discomfort		Wellbutrin	PS		ORAL
Other	1 DAY	Diarrhoea Heart Rate Increased		Glucophage	C		
				Avandia	C		
				Precose	C		
				Atarax	C		
				Ginko	C		

Date:08/07/02ISR Number: 3958753-2Report Type:Expedited (15-DaCompany Report #A0375990A  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 150MG Twice Initial or Prolonged per day	Back Injury Convulsion Headache	Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG See		Zyban	SS	Glaxo Wellcome	ORAL
dosage text	3 MON	Zocor	C		
		Stool Softener	C		

Date:08/07/02ISR Number: 3958754-4Report Type:Expedited (15-DaCompany Report #A0376093A  
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	19 DAY	Agitation Arthralgia Cyanosis Difficulty In Walking Insomnia Oedema Peripheral Rash Urticaria		Zyban        No Concurrent Medication	PS        C	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/07/02ISR Number: 3958755-6Report Type:Expedited (15-DaCompany Report #A0376196A

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hyponatraemia Inappropriate Antidiuretic Hormone Secretion		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:08/07/02ISR Number: 3958756-8Report Type:Expedited (15-DaCompany Report #A0376485A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Retinal Vein Thrombosis		Zyban	PS	Glaxo Wellcome	ORAL

Date:08/07/02ISR Number: 3960398-5Report Type:Expedited (15-DaCompany Report #B0274494A

Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anorexia Bradycardia	Literature Health	Eskalith (Lithium Carbonate)	PS		
1500 MG / PER DAY		Bradyphrenia	Professional				
		Depression Disturbance In Attention Drug Interaction Drug Level Increased Fatigue Lethargy Medication Error Memory Impairment Nausea Neurotoxicity Nystagmus Tremor Weight Decreased Weight Increased		Topiramate (Topiramate) Wellbutrin (Bupropion Hydrochloride) Semisodium Valproate (Divalproex Sodium) Citalopram	SS   SS SS C		



Date:08/08/02ISR Number: 3959455-9Report Type:Expedited (15-DaCompany Report #A0370671A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Twice			Blood Calcium Increased	Wellbutrin	PS	Glaxo Wellcome	ORAL
			Fatigue				
per day	3	MON	Grand Mal Convulsion	Lipitor	C		ORAL
20MG Per day			Loss Of Consciousness	Amitriptyline	C		ORAL
25MG Per day							

Date:08/08/02ISR Number: 3959461-4Report Type:Expedited (15-DaCompany Report #B0275697A  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability							
150MG As			Deafness	Zyban	PS	Glaxo Wellcome	ORAL
			Tinnitus				
directed				Dihydrocodeine	C		ORAL
				Hormone Replacement			
				Therapy	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/08/02ISR Number: 3959687-XReport Type:Direct  
Age:20 YR Gender:Female I/FU:I

Company Report #CTU 173780

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion		Buprion-Wellbutrin-	PS		ORAL
ORAL							

Date:08/08/02ISR Number: 3959912-5Report Type:Direct  
Age:33 YR Gender:Female I/FU:I

Company Report #CTU 173781

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion		Wellbutrin Sr	PS		ORAL
150MG BID PO				Alesse	C		
				Flovent	C		
				Ambien	C		
				Xanax	C		
				Effexor Xr	C		
				Imitrex	C		

Date:08/09/02ISR Number: 3960002-6Report Type:Expedited (15-DaCompany Report #A0376700A  
Age:19 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Coma Incoherent Intentional Misuse Suicide Attempt		Wellbutrin No Concurrent Medication	PS  C	Glaxo Wellcome	ORAL

Date:08/09/02ISR Number: 3960005-1Report Type:Expedited (15-DaCompany Report #B0264307A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 3000MG Single Initial or Prolonged dose		Bronchial Obstruction Clonic Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
1 DAY							

280MG Single			Depressed Level Of	Deroxat	SS	Glaxo Wellcome	ORAL
dose	1	DAY	Consciousness				
1 DAY			Intentional Misuse	Seresta	SS		ORAL
180MG Single			Muscle Contractions	Lexomil	SS		ORAL
dose	1	DAY	Involuntary				
1 DAY			Somnolence	Alcohol	C		
			Suicide Attempt	Nifuroxazide	C	Glaxo Wellcome	ORAL
16MG Single			Tracheal Obstruction	Arestal	C		ORAL
dose	1	DAY					

Date:08/09/02ISR Number: 3960088-9Report Type:Direct Company Report #CTU 173829  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser		Wellbutrin Sr	PS	Glaxcosmithkline	NASAL
150MG DAILY		Medication Error					
NASAL							

Date:08/09/02ISR Number: 3962187-4Report Type:Expedited (15-DaCompany Report #USA-2002-0000807  
Age:28 YR Gender:Female I/FU:F

Outcome	PT
Death	Accidental Overdose
	Drug Toxicity
	Ecchymosis

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Pulmonary Congestion Pulmonary Oedema	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer Health Professional Other	Morphine Sulfate (Similar To Nda 19-516)(Morphine Sulfate)Unknown Hydromorphone Hcl (Similar To Ind 38,424)(Hydromorphon e Hydrochloride)Unknow Fluoxetine (Fluoxetine) Tablet	PS  SS SS		ORAL
ORAL				Diazepam (Diazepam) Injectable Bupropion (Amfebutamone) Unknown Promethazine (Promethazine) Unknown Propranolol (Propranolol ) Tablet Oxepam(Oxazepam) Unknown Caffeine (Caffeine) Unknown Nicotine(Nicotine)Un known	SS  SS  SS  SS  SS  SS		

Date:08/12/02ISR Number: 3960666-7Report Type:Expedited (15-DaCompany Report #B0270679A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State		Zyban	PS	Glaxo Wellcome	
Other	150MG	Unknown		Quetiapine	C		
UNKNOWN		Hyperhidrosis					
1000MG per day		Hypertension					

5MG per day	Loss Of Consciousness	Clonazepam	C
	Overdose	Alcohol	C
	Serotonin Syndrome		
	Somnolence		
	Tachycardia		

Date:08/12/02ISR Number: 3961388-9Report Type:Periodic Company Report #11850310  
 Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 600	Pancreatitis Acute	Health Professional Company Representative	Sustiva Caps 600 Mg (Efavirenz)	PS		ORAL
MILLIGRAM, DAY, ORAL			Stavudine Didanosine Wellbutrin(Bupropion Hcl)	SS SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/13/02ISR Number: 3961228-8Report Type:Expedited (15-DaCompany Report #A0163498A  
 Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Abortion Spontaneous	Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice			Complications Of Maternal				
per day	57	DAY	Exposure To Therapeutic Drugs Insomnia Maternal Drugs Affecting Foetus Pruritus				

Date:08/13/02ISR Number: 3961229-XReport Type:Expedited (15-DaCompany Report #A0376142A  
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -			Alopecia	Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Unknown	5	MON	Chest Pain	Synthroid	C	Glaxo Wellcome	
Initial or Prolonged			Gastrointestinal Disorder	Tenormin	C		
			Hostility	Premarin	C		
			Memory Impairment				
			Mood Swings				

Date:08/13/02ISR Number: 3961230-6Report Type:Expedited (15-DaCompany Report #A0376485A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Retinal Vein Thrombosis	Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:08/13/02ISR Number: 3961232-XReport Type:Expedited (15-DaCompany Report #A0377262A  
 Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Anxiety	Wellbutrin	PS	Glaxo Wellcome	ORAL
300MG Per day 2 YR	Aphasia	Estrogen Replacement	C		ORAL
3MG Per day	Balance Disorder	Advil	C	Glaxo Wellcome	ORAL
	Constipation	Calcium	C		ORAL
	Dry Mouth	Fish Oil	C		ORAL
	Dysgeusia				
	Insomnia				
	Libido Increased				
	Memory Impairment				
	Nausea				
	Pharyngolaryngeal Pain				
	Tinnitus				

Date:08/13/02ISR Number: 3961248-3Report Type:Expedited (15-DaCompany Report #B0276351A  
Age:35 YR Gender:Male I/FU:I

Outcome	PT
Other	Anxiety
	Choking Sensation
	Confusional State
	Histrionic Personality
	Disorder
	Panic Attack
	Paranoia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Sensation Of Foreign Body Vomiting	Report Source	Product	Role	Manufacturer	Route
150MG				Zyban	PS	Glaxo Wellcome	ORAL
Variable dose	8 DAY						

Date:08/15/02ISR Number: 3963976-2Report Type:Expedited (15-DaCompany Report #WAES 01072531  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 25		Amnesia	Consumer	Tab Vioxx 25 Mg	PS		ORAL
Initial or Prolonged Disability		Asthenia	Health				
150 MG/BID		Back Pain	Professional	Wellbutrin 150 Mg	SS		
		Cervical Spinal Stenosis		Astelin	C		
		Depression		Delestrogen	C		
		Electrocardiogram St Segment Depression		Flexeril	C		
		Faecal Incontinence		Lipitor	C		
		Gait Spastic		Nasonex	C		
		Grand Mal Convulsion		Profen Ii	C		
		Hypoaesthesia		Ultram	C		
		Impaired Driving Ability		Clonazepam	C		
		Intervertebral Disc Protrusion					
		Loss Of Consciousness					
		Loss Of Employment					
		Myelopathy					
		Pain					
		Petit Mal Epilepsy					
		Quadripareisis					
		Spinal Cord Injury					
		Syncope					
		Urinary Incontinence					

Date:08/16/02ISR Number: 3962920-1Report Type:Expedited (15-DaCompany Report #A0377435A  
 Age: Gender:Unknown I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL

Date:08/16/02ISR Number: 3962928-6Report Type:Expedited (15-DaCompany Report #B0276132A  
Age:45 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Aspartate
Initial or Prolonged	Aminotransferase
	Increased
	Blood Albumin Decreased
	Blood Potassium Decreased
	Blood Sodium Decreased
	Blood Urea Decreased
	C-Reactive Protein
	Increased
	Diabetes Mellitus
	Erythema Multiforme
	Gamma-Glutamyltransferase
	Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
11 DAY		Heart Rate Increased Lymphocyte Count Decreased					
150MG Twice		Neutrophil Count		Zyban	PS	Glaxo Wellcome	ORAL
per day		Increased		Mediator	SS		ORAL
		Prurigo					
		Pyrexia Rosacea White Blood Cell Count Increased					

Date:08/16/02ISR Number: 3962996-1Report Type:Direct Company Report #CTU 174352  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Decreased Appetite Gun Shot Wound Medication Error		Zyban 150mg Sr Catalytica Pharmaceuticals	PS	Catalytica Pharmaceuticals	ORAL
1 TABLET 3							
DAYS ORAL				Zyban	SS	Catalytica Pharmaceuticals	ORAL
1 TAB TWICE							
DAILY ORAL							

Date:08/19/02ISR Number: 3963401-1Report Type:Expedited (15-DaCompany Report #A0371180A  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Clonic Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day				Marijuana	C		

Naprosyn C  
Flexeril C

Date:08/19/02ISR Number: 3963410-2Report Type:Expedited (15-DaCompany Report #B0276134A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Clonic Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
14UNIT Single							
Initial or Prolonged		Intentional Misuse					
dose	1 DAY						
		Suicide Attempt		Cimicifuga	C		ORAL
1 DAY							
		Toxicologic Test Abnormal		Alcohol	C		

Date:08/19/02ISR Number: 3963419-9Report Type:Direct Company Report #CTU 174419  
Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Intentional Misuse		Topomax	PS		
25MG BID							
Initial or Prolonged		Suicide Attempt		Wellbutrin	SS		
100MG BID							

Date:08/19/02ISR Number: 3964158-0Report Type:Direct Company Report #CTU 174530  
Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Convulsion		Welbutrin-	PS		
Initial or Prolonged				Venlafaxin	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/19/02ISR Number: 3964175-0Report Type:Direct  
Age:30 YR Gender:Female I/FU:I

Company Report #CTU 174511

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Bruxism		Wellbutrin Sr 150mg			
Intervention to		Dizziness		Wellbutrinsr	PS		ORAL
150MG 1X FOR							
Prevent Permanent		Dyskinesia					
3 ORAL, 150MG							
Impairment/Damage		Erythema					
2X PER							
		Headache					
		Hyperacusis					
		Insomnia					
		Joint Stiffness					
		Nausea					
		Pyrexia					

Date:08/19/02ISR Number: 3964735-7Report Type:Expedited (15-DaCompany Report #2002119339GB  
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Foreign	Depo-Medrone			
		Drug Interaction	Other	(Methylprednisolone)			
		Lumbar Vertebral Fracture		Suspension, Sterile	PS		
				Zyban Prolonged			
				Realeased Tablets	SS		ORAL
ORAL							

Date:08/20/02ISR Number: 3964009-4Report Type:Expedited (15-DaCompany Report #A0377456A  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Respiratory Depression		Bupropion			
				Hydrochloride	PS	Glaxo Wellcome	
				Cocaine	SS		
				Morphine	C		
				Heroin	C		
				Codeine	C		

Date:08/20/02ISR Number: 3964081-1Report Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-11850310  
Age:52 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Pancreatitis Acute	Health Professional	Sustiva Caps 600 Mg Stavudine	PS SS	Bristol-Myers Squibb Company Bristol-Myers Squibb Company	ORAL ORAL
unknown start date-prior to 06-Dec-00			Videx Ec Caps	SS	Bristol-Myers Squibb Company	ORAL
at hour of sleep AM + 1600 x 30 days			Wellbutrin	SS		ORAL

Date:08/20/02ISR Number: 3964535-8Report Type:Direct Company Report #CTU 174626  
Age:29 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration ONE DOSE PO QD X 3D OTHER	Tremor		Zyban 150 Mg Tbsr	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

BID

Date:08/20/02 ISR Number: 3964561-9 Report Type:Direct  
 Age:39 YR Gender:Female I/FU:I

Company Report #CTU 174643

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pruritus		Wellbutrin Sr 100mg	PS		ORAL
100MG ORAL	6 DAY	Rash Papular		Calcium	C		
				Depakote	C		
				Multivitamin	C		

Date:08/21/02 ISR Number: 3964459-6 Report Type:Expedited (15-DaCompany Report #A0378334A  
 Age:14 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Serum Sickness		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice							
Initial or Prolonged							
per day	18 DAY			Concerta	C		
1 YR							

Date:08/21/02 ISR Number: 3964460-2 Report Type:Expedited (15-DaCompany Report #B0130344A  
 Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Abdominal Adhesions	Health	Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day		Blood Ethanol Increased	Professional	Dothiepin	SS		ORAL
50MG See		Cardiac Arrest					
dosage text	47 DAY	Coronary Artery		Alcohol	SS		ORAL
		Atherosclerosis		Prempak-C	C		ORAL
.625MG		Intentional Misuse					
Unknown							

RESPIRATORY Pulmonary Congestion Salbutamol C Glaxo Wellcome  
 (INHALATION) 100MCG As Pulmonary Oedema  
 required Ventricular Hypertrophy  
 UNKNOWN 5MG As Diazepam C  
 required  
 UNKNOWN 40MG Per day Fluoxetine C  
 UNKNOWN 7.5MG At Zopiclone C  
 night  
 RESPIRATORY Beclomethasone C Glaxo Wellcome  
 (INHALATION) 100U Unknown

Date:08/21/02ISR Number: 3964467-5Report Type:Expedited (15-DaCompany Report #B0276547A  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 14 DAY		Intentional Misuse	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged 50MG per day		Suicide Attempt		Zoloft	SS		ORAL
4MG per day				Lexomil	C		ORAL

Date:08/21/02ISR Number: 3964473-0Report Type:Expedited (15-DaCompany Report #B0277109A  
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening UNKNOWN Hospitalization - per day 6 DAY	150MG Twice	Monoplegia		Zyban	PS	Glaxo Wellcome	
Initial or Prolonged Other		Respiratory Failure					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/21/02ISR Number: 3965979-0Report Type:Expedited (15-DaCompany Report #A0361723A  
 Age:10 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Abnormal Behaviour Condition Aggravated Grand Mal Convulsion Orchidectomy	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
200 MG/TWICE PER DAY/ORAL							

Date:08/21/02ISR Number: 3965982-0Report Type:Expedited (15-DaCompany Report #A0377696A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Diplopia Strabismus	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
450 MG / IN THE MORNING/ ORAL							

Date:08/23/02ISR Number: 3965386-0Report Type:Expedited (15-DaCompany Report #B0264661A  
 Age:69 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death UNKNOWN	150MG	Acute Myocardial Per day 5 DAY Infarction Bronchopneumonia		Zyban  Tramadol Hydrochloride	PS  C	Glaxo Wellcome	
UNKNOWN		Hypoxia Malaise Nausea					



Date:08/23/02ISR Number: 3965387-2Report Type:Expedited (15-DaCompany Report #B0265519A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day		Cholestasis		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Hepatic Neoplasm Hyperplasia Systemic Lupus Erythematous Rash					

Date:08/23/02ISR Number: 3965388-4Report Type:Expedited (15-DaCompany Report #B0271436A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1 MON		Abortion Spontaneous		Zyban	PS	Glaxo Wellcome	ORAL
		Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/23/02ISR Number: 3965915-7Report Type:Direct  
 Age:23 YR Gender:Female I/FU:I

Company Report #CTU 174894

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Feeling Cold Pain In Extremity		Wellbutrin Sr 150mg Glaxo-Wellcome	PS	Glaxo-Wellcome	
PARENTERAL	150MG	SR QD					
PARENTERAL		Tremor					

Date:08/23/02ISR Number: 3967053-6Report Type:Expedited (15-DaCompany Report #A0377881A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia Chronic Obstructive Pulmonary Disease	Consumer	Nicorette Otc Gum (Nicotine Polacrilex)	PS		ORAL
ORAL							
ORAL		Drug Dependence Emphysema Hyperphagia		Wellbutrin (Bupropion Hydrochloride)	SS		ORAL

Date:08/26/02ISR Number: 3966012-7Report Type:Expedited (15-DaCompany Report #B0264067A  
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Chest Discomfort		Zyban	PS	Glaxo Wellcome	ORAL
300MG per day	62 DAY	Malaise Sudden Death					

Date:08/26/02ISR Number: 3966013-9Report Type:Expedited (15-DaCompany Report #B0271008A  
 Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 15 DAY	Arthralgia	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Erythema	Nicorette	C	Glaxo Wellcome	ORAL
Other	Hepatic Steatosis				
	Joint Swelling				
	Malaise				
	Myalgia				
	Oedema Peripheral				
	Pyrexia				
	Serum Sickness				
	Urticaria				

Date:08/26/02ISR Number: 3966020-6Report Type:Expedited (15-DaCompany Report #B0277513A  
Age:61 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Unknown 26 DAY	Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
Other		Myalgia					
		Oedema Peripheral					
		Red Blood Cell					
		Sedimentation Rate					
		Increased					
		Urticaria					
		White Blood Cell Count					
		Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/27/02ISR Number: 3966329-6Report Type:Expedited (15-DaCompany Report #A0376142A  
 Age:65 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Unknown 5 MON		Abdominal Pain Lower	Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Alopecia	Synthroid	C	Glaxo Wellcome	
		Chest Pain	Tenormin	C		
		Gastrointestinal Disorder	Premarin	C		
		Hostility				
		Memory Impairment				
		Mood Swings				

Date:08/27/02ISR Number: 3966330-2Report Type:Expedited (15-DaCompany Report #A0378086A  
 Age:27 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice		Depression	Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day		Disturbance In Attention				
19 DAY		Fear	Paxil	SS	Glaxo Wellcome	ORAL
		Hallucination, Visual	Ambien	C		
		Irritability	Trazodone	C		
500MG Twice		Sleep Walking	Depakote	C		
per day		Suicide Attempt				
YR			Ritalin	C		

Date:08/27/02ISR Number: 3966336-3Report Type:Expedited (15-DaCompany Report #A0378902A  
 Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly 200MG Per day		Coarctation Of The Aorta	Wellbutrin	PS	Glaxo Wellcome	
		Maternal Drugs Affecting Foetus	Depakote	C		

Pregnancy  
Ventricular Septal Defect

Date:08/27/02ISR Number: 3968149-5Report Type:Expedited (15-DaCompany Report #B0276683A  
Age:26 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Motor Dysfunction	Foreign	Zyban Tablet - Zyban			
Hospitalization -	Respiratory Disorder	Health	(Bupropion			
Initial or Prolonged	Suicide Attempt	Professional	Hydrochloride)	PS		ORAL
150 MG/TWICE		Company				
Disability		Representative	Mineral Oil			
PER DAY/ORAL			Injection (Mineral			
Required			Oil)	SS		
Intervention to						
Prevent Permanent						
INTRAVENOUS	INTRAVENOUS					
Impairment/Damage			Citalopram	C		
			Sertraline	C		

Date:08/27/02ISR Number: 3968171-9Report Type:Expedited (15-DaCompany Report #A0363292A  
Age:44 YR Gender:Male I/FU:F

Outcome	PT
Death	Atherosclerosis Cardiac Disorder Cardiac Failure Acute Cardio-Respiratory Arrest Ischaemia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Myocardial Infarction Ventricular Arrhythmia	Report Source	Product	Role	Manufacturer	Route
ORAL			Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
				Salbutamol Sulphate	C		
				Salmeterol Xinafoate	C		
				Irbesartan	C		

Date:08/27/02ISR Number: 3969095-3Report Type:Expedited (15-DaCompany Report #A0377076A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Antidepressant Drug Level Above Therapeutic	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
4200 MG/SEE		Grand Mal Convulsion					
DOSAGE		Intentional Misuse					
TEXT/ORAL	1 HR	Metabolic Acidosis					

Date:08/27/02ISR Number: 3969109-0Report Type:Expedited (15-DaCompany Report #A0378058A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Loss Of Consciousness Overdose	Health Professional	Wellbutrin Sr Tablet-Controlled Release	PS		ORAL
27 GRAM (S)/							
SINGLE DOSE/							
ORAL							

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 360 MG DAILY Initial or Prolonged PO	Blood Pressure Decreased  Cystitis	Consumer	Cardizem Cd	PS		ORAL
150 MG DAILY  PO	Drug Interaction Medication Error		Zyban Tablet (Bupropion Hcl)	SS		ORAL
50 MG DAILY  ORAL	Myocardial Infarction		Atenolol	SS		ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - Initial or Prolonged	Accident Dyspnoea Echocardiogram Abnormal Hypernatraemia Hypocapnia Hypotonia Hypoxia Loss Of Consciousness		Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/28/02ISR Number: 3968133-1Report Type:Expedited (15-DaCompany Report #D0039102A  
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	5 DAY	Acute Psychosis Disturbance In Attention Insomnia	Foreign Literature Health	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL		Schizophreniform Disorder Suicidal Ideation Vision Blurred	Professional				

Date:08/29/02ISR Number: 3967700-9Report Type:Expedited (15-DaCompany Report #B0118244A  
Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG Twice per day	11 DAY	Angina Unstable Aortic Valve Stenosis Cardiac Disorder Cardiac Failure Congestive Chest Pain Chronic Obstructive Pulmonary Disease Dyspepsia Dyspnoea Dyspnoea Exertional Heart Rate Increased Hypokinesia Left Ventricular Failure Pharyngolaryngeal Pain Ventricular Hypertrophy		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL

Date:08/29/02ISR Number: 3967709-5Report Type:Expedited (15-DaCompany Report #D0038838A  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Disability	Aggression	Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day 6 WK					
Other	Bipolar Disorder	Seroxat	SS	Glaxo Wellcome	
UNKNOWN	92 DAY				
	Crying				
	Diarrhoea				
	Gastrointestinal Disorder				
	Hyperhidrosis				
	Hypomania				
	Migraine				
	Nausea				
	Sleep Disorder				
	Tachycardia				
	Vertigo				
	Weight Decreased				

Date:08/29/02ISR Number: 3968415-3Report Type:Expedited (15-DaCompany Report #C2002-2710.01  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Cystitis	Consumer	Zyban Tablets 150 Mg			
Initial or Prolonged	Hypotension	Other	Glaxosmithkline	PS	Glaxosmithkline	ORAL
150MG DAY						
Other	Medication Error					
ORAL						
			Atenolol Tablets			
			50mg Mylan	SS	Mylan	ORAL
50MG DAY ORAL						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

360MG DAY  
 ORAL  
 Cardizem Cd Aventis SS Aventis ORAL

Date:08/30/02ISR Number: 3968361-5Report Type:Expedited (15-DaCompany Report #B0274152A  
 Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG	Electric Shock		Zyban	PS	Glaxo Wellcome	
UNKNOWN	Twice	Tremor					
per day	26 DAY						

Date:08/30/02ISR Number: 3970052-1Report Type:Expedited (15-DaCompany Report #A103444  
 Age:79 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required	50 MG DAILY,	Anger	Consumer	Zoloft (Sertraline)	PS		ORAL
Intervention to	ORAL	Autoimmune Thyroiditis					
Prevent Permanent	Impairment/Damage	Balance Disorder		Wellbutrin Sr			
300 MG		Blood Pressure Increased		(Bupropion			
(BID), ORAL		Blood Thyroid Stimulating		Hydrochloride)	SS		ORAL
		Hormone Decreased					
		Cerebral Haemorrhage		Coumadin (Warfarin)	SS		
		Contusion		Effexor Xr			
		Convulsion		(Venlafaxine			
300 MG BID,		Cough		Hydrochloride)	SS		ORAL
ORAL		Drug Ineffective					
		Dysphonia		Amantadine	SS		
		Epistaxis		Monopril (Fosinopril			
		Fall		Sodium)	SS		
		Feeling Cold		Toprol Xl			
		Head Injury		(Metoprolol			
		Heart Rate Decreased		Succinate)	C		

Heart Rate Increased	Zebeta (Bisoprolol	C
Hostility	Fumarate)	C
Hyperhidrosis	Lanoxin (Digoxin)	C
Laceration	Lasix (Furosemide)	C
Medication Error	Synthroid	
Meningioma	(Levothyroxine	
Mood Swings	Sodium)	C
Overdose	Vitamin E	
Pain	(Tocopherol)	C
Palpitations	Estrogen (Estrogen	
Parkinson'S Disease	Nos)	C
Prothrombin Time	Zocor (Simvastatin)	C
Prolonged	Estradiol Vaginal	
Pulmonary Oedema	Ring (Estradiol)	C
Rib Fracture	Inderal (Propranolol	
Syncope	Hydrochloride)	C
Transient Ischaemic	Cardizem (Diltiazem	
Attack	Hydrochloride)	C
Vaginal Haemorrhage	Folic Acid	C
	Sinemet (Carbidopa /	
	Levodopa)	C
	Carzon	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/30/02ISR Number: 3970174-5Report Type:Expedited (15-DaCompany Report #A0151352A  
Age:21 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Agitation	Foreign	Zyban Tablet - Zyban	PS		ORAL
Intervention to SINGLE DOSE / Prevent Permanent ORAL Impairment/Damage		Cardiotoxicity Cardiovascular Disorder Electrocardiogram Qrs Complex Prolonged Hallucination Intentional Misuse Sinus Tachycardia Suicide Attempt Tremor	Literature Health Professional	Lorazepam	C		

Date:09/03/02ISR Number: 3970070-3Report Type:Direct Company Report #CTU 175458  
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG TID Initial or Prolonged ORAL		Grand Mal Convulsion		Wellbutrin Sr 100 Mg	PS		ORAL
				Zoloft	C		

Date:09/04/02ISR Number: 3969848-1Report Type:Expedited (15-DaCompany Report #B0114598A  
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Twice Hospitalization - per day 62 DAY Initial or Prolonged Other		Abdominal Pain Abnormal Behaviour Amnesia Anaphylactic Shock Angioneurotic Oedema Blister	Health Professional	Zyban	PS	Glaxo Wellcome	ORAL

Blood Bilirubin Increased  
Blood Pressure Increased  
Chest Pain  
Convulsion  
Diarrhoea  
Dry Skin  
Eczema  
Feeling Drunk  
Hypersensitivity  
Rhinitis  
Tinnitus

Date:09/04/02ISR Number: 3969850-XReport Type:Expedited (15-DaCompany Report #B0259361A  
Age:46 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300MG Per day 21 DAY Initial or Prolonged	Psoriasis		Zyban	PS	Glaxo Wellcome	ORAL

Date:09/04/02ISR Number: 3969853-5Report Type:Expedited (15-DaCompany Report #B0270906A  
Age:38 YR Gender:Male I/FU:F

Outcome	PT
Death	Agitation Delusional Disorder,

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Persecutory Type  
Injury

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
UNKNOWN	150MG As		Zyban	PS	Glaxo Wellcome	
directed	52 DAY					
UNKNOWN			Paracetamol + Codeine	C		
RESPIRATORY			Fenoterol + Ipratropium	C		
(INHALATION)						
UNKNOWN			Cocaine	C		

Date:09/04/02ISR Number: 3971338-7Report Type:Expedited (15-DaCompany Report #B0102029A  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG/PER		Drug Hypersensitivity	Foreign	Zyban	PS		
Initial or Prolonged DAY/	14 DAY	Haematuria	Literature				
		Proteinuria	Health Professional				

Date:09/05/02ISR Number: 3970620-7Report Type:Expedited (15-DaCompany Report #A0370880A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1TAB Twice		Retinal Artery Thrombosis	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
per day	3 WK	Retinal Exudates	Professional				
		Visual Acuity Reduced		Zyrtec	C	Glaxo Wellcome	

Date:09/05/02ISR Number: 3970626-8Report Type:Expedited (15-DaCompany Report #B0134204A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Coronary Artery Occlusion		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Haemochromatosis					
Hospitalization -		Impaired Driving Ability		Lormetazepam			
per day	4	Myalgia		(Noctamide)	C		ORAL
Initial or Prolonged		Myocardial Infarction		Fluoxetine			
1MG per day		Myocardial Ischaemia		Hydrochloride	C		
UNKNOWN	20MG per day			Alprazolam	C		ORAL
.25MG per day							

Date:09/05/02ISR Number: 3970630-XReport Type:Expedited (15-DaCompany Report #B0271161A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Drug Ineffective	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	24	Mean Cell Volume		Miniphase	SS		ORAL
1TAB Per day		Increased		Lexomil	SS		ORAL
.5TAB Twice		Peripheral Sensory					
per day		Neuropathy		Di-Antalvic	SS		ORAL
2UNIT Monthly		Spinal Osteoarthritis					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/05/02ISR Number: 3970631-1Report Type:Expedited (15-DaCompany Report #B0274419A

Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 31 DAY		Abortion Spontaneous		Zyban	PS	Glaxo Wellcome	ORAL
		Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus Pregnancy		Emergency Contraception	C		

Date:09/05/02ISR Number: 3970643-8Report Type:Expedited (15-DaCompany Report #B0278335A

Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG		Headache		Zyban	PS	Glaxo Wellcome	ORAL
Variable dose	2 DAY	Visual Disturbance		Aspirin	C		ORAL
75MG Unknown				Omeprazole	C		ORAL
10MG Unknown				Beclomethasone	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)	100MCG Twice			Salbutamol	C	Glaxo Wellcome	
per day				Simvastatin	C		ORAL
RESPIRATORY (INHALATION)				Diltiazem	C	Glaxo Wellcome	ORAL
20MG Per day							

Date:09/06/02ISR Number: 3972586-2Report Type:Expedited (15-DaCompany Report #A0379437A

Age: Gender: I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Retinal Detachment Vitreous Floaters	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE							
PER DAY/ORAL				Adderall	C		

Date:09/06/02ISR Number: 3972817-9Report Type:Expedited (15-DaCompany Report #A0072356A  
Age:45 YR Gender:Female I/FU:F

Outcome	PT
Disability	Acquired Porphyria
Required	Akathisia
Intervention to	Alopecia
Prevent Permanent	Anxiety
Impairment/Damage	Apathy
	Asthenia
	Autoimmune Thyroiditis
	Cardiolipin Antibody
	Positive
	Cerebellar Syndrome
	Chronic Fatigue Syndrome
	Circulatory Collapse
	Clonic Convulsion
	Cogwheel Rigidity
	Computerised Tomogram
	Abnormal

Freedom Of Information (FOI) Report

Dose	Duration	Depersonalisation	Report Source	Product	Role	Manufacturer	Route
100 MG ORAL		Cytomegalovirus Antibody Positive					
		Diaphragmatic Paralysis	Health Professional	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
		Diarrhoea	Company				
		Disturbance In Attention					
		Drug Withdrawal Syndrome	Representative	Diazepam (Diazepam)	SS		ORAL
VARIABLE DOSE/ ORAL		Dysarthria					
		Dyspnoea		Bismuth			
		Dysstasia		Subsalicylate			
		Dystonia		(Bismuth			
		Encephalopathy		Subsalicylate)	SS		
		Epstein-Barr Virus Antibody Positive		Triamcinolone			
		Erythema		Acetonide	C		
		Facial Palsy		Melatonin	C		
		Fall		Herbal Medication	C		
		Fatigue					
		Fibromyalgia					
		Hemiparesis					
		Loss Of Consciousness					
		Malaise					
		Nail Disorder					
		Nervous System Disorder					
		Nuclear Magnetic Resonance Imaging Abnormal					
		Optic Nerve Disorder					
		Optic Neuritis					
		Porphyria					
		Rash Macular					
		Red Blood Cell Sedimentation Rate Increased					
		Respiratory Alkalosis					
		Tardive Dyskinesia					
		Visual Acuity Reduced					
		Weight Decreased					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 50 MG		Balance Disorder	Consumer	Zoloft (Sertraline)	PS		ORAL
Intervention to (DAILY), ORAL		Cough					
Prevent Permanent Impairment/Damage		Drug Ineffective Fall Irritability		Wellbutrin Sr (Bupropion Hydrochloride)	SS		ORAL
300 MG (BID), ORAL		Pulmonary Oedema					
		Rib Fracture		Coumadin (Warfarin) Effexor Xr (Venlafaxine Hydrochloride)	SS SS		ORAL
300 MG (BID), ORAL				Amantadine Monopril (Fosinopril Sodium) Toprol Xl (Metoprolol Succinate)	SS SS C		

Freedom Of Information (FOI) Report

Zebeta (Bisoprolol Fumarate) C  
 Lanoxin (Digoxin) C  
 Synthroid (Levothyroxine Sodium) C  
 Vitamin E (Tocopherol) C  
 Estrogen (Estrogen Nos) C  
 Zocor(Simvastatin) C  
 Estradiol Vaginal Ring (Estradiol) C  
 Inderal (Propranolol Hydrochloride) C  
 Cardizem (Diltiazem Hydrochloride) C  
 Folic Acid C  
 Stinemet (Carbidopa/Levodopa) C  
 "Carzon" C  
 Lasix (Furosemide) C

Date:09/06/02ISR Number: 3972863-5Report Type:Expedited (15-DaCompany Report #B0270362A  
 Age:64 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Chest Discomfort Headache Hypertension	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
200 MG / PER DAY / ORAL		Obstructive Airways Disorder Orthopnoea Pulmonary Oedema	Company Representative	Labetalol Hydrochloride Tablet (Non-Us Product)	SS		ORAL
				Glimepiride Allopurinol Buflomedil Hydrochloride Quinapril	C C C		

Hydrochloride	C
Atorvastatin Calcium	C
Clopidogrel	
Bisulphate	C
Theophylline	C
Formoterol	C

Date:09/06/02ISR Number: 3973171-9Report Type:Expedited (15-DaCompany Report #B0278040A

Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Cerebrovascular Disorder Grand Mal Convulsion	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ORAL				Captopril Etodolac	C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/09/02ISR Number: 3974158-2Report Type:Direct  
Age:39 YR Gender:Male I/FU:I

Company Report #CTU 175983

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG PO BID Initial or Prolonged		Rhabdomyolysis		Wellbutrin Sr 150 Mg	PS		ORAL

Date:09/10/02ISR Number: 3972720-4Report Type:Expedited (15-DaCompany Report #B0274763A  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 7 WK Other		Keratitis	Health	Zyban	PS	Glaxo Wellcome	ORAL
		Sjogren'S Syndrome	Professional				

Date:09/10/02ISR Number: 3972739-3Report Type:Expedited (15-DaCompany Report #D0039257A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 4TAB per day 2 DAY		Blood Bicarbonate Decreased Blood Ph Abnormal Convulsion Disturbance In Attention Lactic Acidosis		Zyban	PS	Glaxo Wellcome	ORAL

Date:09/11/02ISR Number: 3973364-0Report Type:Expedited (15-DaCompany Report #B0135382A  
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 2 WK Initial or Prolonged		Chills	Health	Zyban	PS	Glaxo Wellcome	ORAL
Disability		Dehydration Dysuria Eosinophil Count	Professional	Nomegestrol	SS		ORAL

Increased  
Headache  
Mucosal Erosion  
Oedema Peripheral  
Pruritus  
Pyrexia  
Rash Generalised  
Skin Exfoliation  
Sleep Disorder  
Stevens-Johnson Syndrome

Date:09/12/02ISR Number: 3974391-XReport Type:Expedited (15-DaCompany Report #B0265019A  
Age:60 YR Gender:Male I/FU:F

Outcome PT  
Hospitalization - Asthenia  
Initial or Prolonged Claustrophobia  
Confusional State  
Convulsion  
Dyspnoea  
Fatigue  
Fear  
Gastrointestinal Disorder  
General Physical Health  
Deterioration

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN	150MG	Initial Insomnia Joint Stiffness Loss Of Consciousness		Zyban	PS	Glaxo Wellcome	
per day	30 DAY	Malaise Twice Memory Impairment					
200MG per day		Pancreatitis Panic Reaction		Simvastatin Seloken	C C		
10MG per day		Peritonitis		Renitec	C		
40MG per day		Purulence		Frusemide	C	Glaxo Wellcome	
75MG per day		Sexual Dysfunction Sleep Disorder		Aspirin	C		

Date:09/13/02ISR Number: 3974845-6Report Type:Expedited (15-DaCompany Report #B0271683A  
Age:49 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day	4 DAY	Ageusia		Zyban	PS	Glaxo Wellcome	ORAL
50MG Unknown			Anosmia		Diclofenac	C		ORAL
2TABS Unknown			Anxiety		Codrydamol	C	Glaxo Wellcome	ORAL
			Dyspnoea Exertional Tachycardia		Prempak C	C		ORAL

Date:09/16/02ISR Number: 3975499-5Report Type:Expedited (15-DaCompany Report #B0134762A  
Age:49 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Unknown		Mydriasis		Zyban	PS	Glaxo Wellcome	ORAL
			Sudden Death					



Outcome	PT
Hospitalization -	Aspartate
Initial or Prolonged	Aminotransferase
	Increased
	Blood Albumin Decreased
	Blood Potassium Decreased
	Blood Sodium Decreased
	Blood Urea Decreased
	C-Reactive Protein
	Increased
	Eosinophilia
	Gamma-Glutamyltransferase
	Increased
	Heart Rate Increased
	Hyperglycaemia
	Lymphocyte Count
	Decreased
	Neutrophil Count
	Increased
	Proteinuria
	Pyrexia
	Rash Generalised
	Rash Maculo-Papular
	Rash Pruritic
	Rosacea
	Toxic Skin Eruption

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	White Blood Cell Count Increased	Report Source	Product	Role	Manufacturer	Route
11 DAY				Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice				Mediator	SS		ORAL
per day							

Date:09/16/02ISR Number: 3975502-2Report Type:Expedited (15-DaCompany Report #B0276547A  
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	14 DAY			Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	50MG per day	Suicide Attempt		Zoloft	SS		ORAL
4MG per day				Lexomil	C		ORAL

Date:09/16/02ISR Number: 3975504-6Report Type:Expedited (15-DaCompany Report #B0277431A  
 Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day			Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -	Initial or Prolonged	Dyspnoea Hypernatraemia Hypoxia Loss Of Consciousness					

Date:09/16/02ISR Number: 3975505-8Report Type:Expedited (15-DaCompany Report #B0278802A  
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization -	Anxiety	Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day 26 DAY					
Initial or Prolonged	Arrhythmia	Kardegic	C		ORAL
	Blood Creatine	Lasilix	C	Glaxo Wellcome	ORAL
	Phosphokinase Increased	Triatec	C		ORAL
	Blood Lactate				
	Dehydrogenase Increased				
	Disorientation				
	Hallucination				
	Hepatic Enzyme Increased				
	Hyponatraemia				
	Malaise				

Date:09/16/02ISR Number: 3975510-1Report Type:Expedited (15-DaCompany Report #B0279427A  
Age:41 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Chest Discomfort		Zyban	PS	Glaxo Wellcome	ORAL
150MG As						
Initial or Prolonged	Dizziness					
directed 10 DAY						
Other	Fatigue					
	Panic Reaction					

Date:09/16/02ISR Number: 3976290-6Report Type:Direct Company Report #CTU 176549  
Age:37 YR Gender:Female I/FU:I

Outcome	PT
	Abnormal Dreams
	Dry Mouth

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Headache Insomnia Tremor	Report Source	Product	Role	Manufacturer	Route
150 MG ONE PO				Zyban 150mg	PS		ORAL
QD X 3 D, THEN BID				Habitrol	C		

Date:09/17/02ISR Number: 3976078-6Report Type:Expedited (15-DaCompany Report #A0371353A  
Age:32 YR Gender:Female I/FU:F

Outcome Dose Other 100MG Twice per day	Duration 4 DAY	PT Agitation Anxiety Depression Euphoric Mood Feeling Abnormal Insomnia Malaise Panic Reaction Performance Status Decreased	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin	PS	Glaxo Wellcome	ORAL
				Ethanol	C		ORAL

Date:09/17/02ISR Number: 3976079-8Report Type:Expedited (15-DaCompany Report #A0378334A  
Age:14 YR Gender:Male I/FU:F

Outcome Dose Hospitalization - 150MG Twice Initial or Prolonged per day Other 1 YR	Duration 18 DAY	PT Serum Sickness	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin	PS	Glaxo Wellcome	ORAL
				Concerta	C		

Date:09/17/02ISR Number: 3976087-7Report Type:Expedited (15-DaCompany Report #B0118244A  
Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG Twice		Abdominal Pain Upper Angina Unstable		Bupropion Hydrochloride	PS	Glaxo Wellcome	ORAL
per day	11	DAY	Cardiac Failure Congestive Chest Discomfort Chest Pain Chronic Obstructive Pulmonary Disease Circulatory Collapse Dyspepsia Dyspnoea Dyspnoea Exertional Heart Rate Increased Pharyngolaryngeal Pain				

Date:09/17/02ISR Number: 3976088-9Report Type:Expedited (15-DaCompany Report #B0134124A  
Age:34 YR Gender:Male I/FU:F

Outcome	PT
Disability	Angioneurotic Oedema Arthralgia Erythema Nodosum

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Oedema Proteinuria Pruritus	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Zyban	PS	Glaxo Wellcome	ORAL
9	DAY						

Date:09/17/02ISR Number: 3976089-0Report Type:Expedited (15-DaCompany Report #B0260806A  
Age:50 YR Gender:Male I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Zyban	PS	Glaxo Wellcome	ORAL
Other		Lichen Planus					
150MG Per day	6 DAY						

Date:09/18/02ISR Number: 3976840-XReport Type:Expedited (15-DaCompany Report #B0267494A  
Age:56 YR Gender:Male I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Zyban	PS	Glaxo Wellcome	
Other		Hypomania					
UNKNOWN	150MG	Unknown		Blood Pressure Medication	C		
		Psychotic Disorder					

Date:09/18/02ISR Number: 3976841-1Report Type:Expedited (15-DaCompany Report #B0272342A  
Age:31 YR Gender:Male I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -		Arthralgia	Health				
3 WK							
Initial or Prolonged		Hypokalaemia Hyponatraemia Insomnia Joint Effusion Renal Colic Tremor	Professional				

Date:09/18/02ISR Number: 3976842-3Report Type:Expedited (15-DaCompany Report #B0272924A  
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	9 DAY	Aggression	Health	Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - 1TAB As Initial or Prolonged required		Agitation	Professional	Apranax	C		ORAL
1.5G Twice per day	9 DAY	Condition Aggravated		Magnesium Pidolate	C		ORAL
		Impulsive Behaviour					
		Orthostatic Hypotension					
200MG per day	9 DAY	Pyromania		Tiapride	C		ORAL
1MG per day	9 DAY			Xanax	C		ORAL

Date:09/18/02ISR Number: 3976846-0Report Type:Expedited (15-DaCompany Report #B0278546A  
Age:58 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	13 DAY	Conjunctival Hyperaemia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown				Simvastatin	C		ORAL
20MG per day				Telmisartan	C	Glaxo Wellcome	ORAL
40MG per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/18/02ISR Number: 3976847-2Report Type:Expedited (15-DaCompany Report #B0278942A

Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	10 DAY	Coma		Zyban	PS	Glaxo Wellcome	ORAL
		Convulsion Cyanosis Dyspnoea Excitability Sudden Death		Oral Contraceptive Pill	C		ORAL

Date:09/18/02ISR Number: 3977437-8Report Type:Direct Company Report #CTU 176745

Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	1T PO QD X 3D , THEN BID 1T Q H UP TO 8 PIECES/D	Abnormal Dreams Dry Mouth Face Oedema Pharmaceutical Product Complaint		Zyban 150mg Tbsr Nicorette 2mg Gum	PS SS		ORAL

Date:09/19/02ISR Number: 3977493-7Report Type:Expedited (15-DaCompany Report #A0373936A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	450MG Unknown	Convulsion		Wellbutrin Sr Unknown	PS C	Glaxo Wellcome	ORAL

Date:09/19/02ISR Number: 3978188-6Report Type:Expedited (15-DaCompany Report #A0380102A

Age:42 YR Gender: I/FU:I



Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Drug Level Increased Intentional Misuse Suicide Attempt	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL					Carbamazepine (Carbamazepine) Ethanol (Alcohol)	SS SS		

Date:09/19/02ISR Number: 3978191-6Report Type:Expedited (15-DaCompany Report #A0380101A  
Age:45 YR Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Drug Level Increased Intentional Misuse Suicide Attempt	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL								

Date:09/19/02ISR Number: 3978293-4Report Type:Expedited (15-DaCompany Report #A0379982A  
Age:36 YR Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL								

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

				Clonazepam (Clonazepam)	SS		
				Trazodone (Trazodone)	SS		

Date:09/19/02ISR Number: 3978294-6Report Type:Expedited (15-DaCompany Report #A0379981A  
Age:37 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Carbamazepine (Carbamazepine) Olanzapine (Olanzapine)	SS SS		

Date:09/19/02ISR Number: 3978296-XReport Type:Expedited (15-DaCompany Report #A0379980A  
Age:36 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL				Hydrocodone + Paracetamol (Hydrocodone + Acetaminoph)	SS		ORAL
ORAL				Carisoprodol (Carisoprodol)	SS		

Date:09/19/02ISR Number: 3978297-1Report Type:Expedited (15-DaCompany Report #A0379979A  
Age:41 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Date:09/19/02ISR Number: 3978298-3Report Type:Expedited (15-DaCompany Report #A0379918A  
Age:17 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/19/02ISR Number: 3978454-4Report Type:Expedited (15-DaCompany Report #A0379990A  
 Age:46 YR Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL					Alprazolam (Alprazolam)	SS		
					Olanzapine (Olanzapine)	SS		

Date:09/19/02ISR Number: 3978455-6Report Type:Expedited (15-DaCompany Report #A0379989A  
 Age:16 YR Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL					Quetiapine (Quetiapine)	SS		
					Valproic Acid (Valproic Acid)	SS		

Date:09/19/02ISR Number: 3978458-1Report Type:Expedited (15-DaCompany Report #A0379988A  
 Age:44 YR Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Death	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL					Fluoxetine (Fluoxetine)	SS		
					Hydrocodone +			

Paracetamol  
(Formulation  
Unknown)  
(Hydrocodone + SS

Date:09/19/02ISR Number: 3978460-XReport Type:Expedited (15-DaCompany Report #A0379987A

Age:35 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL				Venlafaxine Hydrochloride (Formulation Unknown) (Venlafaxine Methylphenidate (Formulation Unknown) (Methylphenidate)	SS  SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/19/02ISR Number: 3978463-5Report Type:Expedited (15-DaCompany Report #A0379986A

Age:21 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Olanzapine (Formulation Unknown) (Olanzapine)	SS		
				Clonazepam (Formulation Unknown) (Clonazepam)	SS		

Date:09/19/02ISR Number: 3978465-9Report Type:Expedited (15-DaCompany Report #A0379985A

Age:46 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Abuser Intentional Misuse	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL				Naproxen (Formulation Unknown) (Naproxen)	SS		ORAL
ORAL				Cocaine (Formulation Unknown) (Cocaine)	SS		

Date:09/19/02ISR Number: 3978466-0Report Type:Expedited (15-DaCompany Report #A0379984A

Age:36 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health	Wellbutrin Unspecified Tablet			

ORAL			Professional	(Bupropion Hydrochloride)	PS		ORAL
ORAL				Ethylene Glycol (Formulation Unknown) (Ethylene Glycol)	SS		ORAL
ORAL				Hypericum (Formulation Unknown) (Hypericum)	SS		ORAL

Date:09/19/02ISR Number: 3978469-6Report Type:Expedited (15-DaCompany Report #A0379983A  
Age:51 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL				Ethanol (Formulation Unknown) (Alcohol)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/19/02ISR Number: 4008125-XReport Type:Periodic  
 Age:74 YR Gender:Female I/FU:I

Company Report #S02-USA-00997-01

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 40 MG QD PO	Stupor	Health Professional	Celexa (Citalopram Hydrobromide)	PS		ORAL
40 MG QD PO			Celexa (Citalopram Hydrobromide)	SS		ORAL
100 MG BID			Wellbutrin (Amfebutamone Hydrochloride)	SS		
			Pamelor (Nortriptyline Hydrochloride)	C		

Date:09/20/02ISR Number: 3977925-4Report Type:Expedited (15-DaCompany Report #A0380991A  
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Anxiety Diarrhoea Headache Hyperhidrosis Impaired Work Ability Vomiting Weight Decreased		Zyban	PS	Glaxo Wellcome	ORAL

Date:09/20/02ISR Number: 3977926-6Report Type:Expedited (15-DaCompany Report #A0381096A  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus		Wellbutrin	PS	Glaxo Wellcome	



Date:09/20/02ISR Number: 3977939-4Report Type:Expedited (15-DaCompany Report #B0279967A

Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hemiparesis		Zyban	PS	Glaxo Wellcome	
UNKNOWN							

Date:09/24/02ISR Number: 3980442-9Report Type:Expedited (15-DaCompany Report #DSA\_21894\_2002

Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Blood Glucose Increased	Consumer	Cardizem Cd	PS		ORAL
360 MG DAILY							
Initial or Prolonged		Blood Pressure Decreased					
PO							
		Blood Pressure Increased		Zyban Tablet			
		Cystitis		(Bupropion Hcl)	SS		ORAL
150 MG DAILY							
		Drug Interaction					
PO							
				Atenolol	SS		ORAL
50 MG DAILY							
PO							
				Glucophage Xr			
				(Metformin)	C		
				Altace	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

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Date:09/25/02ISR Number: 3981329-8Report Type:Direct  
Age:40 YR Gender:Male I/FU:I

Company Report #CTU 177114

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG 1 PO Initial or Prolonged BID ORAL Required Intervention to Prevent Permanent Impairment/Damage		Dizziness		Bupropion	PS		ORAL

Date:09/25/02ISR Number: 3982560-8Report Type:Expedited (15-DaCompany Report #NSADSS2002032610  
Age:33 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death ORAL		Death	Literature Health Professional	Risperdal (Risperidone) Bupropion (Amfebutamone)	PS SS		ORAL

Date:09/26/02ISR Number: 3981983-0Report Type:Direct  
Age:51 YR Gender:Male I/FU:I

Company Report #CTU 177337

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG QD ORAL Initial or Prolonged		Chest Pain		Bupropion 150mg	PS		ORAL

Date:09/27/02ISR Number: 3982877-7Report Type:Expedited (15-DaCompany Report #NSADSS2002032843  
Age:42 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Toxicologic Test Abnormal	Literature Health Professional	Concerta (Sustained Release Tablet) (Methylphenidate Hydrochloride) Bupropion(Amfebutamo ne)	PS SS		

Date:09/27/02ISR Number: 3982889-3Report Type:Expedited (15-DaCompany Report #NSADSS2002032842  
 Age:42 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Literature Health Professional	Concerta (Sustained Release Tablet) (Methylphenidate Hydrochloride) Bupropion	PS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/27/02ISR Number: 3982959-XReport Type:Direct  
Age:73 YR Gender:Male I/FU:I

Company Report #CTU 177501

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - INTRAVENOUS Initial or Prolonged 1MG Q4H PRN	Neuroleptic Malignant Syndrome		Haloperidol	PS		BOLUS
INTRAVENOUS  BO 100 MG QD			Bupropion Sr 100 Mg	SS		ORAL
ORAL			Seroquel	C		

Date:09/27/02ISR Number: 3983538-0Report Type:Expedited (15-DaCompany Report #B0129833A  
Age:42 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG/PER DAY/ORAL	Arthralgia C-Reactive Protein Increased  Eyelid Oedema  Face Oedema Pyrexia Sigmoiditis White Blood Cell Count White Blood Cell Count Increased	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
			Amisulpride	C		

Date:09/27/02ISR Number: 3983543-4Report Type:Expedited (15-DaCompany Report #B0280402A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization -	Latent Tetany	Foreign	Zyban Tablet - Zyban			

Initial or Prolonged

Health  
Professional

(Bupropion  
Hydrochloride)

PS

ORAL

ORAL

Company  
Representative

Date:09/27/02ISR Number: 3983544-6Report Type:Expedited (15-DaCompany Report #B0280371A

Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Confusional State Depression Haematoma	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/PER DAY/ ORAL		Irritability Tremor Vomiting					

Date:09/27/02ISR Number: 3983587-2Report Type:Expedited (15-DaCompany Report #B0280370A

Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Distension Intestinal Hypomotility	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/27/02ISR Number: 3983652-XReport Type:Expedited (15-DaCompany Report #A0380888A  
 Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Blood Potassium Abnormal Blood Sodium Abnormal Drug Withdrawal Syndrome	Foreign Consumer	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL  30 MG PER  DAY/ORAL	Hyperglycaemia Hypoglycaemia		Paxil (Paroxetine Hydrochloride)	SS		ORAL

Date:09/27/02ISR Number: 3984179-1Report Type:Expedited (15-DaCompany Report #B0259361A  
 Age:46 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Psoriasis	Foreign Health Professional	Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE  PER DAY/ORAL						

Date:09/27/02ISR Number: 3984181-XReport Type:Expedited (15-DaCompany Report #B0270415A  
 Age:32 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Required Intervention to 150 MG/AS Prevent Permanent DIRECTED Impairment/Damage	Injury Intentional Misuse Psychotic Disorder  Self Mutilation  Wound	Foreign Health Professional	Zyban (Bupropion Hydrochloride)	PS		
			Oxazepam Mitrazapine	C C		

Date:09/27/02ISR Number: 3984182-1Report Type:Expedited (15-DaCompany Report #B0271008A  
Age:44 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Required SEE DOSAGE Intervention to TEXT / ORAL Prevent Permanent Impairment/Damage	Blood Immunoglobulin G Decreased Hepatic Steatosis Serum Sickness Tooth Infection	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)  Nicotine Polacrilex	PS  C		ORAL

Date:09/27/02ISR Number: 3984828-8Report Type:Expedited (15-DaCompany Report #A0377288A  
Age:29 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Required Intervention to Prevent Permanent 150 MG/SEE Impairment/Damage DOSAGE TEXT/  INTRANASA	Anxiety Chest Discomfort Drug Abuser Medication Error Nasal Discomfort Tremor	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		NASAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/27/02ISR Number: 3984837-9Report Type:Expedited (15-DaCompany Report #A0380744A  
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blindness Unilateral Cholelithiasis Hypoaesthesia	Consumer	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
150 MG / THREE TIMES PER DAY / ORAL	14 YR			Thyroxine Sodium Fioricet Castor Oil Somatropin	C C C C		

Date:09/27/02ISR Number: 3985313-XReport Type:Expedited (15-DaCompany Report #A0173197A  
Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Circulatory Collapse Myocardial Infarction Nausea Urinary Incontinence	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG TWICE PER DAY ORAL		Vomiting		Entex Pse Hydrocodone + Paracetamol Amox. Trihyd+Pot.Clavulan Prednisone	C C C C		

Date:09/27/02ISR Number: 3985801-6Report Type:Expedited (15-DaCompany Report #A103444  
Age:79 YR Gender:Female I/FU:F

Outcome PT



Required	Affective Disorder
Intervention to	Anger
Prevent Permanent	Arrhythmia
Impairment/Damage	Autoimmune Thyroiditis
	Balance Disorder
	Blood Oestrogen Decreased
	Blood Pressure Diastolic Decreased
	Blood Pressure Increased
	Blood Thyroid Stimulating Hormone Decreased
	Contusion
	Cough
	Dementia
	Depression
	Drug Ineffective
	Dyssomnia
	Epistaxis
	Excoriation
	Fall
	Haemorrhagic Stroke
	Hallucination
	Heart Rate Increased
	Hostility
	Meningioma

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Abnormal	Report Source	Product	Role	Manufacturer	Route
50 MG	(DAILY), ORAL	Nuclear Magnetic Resonance Imaging Abnormal Overdose	Consumer	Zoloft (Sertraline)	PS		ORAL
300 MG (BID), ORAL		Pain Palpitations Parkinson'S Disease Prothrombin Time Prolonged		Wellbutrin Sr (Bupropion Hydrochloride)	SS		ORAL
300 MG (BID), ORAL		Rib Fracture Syncope Thinking Abnormal Thyroid Disorder		Coumadin (Warfarin) Effexor Xr (Venlafaxine Hydrochloride)	SS SS		ORAL
		Transient Ischaemic Attack Treatment Noncompliance Vaginal Haemorrhage		Amantadine Monopril (Fosinopril Sodium) Donepezil Hydrochloride Zebeta (Bisoprolol Fumarate) Lanoxin (Digoxin) Lasix (Furosemide) Synthroid (Levothyroxine Sodium) Vitamin E (Tocopherol) Estrogen (Estrogen Nos) Zocor (Simvastatin) Estradiol Vaginal Ring (Estradiol) Inderal (Propranolol Hydrochloride) Cardizem (Diltiazem Hydrochloride) Folic Acid Sinemet (Carbidopa / Levodopa)	SS SS C		

"Carzon" C  
Metoprolol Succinate C

Date:09/30/02ISR Number: 3983757-3Report Type:Direct Company Report #CTU 177601  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Grand Mal Convulsion		Wellbutrin	PS		
200 MG SR BID							
Initial or Prolonged		Pneumonia Aspiration					
X 3 MO PTA							

Date:10/01/02ISR Number: 4008711-7Report Type:Periodic Company Report #2001AP05211  
Age:16 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Quetiapine	PS		
			Health	Valproic Acid	SS		
			Professional	Bupropion	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/02/02ISR Number: 3986583-4Report Type:Expedited (15-DaCompany Report #B0280644A

Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Brain Neoplasm Cerebellar Syndrome Concomitant Disease	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL		Progression Langerhans' Cell Granulomatosis Vertigo					

Date:10/02/02ISR Number: 3987190-XReport Type:Expedited (15-DaCompany Report #A0381384A

Age:16 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Grand Mal Convulsion Intentional Misuse Medication Error	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		
RESPIRATORY (INHALATION)	900 MG/ SINGLE DOSE/ INHALED			Ethanol Cannabis	C C		

Date:10/02/02ISR Number: 3987193-5Report Type:Expedited (15-DaCompany Report #A0381535A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abnormal Behaviour Hallucination, Auditory Impulsive Behaviour Mania	Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ORAL							

Paranoia  
Personality Change  
Psychotic Disorder  
Self Injurious Behaviour  
Thinking Abnormal  
Tinnitus

Date:10/02/02ISR Number: 3987466-6Report Type:Expedited (15-DaCompany Report #A0377706A

Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Encephalopathy Hypothyroidism Metabolic Disorder	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG TWICE							
PER DAY ORAL				Oral Contraceptive Zolpidem Tartrate Quetiapine Fumarate	C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/02/02ISR Number: 3987469-1Report Type:Expedited (15-DaCompany Report #A0379580A  
 Age:59 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Bronchitis Hypotension Pneumonia Haemophilus	Health Professional Company	Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL	Renal Failure Stevens-Johnson Syndrome	Representative	Semisodium Valproate Lithium Carbonate Zolmitriptan Vicodin	C C C C		

Date:10/02/02ISR Number: 3987471-XReport Type:Expedited (15-DaCompany Report #A0380824A  
 Age:32 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Drug Hypersensitivity Dysphagia Oesophageal Stenosis	Consumer	Wellbutrin Sr Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG PER DAY ORAL			Oral Contraceptive Vitamin	C C		

Date:10/02/02ISR Number: 3987982-7Report Type:Expedited (15-DaCompany Report #B0276132A  
 Age:45 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Aspartate Aminotransferase Increased	Foreign	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL
SEE DOSAGE TEXT / ORAL	Blood Albumin Decreased Blood Glucose Increased Blood Potassium Decreased Blood Sodium Decreased		Benfluorex Hydrochloride Tablet (Benfluorex			

150 MG /

TWICE PER DAY

/ ORAL

Blood Sodium Increased

Blood Urea Increased

C-Reactive Protein

Increased

Dermatitis Exfoliative

Dyskinesia

Eosinophilia

Exanthem

Gamma-Glutamyltransferase

Increased

Lymphocyte Count

Decreased

Neutrophil Count

Increased

Proteinuria

Rash

Rash Maculo-Papular

Rosacea

Toxic Skin Eruption

White Blood Cell Count

Increased

Hydrochloride)

SS

ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/03/02ISR Number: 3985628-5Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 177878

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams		Wellbutrin	PS		
150 MGM BID							
		Blood Pressure Systolic		Erythromycin	SS		
333 MGM TID							
		Increased		Ibuprofen	C		
		Headache		Trilevelon	C		
		Tachycardia					
		Tremor					

Date:10/03/02ISR Number: 3989056-8Report Type:Expedited (15-DaCompany Report #A103444  
Age:79 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Anger	Consumer	Zoloft (Sertraline)	PS		ORAL
50 MG							
Intervention to		Arrhythmia					
(DAILY), ORAL							
Prevent Permanent		Balance Disorder		Wellbutrin Sr			
Impairment/Damage		Blood Oestrogen Decreased		(Bupropion			
		Blood Pressure Increased		Hydrochloride)	SS		ORAL
300 MG (BID),							
ORAL		Blood Thyroid Stimulating					
		Hormone Decreased		Coumadin (Warfarin)	SS		
		Cerebral Haemorrhage		Effexor Xr			
		Cognitive Disorder		(Venlafaxine			
		Contusion		Hydrochloride)	SS		ORAL
300 MG (BID),							
ORAL		Cough					
		Depression		Amantadine	SS		
		Drooling		Monopril (Fosinopril			
		Drug Ineffective		Sodium)	SS		
		Epistaxis		Donepezil			
		Fall		Hydrochloride	C		
		Hallucination		Zebeta (Bisoprolol			
		Head Injury		Fumarate)	C		
		Heart Rate Decreased		Lanoxin (Digoxin)	C		
		Heart Rate Increased		Lasix (Furosemide)	C		



Hostility	Synthroid	
Hyperhidrosis	(Levothyroxine	
Medication Error	Sodium)	C
Memory Impairment	Vitamin E	
Meningioma	(Tocopherol)	C
Overdose	Estrogen (Estrogen	
Pain	Nos)	C
Prothrombin Time Abnormal	Zocor (Simvastatin)	C
Pulmonary Oedema	Estradiol Vaginal	
Rib Fracture	Ring (Estradiol)	C
Sedation	Inderal (Propranolol	
Sleep Talking	Hydrochloride)	C
Syncope	Cardizem (Diltiazem	
Transient Ischaemic	Hydrochloride)	C
Attack	Folic Acid	C
Treatment Noncompliance	Sinemet (Carbidopa /	
Tremor	Levodopa)	C
Vaginal Haemorrhage	"Carzon"	C
	Metoprolol Succinate	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/04/02ISR Number: 3988045-7Report Type:Expedited (15-DaCompany Report #B0133703A  
 Age:52 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/PER DAY/ORAL	Angina Unstable Chest Pain Coronary Artery Stenosis	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Omega-3 Marine Triglyceri Simvastatin	C C	
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Date:10/04/02ISR Number: 3988287-0Report Type:Expedited (15-DaCompany Report #A0377881A  
 Age:55 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 4 MG/ TWELVE TIMES PER DAY / ORAL ORAL	Alopecia Chronic Obstructive Pulmonary Disease Emphysema Hyperphagia Medication Error Weight Increased	Consumer	Nicorette Otc Gum (Nicotine Polacrilex)	PS		ORAL
			Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	SS		ORAL

Date:10/04/02ISR Number: 3988345-0Report Type:Expedited (15-DaCompany Report #A0345310A  
 Age:50 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other ORAL	Blood Creatinine Increased Bradyphrenia	Foreign Health Professional	Paxil Tablet (Paroxetine Hydrochloride)	PS		ORAL

Required	Confusional State	Wellbutrin Sr		
Intervention to	Depression	Tablet-Controlled		
Prevent Permanent	Disorientation	Release (Bupropion		
Impairment/Damage	Drug Interaction	Hydrochloride)	SS	ORAL
250 MG / PER				
	Fall			
DAY / ORAL				
	Grand Mal Convulsion	Fluoxetine		
	Hallucination	(Fluoxetine)	SS	ORAL
20 MG / PER				
	Memory Impairment			
DAY / ORAL				
	Movement Disorder	Clonazepam	C	
	Rhabdomyolysis	Oxazepam	C	
	Speech Disorder			
	Thinking Abnormal			

Date:10/07/02ISR Number: 3984417-5Report Type:Expedited (15-DaCompany Report #B0131464A  
Age:25 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Arteriospasm Coronary
Initial or Prolonged	Blood Pressure Decreased
	Cardiac Failure
	Chest Pain
	Cutaneous Vasculitis
	Cytomegalovirus Infection
	Diarrhoea
	Disseminated
	Intravascular Coagulation



Death Depression Consumer Zyban PS Glaxo Wellcome  
UNKNOWN

Sudden Death

Date:10/07/02ISR Number: 3984770-2Report Type:Expedited (15-DaCompany Report #B0268687A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Zyban	PS	Glaxo Wellcome	ORAL
		Fall		Contraceptive			
		Loss Of Consciousness		(Unspecified)	C		
		Malaise					

Date:10/07/02ISR Number: 3987115-7Report Type:Expedited (15-DaCompany Report #2002054675  
Age:35 YR Gender:Male I/FU:F

Outcome	PT
Other	Convulsion
	Drug Level Decreased
	Hemiplegia
	Migraine
	Muscle Twitching

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Paranoia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300 MG (TID), ORAL		Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Bupropion Hydrochloride Citalopram Hydrobromide	SS C		

Date:10/07/02ISR Number: 3987131-5Report Type:Direct  
Age:45 YR Gender:Female I/FU:I Company Report #CTU 178139

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other PO		Rash Erythematous Rash Pruritic		Wellbutrin - 150mg Bid	PS		ORAL
PO				Lipitor -10mg	SS		ORAL

Date:10/07/02ISR Number: 3987300-4Report Type:Expedited (15-DaCompany Report #200219747US  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 180 MG; PO	20 DAY	Agitation Condition Aggravated Convulsion	Consumer	Fexofenadine Hydrochloride (Allegra) Tablets	PS		ORAL
300 MG QD; PO		Difficulty In Walking Insomnia Rash		Amfebutamone Hydrochloride (Wellbutrin Sr)	SS		ORAL
200 MG BID;		Speech Disorder Tongue Spasm Tremor		Amfebutamone Hydrochloride (Wellbutrin Sr)	SS		ORAL

Cefuroxime Axetil  
(Ceftin) C  
Gabapentin  
(Neurontin) C

Date:10/07/02ISR Number: 3990440-7Report Type:Direct  
Age:17 YR Gender:Female I/FU:I

Company Report #CTU 178101

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 200MG BID PO Intervention to Prevent Permanent Impairment/Damage		Grand Mal Convulsion		Wellbutrin Sr 100mg	PS		ORAL
				Celexa	C		
				Zyprexa	C		

Date:10/08/02ISR Number: 3987206-0Report Type:Expedited (15-DaCompany Report #02P-163-0201368-00  
Age:16 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Completed Suicide	Literature Health Professional	Valproic Acid (Depakene) (Valproic Acid) (Valproic Acid)	PS		
				Quetiapine	SS		
				Bupropion	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/08/02ISR Number: 3987364-8Report Type:Expedited (15-DaCompany Report #A0372870A

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other Required Intervention to Prevent Permanent 150 MG / Impairment/Damage THREE TIMES  PER DAY /  ORAL  150 MG / PER  DAY		Grand Mal Convulsion	Health Professional	Wellbutrin Sr  (Bupropion Hydrochloride)  Ludiomil (Formulation Unknown) (Ludiomil)	PS		ORAL
				Acarbose Metformin Hydrochloride Thyroid Celecoxib Omeprazole Risperidone Doxepin Alprazolam Vitamin D Calcium Salt Glucosamine Chondroitin C	C C C C C C C C C C C		

Date:10/08/02ISR Number: 3987378-8Report Type:Expedited (15-DaCompany Report #A0382184A

Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Benign Intracranial Hypertension	Health Professional	Wellbutrin Sr  (Bupropion			



400 MG / PER

Hydrochloride)

PS

ORAL

DAY / ORAL

Date:10/08/02ISR Number: 3989761-3Report Type:Expedited (15-DaCompany Report #02P-163-0201358-00  
Age:36 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Hydrocodone/Acetaminophen (Vicodin)(Hydrocodone/Acetaminophen) Bupropion Carisoprodol	PS SS SS		

Date:10/08/02ISR Number: 3990027-6Report Type:Expedited (15-DaCompany Report #B0281102A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL				Nicotine Substitute	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/08/02ISR Number: 3990048-3Report Type:Expedited (15-DaCompany Report #B0118244A  
Age:63 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/ TWICE PER DAY/ ORAL	Abdominal Pain Upper Aortic Valve Stenosis Blood Pressure Systolic Increased Cardiac Failure Congestive Chronic Obstructive Pulmonary Disease Dilatation Ventricular Dyspepsia Pharyngolaryngeal Pain Ventricular Hypokinesia	Foreign Study Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:10/08/02ISR Number: 3990062-8Report Type:Expedited (15-DaCompany Report #B0274449A  
Age:72 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Hospitalization - Initial or Prolonged SEE DOSAGE TEXT / ORAL	Insomnia Respiratory Failure	Foreign Health Professional Company Representative	Zyban Tablet - Zyban (Bupropion Hydrochloride) Ace Inhibitor	PS C		ORAL

Date:10/08/02ISR Number: 3990073-2Report Type:Expedited (15-DaCompany Report #B0274795A  
Age:30 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Required 150 MG/	Convulsion Rheumatoid Arthritis	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

Intervention to  
VARIABLE  
Prevent Permanent  
DOSE/ORAL  
Impairment/Damage

Nimesulide C

Date:10/08/02ISR Number: 3990074-4Report Type:Expedited (15-DaCompany Report #B0269955A  
Age:50 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to Prevent Permanent 150 MG / PER Impairment/Damage DAY / ORAL	Myalgia Polyarthrititis	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:10/08/02ISR Number: 3990087-2Report Type:Expedited (15-DaCompany Report #B0264074A  
Age:54 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - Initial or Prolonged 150 MG/PER DAY/ ORAL	Angina Unstable Myocardial Infarction	Foreign Health Professional  Company  Representative	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
			Rilmenidine	C		
			Fenofibrate	C		
			Piascledine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Zolpidem C

Date:10/08/02ISR Number: 3990090-2Report Type:Expedited (15-DaCompany Report #B0134232A  
Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ ORAL				Paroxetine Hydrochloride	C		
				Diltiazem Hydrochloride	C		
				Oxygen	C		

Date:10/09/02ISR Number: 3986359-8Report Type:Expedited (15-DaCompany Report #A0380744A  
Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blindness Unilateral		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Three times per day 14 YR		Cholelithiasis		Levothroid	C	Glaxo Wellcome	
		Hypoaesthesia		Fioricet	C		
				Castor Oil	C		
				Human Growth Hormone	C		

Date:10/09/02ISR Number: 3986363-XReport Type:Expedited (15-DaCompany Report #A0382619A  
Age:9 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG Per day 22 DAY Initial or Prolonged		Serum Sickness		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Stevens-Johnson Syndrome					

Date:10/09/02ISR Number: 3986366-5Report Type:Expedited (15-DaCompany Report #B0261797A  
Age:34 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150MG See Hospitalization - dosage text Initial or Prolonged UNKNOWN	Diplopia Epilepsy Loss Of Consciousness Status Epilepticus		Zyban Vitamins Ephedrine Guarana Neuroleptic	PS SS SS	Glaxo Wellcome	ORAL ORAL ORAL
UNKNOWN			Antidepressant (Unspecified)	C		
UNKNOWN			Benzodiazepines	C		

Date:10/09/02ISR Number: 3986371-9Report Type:Expedited (15-DaCompany Report #B0281345A  
Age:31 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150MG Unknown 8 DAY	Aggression Panic Reaction Paranoia		Zyban	PS	Glaxo Wellcome	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/09/02ISR Number: 3986372-0Report Type:Expedited (15-DaCompany Report #B0281421A  
Age:32 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150MG Per day 7 DAY	Muscle Strain		Zyban	PS	Glaxo Wellcome	ORAL

Date:10/09/02ISR Number: 3986373-2Report Type:Expedited (15-DaCompany Report #B0281501A  
Age:32 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG per day 27 DAY Initial or Prolonged	Arthralgia Bone Pain Inflammation Pyrexia		Zyban	PS	Glaxo Wellcome	ORAL

Date:10/09/02ISR Number: 3986418-XReport Type:Periodic Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-11018074  
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration	Alopecia		Buspar Tabs Wellbutrin	PS SS	Bristol-Myers Squibb Company	ORAL

Date:10/09/02ISR Number: 3990863-6Report Type:Expedited (15-DaCompany Report #A0379437A  
Age: Gender:Unknown I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG / TWICE PER DAY	Retinal Detachment Vitreous Floaters	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

/ ORAL

Adderall

C

Date:10/10/02ISR Number: 3987292-8Report Type:Expedited (15-DaCompany Report #A0382701A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus		Bupropion	PS	Glaxo Wellcome	ORAL

Date:10/10/02ISR Number: 3987293-XReport Type:Expedited (15-DaCompany Report #A0382751A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG Per day 1 WK		Cardiac Arrest		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged Other		Ventricular Fibrillation					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/10/02ISR Number: 3987294-1Report Type:Expedited (15-DaCompany Report #B0269609A  
 Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complex Partial Seizures	Health	Zyban	PS	Glaxo Wellcome	
UNKNOWN		3 WK					
		Malaise	Professional				

Date:10/10/02ISR Number: 3991105-8Report Type:Direct Company Report #CTU 178512  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Flat Affect		Zyban 150	PS		
BID							

Date:10/10/02ISR Number: 3991108-3Report Type:Direct Company Report #CTU 178515  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Insomnia		Wellbutrin 150mg	PS		
BID		Tremor					

Date:10/10/02ISR Number: 3991109-5Report Type:Direct Company Report #CTU 178516  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Nightmare		Wellbutrin 150mg	PS		
BID							

Date:10/10/02ISR Number: 3991110-1Report Type:Direct Company Report #CTU 178517  
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Rash Wellbutrin 150 PS  
 BID  
 Date:10/10/02ISR Number: 3991111-3Report Type:Direct Company Report #CTU 178518  
 Age:48 YR Gender:Female I/FU:I  
 Outcome PT Report Source Product Role Manufacturer Route  
 Dose Duration Palpitations Wellbutrin 150mg PS  
 BID

Date:10/10/02ISR Number: 3991113-7Report Type:Direct Company Report #CTU 178520  
 Age:27 YR Gender:Female I/FU:I  
 Outcome PT Report Source Product Role Manufacturer Route  
 Dose Duration Urticaria Wellbutrin PS  
 Other

Date:10/11/02ISR Number: 3988128-1Report Type:Expedited (15-DaCompany Report #A0382851A  
 Age:36 YR Gender:Male I/FU:F  
 Outcome PT Report Source Product Role Manufacturer Route  
 Dose Duration Hospitalization - Drug Abuser Wellbutrin PS Glaxo Wellcome  
 INTRAVENOUS  
 Initial or Prolonged Medication Error

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/14/02ISR Number: 3989442-6Report Type:Expedited (15-DaCompany Report #B0281921A  
 Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	18 DAY	Cyanosis Discomfort Influenza Like Illness Oedema Physical Disability Urticaria		Zyban  Salatel	PS  C	Glaxo Wellcome	ORAL

Date:10/14/02ISR Number: 3989444-XReport Type:Expedited (15-DaCompany Report #B0282033A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Atrial Fibrillation Hyperthyroidism Insomnia Nervousness Palpitations Tremor		Zyntabac	PS	Glaxo Wellcome	ORAL

Date:10/15/02ISR Number: 3989827-8Report Type:Expedited (15-DaCompany Report #A0382885A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Hypersensitivity Oxygen Saturation Decreased Rash Pustular		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:10/15/02ISR Number: 3993747-2Report Type:Expedited (15-DaCompany Report #NSADSS2002032582  
 Age:42 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death Hospitalization - Initial or Prolonged SEE IMAGE Required Intervention to Prevent Permanent 60 TABLE, Impairment/Damage ORAL	Acute Respiratory Distress Syndrome Aspiration  Coma Completed Suicide Convulsion  Depressed Level Of Consciousness Hypotension  Intentional Misuse	Literature Health Professional	Topamax (Unspecified) (Topiramate)  Wellbutrin Sr (Amfebutamone Hydrochloride)  Acetaminophen/Propox yphene (Aporex)	PS  SS  SS	ORAL  ORAL  ORAL
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Date:10/15/02ISR Number: 4027522-XReport Type:Periodic Company Report #2001-09-1438  
Age: Gender: I/FU:I

Outcome Dose Hospitalization - Initial or Prolonged	Duration	PT Convulsion	Report Source Health Professional	Product Claritin-D 24 Hour (Loratadine 10mg/Pseudoephedrin Extended Release Tablet	Role PS	Manufacturer	Route ORAL
ORAL				Wellbutrin Sr (Bupropion)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/02ISR Number: 4027579-6Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #HQ2565804JUN2002

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 0.5 MG 1X PER Other 1 DAY, ORAL	Drug Withdrawal Syndrome Grand Mal Convulsion		Ativan (Lorazepam, Tablet)	PS		ORAL
200 MG 2X PER  1 DAY, ORAL			Wellbutrin - Slow Release (Amfebutamone Hydrochloride)	SS		ORAL
			Nortriptyline (Nortriptyline)	C		

Date:10/16/02ISR Number: 3990616-9Report Type:Expedited (15-DaCompany Report #A0382751A  
Age:52 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150MG Per day 1 WK Hospitalization - 80MG Per day Initial or Prolonged 25MG Per day Other	Cardiac Arrest  Ventricular Fibrillation	Health  Professional	Zyban  Micardis  Hydrochlorothiazide	PS  C  C	Glaxo Wellcome  Glaxo Wellcome	ORAL  ORAL  ORAL

Date:10/16/02ISR Number: 3990617-0Report Type:Expedited (15-DaCompany Report #A0383055A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Breast Cancer		Zyban	PS	Glaxo Wellcome	ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Unknown 2 DAY	Abdominal Pain	Health	Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged		Aggression Agitation Anxiety Autonomic Nervous System Imbalance Blood Bilirubin Increased Blood Lactate Dehydrogenase Increased C-Reactive Protein Decreased Haptoglobin Decreased Headache Jaundice Loss Of Consciousness Malaise Normochromic Normocytic Anaemia Pallor Thrombotic Thrombocytopenic Purpura Treatment Noncompliance	Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 3990626-1Report Type:Expedited (15-DaCompany Report #B0282017A  
Age:35 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 10 DAY	Abortion Missed	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged 50MG Per day 8 DAY	Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus Pregnancy		Triflucan	SS		ORAL

Date:10/16/02ISR Number: 3993626-0Report Type:Direct Company Report #CTU 178854  
Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG PO QD	Erythema Multiforme		Wellbutrin Sr	PS		ORAL
Initial or Prolonged	Stevens-Johnson Syndrome					

Date:10/17/02ISR Number: 3991181-2Report Type:Expedited (15-DaCompany Report #B0282085A  
Age:47 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 3 WK	Drug Interaction	Health	Zyban	PS	Glaxo Wellcome	
Hospitalization - Initial or Prolonged	Heart Injury International Normalised Ratio Increased Intracardiac Thrombus Pericardial Haemorrhage Procedural Complication	Professional	Previscan Aspirine Lasilix Diffu K Beta-Blocker(Unspeci fied) Triatec Vasten Discotriner	SS SS C C C C C C	Glaxo Wellcome	ORAL ORAL ORAL ORAL ORAL ORAL ORAL

TRANSDERMAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death		Cerebral Haemorrhage	Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice						
Hospitalization -	Nausea					
per day	29 DAY					
Initial or Prolonged	Oedema		Aspirin	C		ORAL
75MG Per day						
	Urticaria		Perindopril	C		ORAL
2MG Unknown						
			Salbutamol	C	Glaxo Wellcome	
RESPIRATORY						
(INHALATION)	2UNIT Four					
times per day						
			Ipratropium	C	Glaxo Wellcome	
RESPIRATORY						
(INHALATION)	2UNIT Three					
times per day						

Outcome	PT
Death	Abdominal Pain
Required	Abnormal Faeces
Intervention to	Blood Sodium Increased
Prevent Permanent	Cardiac Arrest
Impairment/Damage	Catatonia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Circulatory Collapse Disseminated Intravascular Coagulation	Report Source	Product	Role	Manufacturer	Route
ORAL		Haematocrit Decreased Heart Rate Increased	Literature Health	Risperdal (Risperidone)	PS		ORAL
ORAL		Hypertension Medication Error	Professional	Bupropion (Amfebutamone)	SS		ORAL
ORAL		Mental Status Changes Pyrexia		Diphenhydramine (Diphenhydramine)	SS		ORAL
		Tremor					

Date:10/18/02ISR Number: 3992011-5Report Type:Expedited (15-DaCompany Report #B0134232A  
Age:56 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG	Unknown	Fatigue	Health	Zyban	PS	Glaxo Wellcome	ORAL
RESPIRATORY				Professional	Ventoline	SS	Glaxo Wellcome	
(INHALATION)		4UNIT Twice						
per day					Seretide	SS	Glaxo Wellcome	
RESPIRATORY								
(INHALATION)					Paroxetine Hydrochloride	SS	Glaxo Wellcome	ORAL
UNKNOWN					Nicotine Polacrilex	SS	Glaxo Wellcome	
300MG Three					Theophylline	SS		ORAL
times per day								
.75L per day					Diltiazem Hydrochloride	C	Glaxo Wellcome	ORAL
					Wine	C		ORAL
					Oxygen Therapy	C		



Date:10/18/02ISR Number: 3992012-7Report Type:Expedited (15-DaCompany Report #B0279967A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hemiparesis	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
1TAB Variable							
dose	5	WK					
		Sensory Loss					

Date:10/18/02ISR Number: 3992015-2Report Type:Expedited (15-DaCompany Report #B0282033A  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Arrhythmia Atrial Fibrillation Hyperthyroidism Insomnia Nervousness Palpitations Tremor	Health Professional	Zyntabac	PS	Glaxo Wellcome	ORAL

Date:10/18/02ISR Number: 3992016-4Report Type:Expedited (15-DaCompany Report #B0282196A  
Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening UNKNOWN		Panic Attack Suicidal Ideation		Zyban	PS	Glaxo Wellcome	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/18/02ISR Number: 3992019-XReport Type:Expedited (15-DaCompany Report #B0282592A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
1TAB Unknown	29 DAY		Influenza Like Illness	Zyban	PS	Glaxo Wellcome	ORAL
			Pain In Extremity				
			Paraesthesia				

Date:10/21/02ISR Number: 3992742-7Report Type:Expedited (15-DaCompany Report #A0383421A  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death			Drug Interaction	Zyban	PS	Glaxo Wellcome	ORAL
			Malaise	Effexor	SS		ORAL
				Ativan	C		
				Elavil	C	Glaxo Wellcome	
				Salbutamol	C	Glaxo Wellcome	
				Tylenol	C	Glaxo Wellcome	

Date:10/21/02ISR Number: 3992743-9Report Type:Expedited (15-DaCompany Report #A0383562A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
8 MON			Aneurysm	Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged			Convulsion				
Other			Libido Increased				

Date:10/21/02ISR Number: 3992998-0Report Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-12075131  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
"As required"			Drug Interaction	Trazodone Hcl Tabs	PS	Apothecon	ORAL
Initial or Prolonged			International Normalised	Enoxaparin	C		
			Ratio Decreased	Olanzapine	C		

Clonazepam	C		
Sertraline Hcl	C		
Ibuprofen	C		
Metoclopramide Hcl	C		
Diphenhydramine	C		
Paracetamol Elixir	C		
Docusate Sodium	C		
Warfarin Sodium	I	Bristol-Myers Squibb	
		Company	
Wellbutrin	I		ORAL

Date:10/21/02ISR Number: 3995534-8Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 179097

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State		Zyban 150mg	PS		ORAL
150 MG ORAL		Crying					
		Depression					
		Dyspnoea					
		Feeling Abnormal					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/21/02ISR Number: 3995774-8Report Type:Direct  
Age:48 YR Gender:Female I/FU:I

Company Report #CTU 179088

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Sr 150 Mg	PS		ORAL
150 MG BID		Feeling Jittery					
ORAL		Hypersensitivity					
		Insomnia					

Date:10/21/02ISR Number: 3998273-2Report Type:Expedited (15-DaCompany Report #S02-USA-02227-01  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Health Professional	Lexapro (Escitalopram)	PS		ORAL
Hospitalization - Initial or Prolonged			Company Representative	Wellbutrin (Bupropion Hydrochloride)	SS		
10 MG QD PO				Prozac (Fluoxetine Hydrochloride)	SS		
80 MG QD				Prozac (Fluoxetine Hydrochloride)	SS		
40 MG QD							

Date:10/21/02ISR Number: 3998619-5Report Type:Expedited (15-DaCompany Report #2002061185  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abortion Spontaneous Complications Of Maternal	Foreign Health	Fluconazole (Fluconazole)	PS		ORAL
Hospitalization - Initial or Prolonged		Exposure To Therapeutic	Professional				
50 MG		Drugs		Bupropion	SS		ORAL
(DAILY), ORAL		Intra-Uterine Death					
(DAILY), ORAL							

Maternal Drugs Affecting  
Foetus

Date:10/22/02ISR Number: 3996071-7Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 179183

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Disturbance In Attention		Wellbutrin	PS		
BUCCAL	100 SR	BUCCAL					
		Drug Ineffective		Tricyclic	SS		
		Hallucination		Zoloft	SS		
		Pharmaceutical Product					
		Complaint					

Date:10/22/02ISR Number: 3997826-5Report Type:Expedited (15-DaCompany Report #NSADSS2002032843  
Age:42 YR Gender:Male I/FU:F

Outcome	PT
Death	Brain Herniation
Hospitalization -	Brain Oedema
Initial or Prolonged	Cardiomegaly
Required	Cerebral Artery Occlusion
Intervention to	Coma
Prevent Permanent	Completed Suicide
Impairment/Damage	Cyanosis
	Hepatosplenomegaly
	Loss Of Consciousness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Moaning Pulmonary Congestion Renal Failure					
ORAL		Respiratory Failure Retroperitoneal Haemorrhage Shock	Literature Health Professional	Concerta (Sustained Release Tablet) (Methylphenidate Hydrochloride)	PS		ORAL
		Ventricular Hypertrophy		Bupropion (Amfebutamone)	SS		

Date:10/23/02ISR Number: 3994909-0Report Type:Expedited (15-DaCompany Report #B0280070A  
Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 13 DAY Initial or Prolonged		Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
		C-Reactive Protein Increased Fibrin Increased Hilar Lymphadenopathy Metastases To Lung Neoplasm Oedema Peripheral Paraesthesia Pyrexia Red Blood Cell Sedimentation Rate Increased					

Date:10/23/02ISR Number: 3998905-9Report Type:Expedited (15-DaCompany Report #1662554A  
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - ORALLY Initial or Prolonged ORALLY		Convulsion Disseminated	Literature Health	Regular Strength Tylenol 325 Mg	PS		ORAL
		Intravascular Coagulation Hepatic Failure	Professional	Extra Strength Tylenol Pm Product	SS		ORAL

Hyperpyrexia	Effexor	SS
Posturing	Wellbutrin Sr	SS
Pupil Fixed	Librium	SS
Renal Failure	Ambien	SS
Rhabdomyolysis	Skelaxin	SS
Tachycardia	Midol	SS
	Pyridium	SS
	Aleve	SS
	Relafen	SS

Date:10/23/02ISR Number: 3999404-0Report Type:Expedited (15-DaCompany Report #A0383144A

Age:49 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Warfarin Sodium (Formulation Unknown) (Warfarin Sodium)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Carbamazepine  
 (Formulation  
 Unknown)  
 (Carbamazepine) SS

Date:10/23/02ISR Number: 3999406-4Report Type:Expedited (15-DaCompany Report #A0383146A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Olanzapine  
 (Formulation  
 Unknown)  
 (Olanzapine) SS

Date:10/23/02ISR Number: 3999408-8Report Type:Expedited (15-DaCompany Report #A0383148A

Age:43 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Quetiapine  
 (Formulation  
 Unknown)  
 (Quetiapine) SS

Date:10/23/02ISR Number: 3999410-6Report Type:Expedited (15-DaCompany Report #A0383149A

Age:33 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Literature	Wellbutrin			



ORAL			Health Professional	Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
				Risperidone (Formulation Unknown)			
				(Risperidone)	SS		
				Diphenhydramine (Formulation Unknown)			
				(Diphenhydramine)	SS		

Date:10/23/02ISR Number: 3999566-5Report Type:Expedited (15-DaCompany Report #A0383151A  
Age:52 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
				Coreg (Formulation			

Freedom Of Information (FOI) Report

Unknown) (Cavedilol) SS  
 Amlodipine +  
 Benazepril  
 (Formulation  
 Unknown) (Amlodipine  
 + Benazepril) SS

Date:10/23/02ISR Number: 3999568-9Report Type:Expedited (15-DaCompany Report #A0383128A  
 Age:52 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Fluvoxamine (Formulation Unkown) (Fluvoxamine) Ethanol (Formulation Unknown) (Alcohol)	SS SS		

Date:10/23/02ISR Number: 3999571-9Report Type:Expedited (15-DaCompany Report #A0383129A  
 Age:34 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecfied Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Isotretinoin (Formulation Unknown) (Isotretinoin) Diphenhydramine (Formulation Unknown) (Diphenhydramine)	SS SS		

Date:10/23/02ISR Number: 3999575-6Report Type:Expedited (15-DaCompany Report #A0383132A  
Age:67 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Sertraline (Formulation Unknown) (Sertraline)	SS		

Date:10/23/02ISR Number: 3999578-1Report Type:Expedited (15-DaCompany Report #A0383133A  
Age:42 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL				Hydrochloride)	PS		ORAL
				Topiramate (Formulation Unknown) (Topiramate)	SS		
				Dextroprop. + Paracetamol (Formulation Unknown) (Propoxyphene +	SS		

Date:10/23/02ISR Number: 3999588-4Report Type:Expedited (15-DaCompany Report #A0383134A  
Age:19 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Valproic Acid (Valproic Acid) Olanzapine (Olanzapine)	SS SS		

Date:10/23/02ISR Number: 3999590-2Report Type:Expedited (15-DaCompany Report #A0383135A  
Age:31 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Citalopram (Formulation Unknown) (Citalopram) Amphetamine (Formulation Unknown)	SS		

(Amphetamine) SS  
Dexedrine  
(Formulation  
Unknown)  
(Dextroamphetamine  
Sulfate) SS

Date:10/23/02ISR Number: 3999592-6Report Type:Expedited (15-DaCompany Report #A0383136A

Age:36 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Doxepin (Doxepin)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/23/02ISR Number: 3999594-XReport Type:Expedited (15-DaCompany Report #A0383137A

Age:28 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Trazodone (Formulation Unknown) (Trazodone) Venlafaxine Hydrochloride (Formulation Unknown) (Venlafaxine	SS  SS		

Date:10/23/02ISR Number: 3999596-3Report Type:Expedited (15-DaCompany Report #A0383140A

Age:30 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Diphenhydramine + Paraceta. (Formulation Unknown) (Acetaminophen + Venlafaxine Hydrochloride (Formulation Unknown) (Venlafaxine	SS  SS		

Date:10/23/02ISR Number: 3999598-7Report Type:Expedited (15-DaCompany Report #A0383141A

Age:42 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Hydrocodone + Paracetamol (Formulation Unknown) (Hydrocodone + Dextropropoxyphene (Formulation Unknown) (Dextropropoxyphene)	SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/23/02ISR Number: 3999600-2Report Type:Expedited (15-DaCompany Report #A0383142A

Age:37 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Aspirin (Formulation Unknown) (Aspirin)	SS		
				Olanzapine (Formulation Unknown) (Olanzapine)	SS		

Date:10/23/02ISR Number: 3999602-6Report Type:Expedited (15-DaCompany Report #A0383143A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Tramadol Hydrochloride (Formulation Unknown) (Tramadol Hydrochloride)	SS		
				Metoclopramide (Formulation Unknown) (Metoclopramide)	SS		

Date:10/23/02ISR Number: 3999606-3Report Type:Expedited (15-DaCompany Report #A0383124A

Age:15 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Wellbutrin			



ORAL	Intentional Misuse	Health Professional	Unspecified Tablet (Bupropion Hydrochloride)	PS	ORAL
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Date:10/23/02ISR Number: 3999609-9Report Type:Expedited (15-DaCompany Report #A0383127A  
 Age:17 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Fluvoxamine (Formulation Unknown) (Fluvoxamine)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/23/02ISR Number: 3999612-9Report Type:Expedited (15-DaCompany Report #A0383147A

Age:33 YR Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL					Quetiapine (Formulation Unknown) (Quetiapine)	SS		

Date:10/23/02ISR Number: 3999617-8Report Type:Expedited (15-DaCompany Report #A0383130A

Age:42 YR Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL					Methylphenidate (Formulation Unknown) (Methylphenidate)	SS		

Date:10/23/02ISR Number: 3999628-2Report Type:Expedited (15-DaCompany Report #A0383131A

Age:42 YR Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide Intentional Misuse	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL					Methylphenidate (Formulation Unknown)			

Date:10/23/02ISR Number: 4027648-0Report Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #NSADSS2002015369

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Chest Pain	Health	Remicade ( 5 Mg/Ml			
Hospitalization -		Headache	Professional	Lyophilized Powder)			
Initial or Prolonged		Hypersensitivity		(Infliximab,			
Other		Hypertension		Recombinant)	PS		
INTRAVENOUS	500MG	, 1 IN					
1 TIMES, IV		Pain					
		Sexual Dysfunction		Wellbutrin			
		Vertigo		(Amfebutamone			
				Hydrochloride)	SS		
				Vioxx (Rofecoxib)	SS		
				Prednisone	C		
				Methotrexate	C		
				Celebrex (Celecoxib)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/25/02ISR Number: 3996314-XReport Type:Expedited (15-DaCompany Report #A0383992A

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Atrial Fibrillation		Zyban	PS	Glaxo Wellcome	ORAL

Date:10/28/02ISR Number: 3997563-7Report Type:Expedited (15-DaCompany Report #A0383140A

Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health	Bupropion	PS	Glaxo Wellcome	ORAL
Hospitalization - Initial or Prolonged		Convulsion	Professional	Relafen	SS	Glaxo Wellcome	ORAL
UNKNOWN		Disseminated Intravascular Coagulation		Acetaminophen + Diphenhydramine	SS		
UNKNOWN		Hepatic Failure		Venlafaxine	SS		
UNKNOWN		Intentional Misuse		Tylenol	SS	Glaxo Wellcome	
UNKNOWN		Nystagmus		Librium	SS		
UNKNOWN		Posturing		Ambien	SS		
UNKNOWN		Pupil Fixed		Skelaxin	SS		
UNKNOWN		Pyrexia		Midol	SS		
UNKNOWN		Renal Failure		Pyridium	SS		
UNKNOWN		Rhabdomyolysis		Aleve	SS		
UNKNOWN		Tachycardia					

Date:10/28/02ISR Number: 3997567-4Report Type:Expedited (15-DaCompany Report #B0135045A

Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		No Adverse Drug Effect	Health	Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown			Professional				

Date:10/28/02ISR Number: 3997580-7Report Type:Expedited (15-DaCompany Report #B0282937A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Drug Tolerance Decreased Malaise Muscle Spasms Visual Disturbance		Zyban	PS	Glaxo Wellcome	ORAL

Date:10/28/02ISR Number: 3997586-8Report Type:Expedited (15-DaCompany Report #B0283367A  
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 15 DAY		Papilloedema Visual Acuity Reduced		Zyban	PS	Glaxo Wellcome	ORAL

Date:10/28/02ISR Number: 3997587-XReport Type:Expedited (15-DaCompany Report #B0283519A  
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG Per day 10 DAY		Oedema Peripheral Urticaria Wheezing		Zyban	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/29/02ISR Number: 3998526-8Report Type:Expedited (15-DaCompany Report #A0384610A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aortic Valve Stenosis		Wellbutrin	PS	Glaxo Wellcome	ORAL
		Condition Aggravated					

Date:10/29/02ISR Number: 3998534-7Report Type:Expedited (15-DaCompany Report #B0283267A

Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Conversion Disorder		Zyban	PS	Glaxo Wellcome	
UNKNOWN		56 DAY					
Hospitalization -		Dysphonia					
Initial or Prolonged		Urticaria					

Date:10/29/02ISR Number: 3998535-9Report Type:Expedited (15-DaCompany Report #B0283291A

Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Anxiety		Zyban	PS	Glaxo Wellcome	ORAL
3 DAY		Meniere'S Disease					
		Nausea					
		Palpitations					
		Rash					
		Visual Disturbance					

Date:10/29/02ISR Number: 3998536-0Report Type:Expedited (15-DaCompany Report #B0283322A

Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dyspnoea		Zyban	PS	Glaxo Wellcome	ORAL
10 DAY							
Initial or Prolonged		Nephritis					

Date:10/29/02ISR Number: 3998543-8Report Type:Expedited (15-DaCompany Report #B0283515A  
Age:60 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Dizziness		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Headache		Thyroid Medication	C		
	Inner Ear Disorder					
	Psychosomatic Disease					

Date:10/29/02ISR Number: 4000461-6Report Type:Direct Company Report #CTU 179774  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Rash Pruritic		Gabapril 2 Mg	PS		ORAL
2 MG QHS ORAL	Rash Scaly		Tenormin	C		
			Valium	C		
			Zoloft	C		
			Lomotil	C		
			Synthroid	I		
			Kcl	I		
			Wellbutrin	I		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/30/02ISR Number: 3999548-3Report Type:Expedited (15-DaCompany Report #D0039368A  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Aortic Valve Stenosis Disturbance In Attention Drug Dependence Memory Impairment Restlessness Visual Disturbance		Zyban Diclofenac	PS C	Glaxo Wellcome	ORAL ORAL

Date:10/31/02ISR Number: 4000512-9Report Type:Expedited (15-DaCompany Report #A0382619A  
Age:9 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG Per day 22 DAY Initial or Prolonged Other		Hypersensitivity Serum Sickness Stevens-Johnson Syndrome	Health Professional	Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:10/31/02ISR Number: 4000517-8Report Type:Expedited (15-DaCompany Report #B0282735A  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day 3 WK		Alanine Aminotransferase Increased Arthralgia Blood Creatine Phosphokinase Increased Difficulty In Walking Myositis		Zyban	PS	Glaxo Wellcome	ORAL

Date:10/31/02ISR Number: 4003925-4Report Type:Expedited (15-DaCompany Report #HQ4742622OCT2002  
Age:30 YR Gender:Male I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Completed Suicide Convulsion	Literature	Effexor (Venlafaxine Hydrochloride, Tablet)	PS		ORAL
OVERDOSE		Disseminated					
AMOUNT		Intravascular Coagulation					
UNKNOWN, ORAL		Electroencephalogram Abnormal		Aleve (Naproxen Sodium, )	SS		ORAL
OVERDOSE		Hepatic Failure					
AMOUNT		Mydriasis					
UNKNOWN, ORAL		Nystagmus Posturing		Ambien (Zolpidem Tartrate, )	SS		ORAL
OVERDOSE		Pupil Fixed					
AMOUNT		Pyrexia					
UNKNOWN, ORAL		Renal Failure Rhabdomyolysis Tachycardia		Librium (Chlordiazepoxide Hydrochloride, )	SS		ORAL
OVERDOSE							
AMOUNT							
UNKNOWN, ORAL							
OVERDOSE				Metaxalone (Metaxalone, )	SS		ORAL
AMOUNT							
UNKNOWN, ORAL							
				Midol (Acetylsalicylic			

Freedom Of Information (FOI) Report

OVERDOSE			Acid/Caffeine/Cinnam edrine/Phenacetin, )	SS	
AMOUNT			Pyridium (Phenazopyridine Hydrochloride, )	SS	ORAL
UNKNOWN, ORAL					
OVERDOSE			Relafen (Nabumetone, )	SS	ORAL
AMOUNT					
UNKNOWN, ORAL					
OVERDOSE			Tylenol (Paracetamol, )	SS	ORAL
AMOUNT					
UNKNOWN, ORAL					
OVERDOSE			Tylenol Pm (Diphenhydramine/Par acetamol, )	SS	ORAL
AMOUNT					
UNKNOWN, ORAL					
OVERDOSE			Wellbutrin (Amfebutamone Hydrochloride, )	SS	ORAL
AMOUNT					
UNKNOWN, ORAL					

Date:10/31/02ISR Number: 4004475-1Report Type:Expedited (15-DaCompany Report #HQ4898628OCT2002  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Death	Drug Interaction	Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS	ORAL
75 MG 1 X PER					
1 DAY, ORAL					

Zyban (Amfebutamone Hydrochloride)	SS
Tylenol (Paracetamol)	C
Salbutamol (Salbutamol)	C
Ativan (Lorazepam)	C
Elavil (Amitriptyline Hydrochloride)	C

Date:10/31/02ISR Number: 4004514-8Report Type:Expedited (15-DaCompany Report #200213918BCC  
Age:30 YR Gender:Male I/FU:I

Outcome	PT
Death	Cardiac Arrest
Hospitalization -	Completed Suicide
Initial or Prolonged	Convulsion
	Disseminated
	Intravascular Coagulation
	Hepatic Failure
	Nystagmus
	Posturing
	Pupil Fixed
	Pyrexia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Renal Failure Rhabdomyolysis Tachycardia	Report Source	Product	Role	Manufacturer	Route
ORAL			Literature Health	Aleve (Naproxen Sodium)	PS		ORAL
			Professional Other	Midol	SS		
				Tylenol Pm	SS		
				Effexor (Venlafaxine Hydrochloride)	SS		
				Wellbutrin Sr (Amfebutamone Hydrochloride)	SS		
				Librium	SS		
				Ambien (Zolpidem Tartrate)	SS		
				Skelaxin (Metaxalone Pyridium (Phenazopyridine Hydrochloride))	SS		
				Tylenol (Paracetamol)	SS		
				Relafen (Nabumetone)	SS		

Date:10/31/02ISR Number: 4004888-8Report Type:Expedited (15-DaCompany Report #2002AP03609  
Age:43 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature Health	Quetiapine	PS		
		Completed Suicide	Professional	Bupropion	SS		
		Intentional Misuse					

Date:10/31/02ISR Number: 4004924-9Report Type:Expedited (15-DaCompany Report #2002AP03602  
Age:33 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature Health	Quetiapine	PS		
		Completed Suicide	Professional	Bupropion	SS		
		Intentional Misuse					

Date:11/01/02ISR Number: 4001455-7Report Type:Expedited (15-DaCompany Report #B0283370A  
Age:26 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 60TABS Single Initial or Prolonged dose 1 DAY	Amnesia Confusional State Delirium		Zyban Equanil	PS C	Glaxo Wellcome	ORAL ORAL
1 DAY	Intentional Misuse Somnolence Suicide Attempt					

Date:11/01/02ISR Number: 4001460-0Report Type:Expedited (15-DaCompany Report #D0039698A  
Age:28 YR Gender:Male I/FU:I

Outcome	PT
Other	Dry Mouth Intentional Misuse Overdose

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Suicide Attempt Tremor				
Dose	Duration		Report Source	Product	Role	Manufacturer
12TABS Single			Health	Zyban	PS	Glaxo Wellcome
dose	1 DAY		Professional			
1 DAY				Alcohol	SS	
						ORAL

Date:11/01/02ISR Number: 4005534-XReport Type:Expedited (15-DaCompany Report #HQ4742622OCT2002  
Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Disseminated Intravascular Coagulation	Literature	Effexor (Venlafaxine Hydrochloride, Tablet)	PS		ORAL
OVERDOSE		Hepatic Failure					
AMOUNT		Hyperpyrexia					
UNKNOWN, ORAL		Mydriasis Nervous System Disorder		Aleve (Naproxen Sodium,)	SS		ORAL
OVERDOSE		Pupil Fixed					
AMOUNT		Renal Failure					
UNKNOWN, ORAL		Rhabdomyolysis		Ambien (Zolpidem Tartrate)	SS		ORAL
OVERDOSE							
AMOUNT							
UNKNOWN, ORAL				Librium (Chlordiazepoxide Hydrochloride)	SS		ORAL
OVERDOSE							
AMOUNT							

UNKNOWN, ORAL

Metaxalone  
(Metaxalone) SS ORAL

OVERDOSE

AMOUNT

UNKNOWN, ORAL

Midol  
(Acetylsalicylic  
Acid/Caffeine/Cinnam  
edrine/Phenacetin,) SS  
Pyridium  
(Phenazopyridine  
Hydrochloride,) SS ORAL

OVERDOSE

AMOUNT

UNKNOWN, ORAL

Relafen (Nabumetone) SS ORAL

OVERDOSE

AMOUNT

UNKNOWN, ORAL

Tylenol  
(Paracetamol) SS ORAL

OVERDOSE

AMOUNT

UNKNOWN, ORAL

Tylenol Pm  
(Diphenhydramine/Par  
acetamol) SS ORAL

OVERDOSE

AMOUNT

UNKNOWN, ORAL

Wellbutrin  
(Amfebutamone  
Hydrochloride) SS ORAL

OVERDOSE

AMOUNT

UNKNOWN, ORAL





FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/04/02ISR Number: 4006263-9Report Type:Expedited (15-DaCompany Report #HQ4984731OCT2002  
 Age:34 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature	Diphenhydramine Hcl (Diphenhydramine Hydrochloride, Injection)	PS		ORAL
ORAL				Bupropion (Amfebutamone, )	SS		ORAL
ORAL				Isotretinoin (Isotretinoin, )	SS		ORAL

Date:11/05/02ISR Number: 4006764-3Report Type:Expedited (15-DaCompany Report #HQ4986031OCT2002  
 Age:33 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse	Literature	Diphenhydramine Hcl (Diphenhydramine Hydrochloride, Injection)	PS		
OVERDOSE							
AMOUNT							
UNKNOWN				Bupropion (Amfebutamone, )	SS		
OVERDOSE							
AMOUNT							
UNKNOWN				Risperidone (Risperidone, )	SS		
OVERDOSE							
AMOUNT							
UNKNOWN							

Date:11/05/02ISR Number: 4006923-XReport Type:Expedited (15-DaCompany Report #HQ4926530OCT2002

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Overdose	Literature	Metoclopramide (Metoclopramide Hydrochloride,	PS		ORAL
OVEDOSE							
AMOUNT							
UNKNOWN, ORAL							
OVERDOSE				Bupropion (Amfebutamone, )	SS		ORAL
AMOUNT							
UNKNOWN, ORAL							
OVERDOSE				Tramadol (Tramadol, )	SS		ORAL
AMOUNT							
UNKNOWN, ORAL							

Date:11/06/02ISR Number: 4004617-8Report Type:Expedited (15-DaCompany Report #A0375799A

Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Arthralgia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Dysgeusia					
per day	4	WK					
		Dyspepsia Feeling Jittery		Triamterene/Hctz 37.5mg/25mg	C	Glaxo Wellcome	ORAL
1UNIT Per day							
.5MG Per day		Pain In Extremity		Xanax	C		ORAL
		YR					

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Freedom Of Information (FOI) Report

1.25MG Per  
 day YR  
 Date:11/06/02ISR Number: 4004618-XReport Type:Expedited (15-DaCompany Report #A0376093A  
 Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Arthralgia					
per day	19	DAY		No Concurrent Medication	C		
		Cyanosis					
		Difficulty In Walking					
		Insomnia					
		Oedema Peripheral					
		Rash					
		Urticaria					

Date:11/06/02ISR Number: 4004620-8Report Type:Expedited (15-DaCompany Report #A0383335A  
 Age:74 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hyponatraemia		Wellbutrin	PS	Glaxo Wellcome	ORAL
300MG Per day		Inappropriate		Micardis	C	Glaxo Wellcome	ORAL
40MG Per day	2	MON		Pravachol	C		ORAL
40MG Per day	10	YR	Antidiuretic Hormone	Aspirin	C		ORAL
81MG Per day	10	YR	Secretion	Neurontin	C		ORAL
300MG As							
required	YR			Niaspan	C		
UNKNOWN	500MG Per day						

Date:11/06/02ISR Number: 4004623-3Report Type:Expedited (15-DaCompany Report #A0385250A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	6	MON		Wellbutrin	PS	Glaxo Wellcome	ORAL
UNKNOWN				Remeron	C		
UNKNOWN				Xanax	C		
UNKNOWN				Gas-X	C		

Date:11/06/02ISR Number: 4004645-2Report Type:Expedited (15-DaCompany Report #B0283868A  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day	8	DAY		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Increased C-Reactive Protein Increased Csf Protein Increased Grand Mal Convulsion Pyrexia Red Blood Cell Count Abnormal		Urbanyl Alcohol	C C		ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/06/02ISR Number: 4004648-8Report Type:Expedited (15-DaCompany Report #B0283974A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphonia		Zyban	PS	Glaxo Wellcome	ORAL
24 DAY		Dysphonia Laryngitis					

Date:11/06/02ISR Number: 4004649-XReport Type:Expedited (15-DaCompany Report #B0283976A  
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Emotional Distress		Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown 12 DAY		Fear					

Date:11/06/02ISR Number: 4004650-6Report Type:Expedited (15-DaCompany Report #B0283979A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Drug Withdrawal Syndrome		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice Other per day	54 DAY	Panic Attack					

Date:11/06/02ISR Number: 4007804-8Report Type:Expedited (15-DaCompany Report #2002AP03419  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 50 MG DAILY		Blood Glucose Increased	Consumer	Atenolol	PS		ORAL
Initial or Prolonged PO		Cardiac Disorder					
360 MG DAILY		Cystitis		Cardizem	SS		ORAL

PO	Hypertension				
150 MG DAILY	Hypotension		Zyban	SS	ORAL
PO	Medication Error				
	Myocardial Infarction		Glucophage "Abic"	C	
			Altace	C	
			Lantus	C	

Date:11/06/02ISR Number: 4023761-2Report Type:Periodic Company Report #A0161304A  
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required	150 MG /	Grand Mal Convulsion Loss Of Consciousness	Health Professional Company Representative	Wellbutrin Sr (Bupropion Hydrochloride)	PS		ORAL
Intervention to THREE TIMES Prevent Permanent PER DAY / Impairment/Damage ORAL				Dexamphetamine Sulphate	C		

Date:11/06/02ISR Number: 4023764-8Report Type:Periodic Company Report #A0160041A  
 Age:36 YR Gender:Female I/FU:I

Outcome  
Life-Threatening  
Required  
Intervention to  
Prevent Permanent

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG / PER DAY / ORAL		Asthma Oedema Peripheral Pruritus	Health Professional Company	Wellbutrin Sr (Bupropion Hydrochloride)	PS		ORAL
		Rash	Representative				

Date:11/06/02ISR Number: 4023765-XReport Type:Periodic Company Report #A0152014A  
 Age:61 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1 TABLET / PER DAY / ORAL		Dyspnoea Hypersensitivity Urticaria	Consumer	Wellbutrin Sr (Bupropion Hydrochloride)	PS		ORAL
			Visual Acuity Reduced					
					Blood Pressure Medication	C		

Date:11/06/02ISR Number: 4023767-3Report Type:Periodic Company Report #A0141485A  
 Age:25 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	150 MG / TWICE PER DAY / ORAL		Arthralgia Asthenia Burning Sensation	Consumer	Wellbutrin Sr (Bupropion Hydrochloride)	PS		ORAL
			Dyspnoea					
			Face Oedema					
			Fear Hypersensitivity Mobility Decreased					

Oedema Peripheral  
Oropharyngeal Swelling  
Pain  
Pain In Extremity  
Pruritus  
Rash  
Swelling  
Tenderness  
Urticaria

Date:11/06/02ISR Number: 4028909-1Report Type:Periodic Company Report #A0380851A  
Age:30 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other	Convulsion	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		
200 MG/ PER DAY/						



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/06/02ISR Number: 4028910-8Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #A0380199A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Migraine Pulmonary Embolism	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
250 MG/ TWICE						
PER DAY/ ORAL 2 YR						

Date:11/06/02ISR Number: 4028911-XReport Type:Periodic  
Age:27 YR Gender:Male I/FU:I

Company Report #A0372787A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Hypersensitivity	Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE						
PER DAY/ ORAL						

Date:11/06/02ISR Number: 4028912-1Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #A0370877A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Fatigue Rhabdomyolysis	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ TWICE						
PER DAY/ ORAL 4 YR						
			Alprazolam	C		
			Zolpidem Tartrate	C		

Date:11/06/02ISR Number: 4028913-3Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0370550A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Cardiac Disorder	Health	Wellbutrin Sr			
Hospitalization -	Food Intolerance	Professional	Tablet-Controlled			
Initial or Prolonged	Hypersensitivity		Release (Bupropion			
Required	Psychomotor Hyperactivity		Hydrochloride)	PS		ORAL
150 MG/ TWICE						
Intervention to	Throat Tightness					
PER DAY/ ORAL						
Prevent Permanent	Urticaria		Diphenhydramine			
Impairment/Damage			Hydrochloride			
			(Formulation			
			Unknown)			
			(Diphenhydramine	SS		
			Alprazolam	C		
			Diuretic	C		

Date:11/06/02ISR Number: 4028914-5Report Type:Periodic  
Age:20 YR Gender:Female I/FU:I

Company Report #A0370004A

Outcome  
Hospitalization -  
Initial or Prolonged  
Required  
Intervention to  
Prevent Permanent

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/ PER DAY/ ORAL	6 MON	Convulsion	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

Date:11/06/02ISR Number: 4028915-7Report Type:Periodic Company Report #A0369151A  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other Required Intervention to SEE DOSAGE Prevent Permanent Impairment/Damage		Convulsion	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

Date:11/06/02ISR Number: 4028916-9Report Type:Periodic Company Report #A0368833A  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

(THERAPY

DATES: SEE

TEXT)

5 MG/ IN THE

MORNING/ ORAL

(THERAPY

DATES: YEARS)

Lorazepam  
(Formulation  
Unknown) (Lorazepam) SS

ORAL

Nortriptyline C

Date:11/06/02ISR Number: 4028917-0Report Type:Periodic  
Age:52 YR Gender:Male I/FU:I

Company Report #A0368730A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Pancreatitis	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL				Lamivudine	C		
				Efavirenz	C		
				Didanosine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/06/02ISR Number: 4028918-2Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #A0368647A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Grand Mal Convulsion	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
100 MG/ TWICE  PER DAY/ ORAL			Fluvoxamine Maleate Olanzapine	C C		

Date:11/06/02ISR Number: 4028919-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #A0366264A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Convulsion Hypoglycaemia	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
UNK / UNK /  ORAL			Sulfonylurea Agent Insulin	C C		

Date:11/06/02ISR Number: 4028920-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0366204A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Stevens-Johnson Syndrome	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
UNK / UNKNOWN  / ORAL						

Date:11/06/02ISR Number: 4028921-2Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #A0365838A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Life-Threatening	Grand Mal Convulsion	Health	Wellbutrin Sr			
Hospitalization -	Joint Dislocation	Professional	Tablet-Controlled			
Initial or Prolonged		Company	Release (Bupropion			
Disability		Representative	Hydrochloride)	PS		ORAL
150 MG/ TWICE						
Required						
PER DAY /						
Intervention to						
ORAL						
Prevent Permanent						
Impairment/Damage						

Date:11/06/02ISR Number: 4028922-4Report Type:Periodic  
Age:74 YR Gender:Female I/FU:I

Company Report #A0365478A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Cerebrovascular Accident	Health	Wellbutrin Sr			
Initial or Prolonged	Drug Interaction	Professional	Tablet-Controlled			
			Release (Bupropion			
			Hydrochloride)	PS		ORAL
100 MG/ TWICE						
PER DAY /						
ORAL						
			Nortriptyline Hcl			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

UNK / UNK /	Capsule (Nortriptyline Hcl)	SS	ORAL
ORAL			
UNK / UNK /	Citaloram Hydrobromide (Formulation Unknown) (Citalopram Hydrobromide)	SS	ORAL
ORAL			

Date:11/06/02ISR Number: 4028923-6Report Type:Periodic Company Report #A0171156A  
 Age:46 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Delusion Hallucination Medication Error Paranoia	Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
50 MG / TWICE PER DAY /	Psychotic Disorder					
ORAL						

Thiothixene C  
 Methysergide Maleate C  
 Benztropine Mesylate C

Date:11/06/02ISR Number: 4028924-8Report Type:Periodic Company Report #A0169144A  
 Age:11 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Abasia Arthralgia Oedema Peripheral Pain In Extremity	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
100 MG / PER DAY / ORAL	Rash					

Date:11/06/02ISR Number: 4028925-XReport Type:Periodic  
 Age:33 YR Gender:Female I/FU:F

Company Report #A0168576A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Required UNK / SEE Intervention to DOSAGE TEXT / Prevent Permanent ORAL Impairment/Damage		Coma Convulsion Overdose	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
UNK/ SEE DOSAGE TEXT / ORAL ORAL				Fluoxetine Hydrochloride Tablet (Fluoxetine Hydrochloride)	SS		ORAL
				Ethanol (Formulation Unknown) (Alcohol)	SS		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/06/02ISR Number: 4028926-1Report Type:Periodic  
Age:8 YR Gender:Female I/FU:I

Company Report #A0168351A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Required Intervention to UNK / SEE Prevent Permanent DOSAGE TEXT / Impairment/Damage ORAL	Amnesia Complex Partial Seizures Confusional State Coordination Abnormal Dyskinesia Grand Mal Convulsion Tardive Dyskinesia	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)  Clonidine	PS  C		ORAL

Date:11/06/02ISR Number: 4028927-3Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #A0167595A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged  150 MG / PER DAY / ORAL	Chest Pain Rhabdomyolysis	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)  Semisodium Valproate Risperidone	PS  C C		ORAL

Date:11/06/02ISR Number: 4028928-5Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #A0167591A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged  150 MG / TWICE PER DAY	Abdominal Pain Upper Chest Discomfort Chest Pain Diarrhoea Ear Pruritus	Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

/ ORAL  
Erythema  
Flushing

Date:11/06/02ISR Number: 4028929-7Report Type:Periodic Company Report #A0171410A  
Age:54 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/PER DAY/ORAL	Cerebral Infarction Cerebrovascular Accident Difficulty In Walking Dysarthria Dysgraphia Hemiparesis Nausea Speech Disorder Vomiting	Health Professional Company Representative	Wellbutrin Sr (Bupropion Hydrochloride)  Quinapril Hydrochloride Carbamazepine Darvocet-N Glipizide Metformin Hydrochloride	          C C C C  C		          ORAL

Date:11/06/02ISR Number: 4028930-3Report Type:Periodic Company Report #A0166131A  
Age: Gender: I/FU:I

Outcome	PT	Report Source
Hospitalization - Initial or Prolonged	Abdominal Pain	Health Professional

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration		Company Representative	Product	Role	Manufacturer	Route
ORAL				Wellbutrin Sr (Bupropion Hydrochloride)	PS		ORAL

Date:11/06/02ISR Number: 4028931-5Report Type:Periodic Company Report #A0166132A  
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - Initial or Prolonged	Abdominal Pain	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL						

Date:11/06/02ISR Number: 4028932-7Report Type:Periodic Company Report #A0166133A  
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - Initial or Prolonged	Abdominal Pain	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL						

Date:11/06/02ISR Number: 4028933-9Report Type:Periodic Company Report #A0164485A  
Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - Initial or Prolonged Disability 150 MG/ ORAL Required	Grand Mal Convulsion Joint Dislocation Upper Limb Fracture	Health Professional Company Representative	Wellbutrin Sr (Bupropion Hydrochloride)	PS		ORAL

Intervention to  
Prevent Permanent  
Impairment/Damage

Date:11/06/02ISR Number: 4028934-0Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #A0164455A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Required Intervention to 150 MG / PER Prevent Permanent DAY / ORAL Impairment/Damage	Anorexia Convulsion Diarrhoea	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)  Olanzapine	PS  C		ORAL

Date:11/06/02ISR Number: 4028935-2Report Type:Periodic  
Age:13 YR Gender:Female I/FU:I

Company Report #A0162501A

Outcome	PT
Hospitalization - Initial or Prolonged	Blood Pressure Systolic Increased Grand Mal Convulsion Hypoacusis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Intentional Misuse Irritability Suicide Attempt	Report Source	Product	Role	Manufacturer	Route
150 MG/ORAL		Tachycardia Tremor	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

Date:11/06/02ISR Number: 4028936-4Report Type:Periodic Company Report #A0162571A  
Age:58 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Accidental Overdose Syncope Tremor	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL					Thyroxine Sodium Valsartan	C C		

Date:11/06/02ISR Number: 4028937-6Report Type:Periodic Company Report #A0162687A  
Age:43 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged	150 MG /		Angioneurotic Oedema	Health Professional Company Representative	Wellbutrin Sr  (Bupropion Hydrochloride)	PS		ORAL
TWICE PER DAY								
/ ORAL		4 WK						

Date:11/06/02ISR Number: 4028938-8Report Type:Periodic Company Report #A0162688A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Oedema Peripheral	Consumer Health Professional Company	Wellbutrin Sr (Bupropion Hydrochloride)	PS		ORAL
150 MG /			Representative				
TWICE PER DAY							
/ ORAL,							
MONTHS							

Date:11/06/02ISR Number: 4028939-XReport Type:Periodic Company Report #A0360567A  
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Rash	Health Professional	Wellbutrin Sr (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE							
PER DAY/ORAL				Etanercept	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/06/02ISR Number: 4028940-6Report Type:Periodic  
Age:55 YR Gender:Male I/FU:I

Company Report #A0360368A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/TWICE PER DAY/ORAL	Grand Mal Convulsion	Health Professional	Wellbutrin Sr (Bupropion Hydrochloride)	PS		ORAL

Date:11/06/02ISR Number: 4028941-8Report Type:Periodic  
Age:19 YR Gender:Male I/FU:I

Company Report #A0359697A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Required Intervention to 200 MG/TWICE Prevent Permanent PER DAY/ORAL 6 MON Impairment/Damage 200 MG/PER DAY/ORAL	Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Sr (Bupropion Hydrochloride)	PS		ORAL
			Desipramine (Desipramine)	SS		ORAL
			Carbamazepine	C		

Date:11/06/02ISR Number: 4028942-XReport Type:Periodic  
Age:87 YR Gender:Female I/FU:I

Company Report #A0175740A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Disability 150 MG/PER DAY/ORAL	Asthenia Blood Pressure Fluctuation Decreased Appetite Difficulty In Walking Dysstasia	Consumer	Wellbutrin Sr (Bupropion Hydrochloride)	PS		ORAL
			Citalopram			

Feeding Disorder  
Heart Rate Irregular  
Tremor

Hydrobromide C  
Megestrol Acetate C  
Atenolol C

Date:11/06/02ISR Number: 4028943-1Report Type:Periodic Company Report #A0175691A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Convulsion	Health Professional Company Representative	Wellbutrin Sr  (Bupropion Hydrochloride)	PS		ORAL
150 MG/PER DAY/ORAL						

Date:11/06/02ISR Number: 4028944-3Report Type:Periodic Company Report #A0175373A  
Age:37 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - Initial or Prolonged Required	Grand Mal Convulsion	Health Professional	Wellbutrin Sr  (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE Intervention to PER DAY/ORAL Prevent Permanent Impairment/Damage ORAL			Theophylline (Theophylline)  Prednisone	SS		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

VARIABLE	(Prednisone)	SS	ORAL
DOSE/ORAL	Gatifloxacin		
400 MG/PER	(Gatifloxacin)	SS	ORAL
DAY/ORAL			

Date:11/06/02ISR Number: 4028945-5Report Type:Periodic Company Report #A0174702A  
 Age:54 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Anaphylactic Reaction	Health Professional Company Representative	Wellbutrin Sr  (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE			Adderall	C		
PER DAY/ORAL						

Date:11/06/02ISR Number: 4028946-7Report Type:Periodic Company Report #A0174492A  
 Age:46 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Flushing Insomnia Shock	Consumer	Wellbutrin Sr  (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE						
PER DAY/ORAL						

Date:11/06/02ISR Number: 4028947-9Report Type:Periodic Company Report #A0173914A  
 Age:25 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization -	Convulsion	Health	Wellbutrin Sr			

Initial or Prolonged  
Disability

Professional  
Company  
Representative

Tablet-Controlled  
Release (Bupropion  
Hydrochloride)

PS

ORAL

150 MG/ORAL 1 WK

Date:11/06/02ISR Number: 4028948-0Report Type:Periodic  
Age:18 YR Gender:Male I/FU:I

Company Report #A0173089A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - Initial or Prolonged Required SEE DOSAGE Intervention to TEXT/ORAL Prevent Permanent Impairment/Damage	Convulsion	Health Professional Company Representative	Wellbutrin Sr  (Bupropion Hydrochloride)	PS		ORAL
			Adderall Cannabis	C C		

Date:11/06/02ISR Number: 4028949-2Report Type:Periodic  
Age:23 YR Gender:Male I/FU:I

Company Report #A0171989A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Required Intervention to Prevent Permanent Impairment/Damage 150 MG/TWICE PER DAY/ORAL	Convulsion Postictal State	Health Professional	Wellbutrin Sr  (Bupropion Hydrochloride)	PS		ORAL
			Liotrix	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/06/02ISR Number: 4028950-9Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #A0379128A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other	Convulsion	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
200 MG/ TWICE  PER DAY/ ORAL						

Date:11/06/02ISR Number: 4028951-0Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #A0377112A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other	Convulsion	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
300 MG/ PER  DAY/ ORAL						

Date:11/06/02ISR Number: 4028952-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0376857A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Cardiac Enzymes Increased Overdose	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
SEE DOSAGE  TEXT / ORAL						

Date:11/06/02ISR Number: 4028953-4Report Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #A0376579A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Required		Convulsion	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage							

Date:11/06/02ISR Number: 4028954-6Report Type:Periodic Company Report #A0375560A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Headache Migraine	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Date:11/06/02ISR Number: 4028955-8Report Type:Periodic Company Report #A0374621A  
Age:53 YR Gender:Female I/FU:I

Outcome  
Life-Threatening  
Required

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Intervention to  
Prevent Permanent  
Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG/ PER/ DAY/ ORAL		Decreased Appetite Depression Drug Ineffective Suicidal Ideation	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

Date:11/06/02ISR Number: 4028956-XReport Type:Periodic Company Report #A0363301A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion Intentional Misuse Suicide Attempt	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride) Fluoxetine Hydrochloride (Formulation Unknown (Fluoxetine Hydrochloride) Ibuprofen (Formulation Unknown) (Ibuprofen)	PS     SS SS		

Date:11/06/02ISR Number: 4028957-1Report Type:Periodic Company Report #A0362683A  
Age:19 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required Intervention to 150 MG/ THREE Prevent Permanent TIMES PER		Convulsion	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

Impairment/Damage  
DAY/ ORAL

Clonazepam C  
Adderall C

Date:11/06/02ISR Number: 4028958-3Report Type:Periodic Company Report #A0362388A  
Age:55 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - Initial or Prolonged Required 200 MG/ TWICE Intervention to PER DAY/ ORAL Prevent Permanent Impairment/Damage	Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
			Fluvoxamine Maleate Fluoxetine Hydrochloride	C C		



Death	Death	Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Per day					
		Claritin	C		
		Monopril	C		
20MG Per day		Iud	C		

Date:11/07/02ISR Number: 4005283-8Report Type:Expedited (15-DaCompany Report #A0385531A  
 Age:38 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Chest Discomfort		Wellbutrin	PS	Glaxo Wellcome	ORAL
400MG Per day						
Initial or Prolonged	Hyperhidrosis		Effexor Xr	C		
	Palpitations		Xanax	C		
	Ventricular Tachycardia		Atarax	C		

Date:11/07/02ISR Number: 4005292-9Report Type:Expedited (15-DaCompany Report #A0385736A  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Other	Cardiac Failure		Wellbutrin	PS	Glaxo Wellcome	ORAL
	Congestive					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/02ISR Number: 4005297-8Report Type:Expedited (15-DaCompany Report #B0280070A  
Age:31 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 13 DAY Initial or Prolonged	Arthralgia	Health	Zyban	PS	Glaxo Wellcome	ORAL
	C-Reactive Protein Increased Fibrin Increased Hilar Lymphadenopathy Metastases To Lung Neoplasm Oedema Peripheral Paraesthesia Pyrexia Red Blood Cell Sedimentation Rate Increased	Professional				

Date:11/07/02ISR Number: 4005302-9Report Type:Expedited (15-DaCompany Report #B0284138A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - UNKNOWN Initial or Prolonged	Agitation		Zyban	PS	Glaxo Wellcome	
	Balance Disorder Contusion Difficulty In Walking Discomfort Dyspnoea Emotional Distress Face Oedema Fatigue Malaise Memory Impairment Oedema Peripheral Rash Generalised Rash Papular Tinnitus Urticaria					

Date:11/07/02ISR Number: 4005303-0Report Type:Expedited (15-DaCompany Report #B0284170A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspnoea		Zyban	PS	Glaxo Wellcome	ORAL
14 DAY		Haemoptysis					

Date:11/07/02ISR Number: 4005555-7Report Type:Direct Company Report #CTU 180550  
Age:41 YR Gender:Female I/FU:I

Outcome	PT
Other	Anxiety
	Breathing-Related Sleep
	Disorder
	Crying
	Depersonalisation
	Dyspnoea
	Muscle Twitching
	Pain
	Pruritus

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Sensation Of Pressure Tinnitus Vertigo	Report Source	Product	Role	Manufacturer	Route
150 MG 1X				Wellbutrin Sr 150 Mg Glaxosmithkline	PS	Glaxosmithkline	ORAL
ORAL				Claritind	C		
				Vitamins	C		
				Minerals	C		

Date:11/08/02ISR Number: 4006092-6Report Type:Expedited (15-DaCompany Report #B0281421A  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day 6 DAY	Muscle Rupture	Health	Zyban	PS	Glaxo Wellcome	ORAL
Other			Professional				

Date:11/08/02ISR Number: 4006097-5Report Type:Expedited (15-DaCompany Report #B0283515A  
Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice	Initial or Prolonged per day 64 DAY	Dizziness	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
150MCG per day		Headache		Thyroid Medication	C		ORAL
		Hypomania					
		Inner Ear Disorder					

Date:11/08/02ISR Number: 4006100-2Report Type:Expedited (15-DaCompany Report #D0039689A  
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Chest Pain	Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown 3 DAY	Dizziness	Trimipramine	C		ORAL
25MG Per day	Dyspnoea				
	Exanthem				
	Gastrointestinal Disorder				

Date:11/08/02ISR Number: 4007486-5Report Type:Direct Company Report #CTU 180626  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG 2X Initial or Prolonged DAILY ORAL		Anxiety		Wellbutrin 150 Mg.	PS		ORAL
		Chest Pain					
		Dyspnoea		Nicoderm Patch Step			
		Heart Rate Increased		1	SS		
TRANSDERMAL	1 PATCH EVERY	Hypoaesthesia					
24		Panic Disorder					
TRANSDERMAL							

Date:11/11/02ISR Number: 4006927-7Report Type:Expedited (15-DaCompany Report #A0075655A  
 Age:49 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Amnesia
Initial or Prolonged	Depression
	Dyspnoea
	Emotional Disorder
	Grand Mal Convulsion

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Status Epilepticus  
Suicidal Ideation

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	5 DAY		Zyban	PS	Glaxo Wellcome	ORAL
500MG Per day			Calcium	C		
1.25MG Per day			Ces	C		ORAL
			Vitamin C	C		

Date:11/12/02ISR Number: 4008406-XReport Type:Direct Company Report #CTU 180799  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Bupropion	PS		

Date:11/13/02ISR Number: 4008407-1Report Type:Expedited (15-DaCompany Report #A0380824A  
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dysphagia		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Per day							
Initial or Prolonged		Hypersensitivity		Birth Control Pill	C		
1TAB Per day							
Other		Oesophageal Stenosis Pain		Vitamin	C		

Date:11/13/02ISR Number: 4008409-5Report Type:Expedited (15-DaCompany Report #A0386005A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Glucose Increased		Wellbutrin	PS	Glaxo Wellcome	ORAL
300MG Per day	1 YR						

60MG Per day	Conversion Disorder	Prozac	C		ORAL
	Hypertension	Remeron	C		ORAL
45MG Per day	Lipids Increased	Synthroid	C	Glaxo Wellcome	ORAL
		Hypertension Med	C		
		Hypertension Med	C		
		Anti-Hyperlipidaemic (Unspecified)	C		

Date:11/13/02ISR Number: 4008411-3Report Type:Expedited (15-DaCompany Report #A0386169A  
Age:13 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Autoimmune Thyroiditis		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Per day	9 MON	Dysmenorrhoea					
		Headache					

Date:11/14/02ISR Number: 4009026-3Report Type:Expedited (15-DaCompany Report #A0385936A  
Age:4 DY Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Bradycardia
Other	Drug Withdrawal Syndrome
	Neonatal
	Feeling Jittery
	Maternal Drugs Affecting Foetus
	Neonatal Apnoeic Attack

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Neonatal Disorder  
Sepsis Neonatal

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
3 WK			Wellbutrin	PS	Glaxo Wellcome	
UNKNOWN			Ampicillin	C	Glaxo Wellcome	
UNKNOWN			Kanamycin	C		
			Prenatal Vitamin Iron	C C		

Date:11/14/02ISR Number: 4009028-7Report Type:Expedited (15-DaCompany Report #A0386329A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day		Grand Mal Convulsion Tic		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:11/14/02ISR Number: 4009031-7Report Type:Expedited (15-DaCompany Report #B0275736A  
Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN Initial or Prolonged	2TAB per day 9 DAY	Aggression Agitation Depression Palpitations Panic Attack		Zyban	PS	Glaxo Wellcome	

Date:11/14/02ISR Number: 4009032-9Report Type:Expedited (15-DaCompany Report #B0282085A  
Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Life-Threatening 90 DAY	Cardiac Disorder	Zyban	PS	Glaxo Wellcome	
Hospitalization - 25MG Per day	Drug Interaction	Previscan	SS		ORAL
Initial or Prolonged	Haematoma	Aspegic 100	SS		ORAL
	Iatrogenic Injury	Lasilix	C	Glaxo Wellcome	ORAL
	International Normalised Ratio Increased	Diffu K	C		ORAL
300MG per day		Sectral	C		ORAL
2.5MG Twice	Intracardiac Thrombus	Triatec	C		ORAL
per day	Pericardial Haemorrhage				
20MG Twice	Post Procedural	Vasten	C		ORAL
per day	Complication				
TRANSDERMAL		Discotriner	C	Glaxo Wellcome	

Date:11/14/02ISR Number: 4009039-1Report Type:Expedited (15-DaCompany Report #B0284793A  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Per day 18 DAY Initial or Prolonged	Atrial Fibrillation		Zyban	PS	Glaxo Wellcome	ORAL
	Depression					
	Supraventricular					
	Extrasystoles					
	Tachyarrhythmia					
	Tachycardia					
	Ventricular Extrasystoles					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/14/02 ISR Number: 4009975-6 Report Type:Direct  
 Age:24 YR Gender:Female I/FU:I

Company Report #CTU 180969

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Migraine		Wellbutrin Sr 100 Mg	PS		ORAL
100 MG BID		Nausea					
ORAL		Pyrexia					
		Vomiting					

Date:11/14/02 ISR Number: 4012907-8 Report Type:Expedited (15-DaCompany Report #2002133101US  
 Age:34 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Literature Health	Dramamine (Dimenhydrinate) Tablet	PS		ORAL
ORAL			Professional	Isotretinoin (Isotretinoin)			
				Acetaminophen (Paracetamol)	SS		
				Bupropion (Amfebutamone)	SS		

Date:11/15/02 ISR Number: 4010024-4 Report Type:Expedited (15-DaCompany Report #A0382851A  
 Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - INTRAVENOUS		Drug Abuser		Wellbutrin	PS	Glaxo Wellcome	
Initial or Prolonged		Medication Error		Seroquel	C		ORAL
100MG Per day		Phlebitis		Wellbutrin	C	Glaxo Wellcome	ORAL
200MG Per day				Lithium	C	Glaxo Wellcome	ORAL
150MG Per day							

Date:11/15/02ISR Number: 4010029-3Report Type:Expedited (15-DaCompany Report #A0386403A  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin	PS	Glaxo Wellcome	
150MG Per day		Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus					

Date:11/15/02ISR Number: 4010030-XReport Type:Expedited (15-DaCompany Report #A0386477A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Zyban	PS	Glaxo Wellcome	ORAL
1 WK							

Date:11/18/02ISR Number: 4010894-XReport Type:Expedited (15-DaCompany Report #B0284567A  
Age:53 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Blood Fibrinogen Increased C-Reactive Protein Increased Csf Glucose Abnormal

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	8 DAY	Csf Protein Abnormal Csf White Blood Cell Count Positive Grand Mal Convulsion Pyrexia Red Blood Cells Csf Positive		Zyban	PS	Glaxo Wellcome	ORAL

Date:11/18/02ISR Number: 4013271-0Report Type:Expedited (15-DaCompany Report #2002065752  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL Initial or Prolonged		Confusional State Diarrhoea Drooling Fall Hip Fracture Pelvic Fracture Weight Decreased	Consumer	Zoloft (Sertraline) Aricept (Donepezil) Bupropion Hydrochloride	PS SS SS		ORAL

Date:11/18/02ISR Number: 4014589-8Report Type:Expedited (15-DaCompany Report #B0284705A  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthralgia Blood Alkaline Phosphatase Increased Blood Bilirubin Increased Cardiac Murmur Dyspnoea Gamma-Glutamyltransferase Increased Haematuria Hypersensitivity Pericardial Effusion Urinary Casts White Blood Cells Urine Positive	Foreign Health Professional	Zyban Tablet - Zyban (Bupropion Hydrochloride)	PS		

Date:11/19/02ISR Number: 4011453-5Report Type:Expedited (15-DaCompany Report #B0280687A  
Age:60 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 7 WK	Chest Pain		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged 200MG Per day	Phlebitis		Allopurinol	C	Glaxo Wellcome	ORAL
300MG Per day	Pulmonary Embolism		Fenofibrate	C		ORAL

Date:11/19/02ISR Number: 4011708-4Report Type:Direct Company Report #CTU 181223  
Age:40 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG BID PO Initial or Prolonged Disability	Impaired Gastric Emptying		Zyban 150 Bid	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/19/02ISR Number: 4014600-4Report Type:Expedited (15-DaCompany Report #B0276683A

Age:25 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - Initial or Prolonged 150 MG / PER Disability DAY / ORAL Required Intervention to Prevent Permanent INTRA VENOUS Impairment/Damage	Aggression Agitation Anxiety Coma Conjunctival Hyperaemia Disturbance In Attention Hyperhidrosis Indifference Lung Disorder Motor Dysfunction Multi-Organ Failure Suicide Attempt Tachycardia Tremor	Foreign Health Professional Company Representative	Zyban Tablet - Zyban (Bupropion Hydrochloride) Mineral Oil Injection (Mineral Oil) Sertraline	PS SS C		ORAL

Date:11/20/02ISR Number: 4012165-4Report Type:Expedited (15-DaCompany Report #A0386665A

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	Asthenia Blindness Unilateral Hypoaesthesia		Wellbutrin Metadate Cd Effexor Xr Geodon	PS C C C	Glaxo Wellcome	ORAL

Date:11/20/02ISR Number: 4028995-9Report Type:Periodic Company Report #001-0981-M0108765

Age:55 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization -	Eosinophilia	Health	Atorvastatin			

Initial or Prolonged	Rash	Professional	(Atorvastatin)	PS	ORAL
10 MG					
(DAILY), ORAL			(Bupropion)	SS	ORAL
ORAL			Timolol Maleate	C	
			Sulindac	C	
			Ibuprofen	C	

Date:11/21/02ISR Number: 4012726-2Report Type:Expedited (15-DaCompany Report #A0386663A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Libido Decreased		Paxil	PS	Glaxo Wellcome	ORAL
60MG Per day		Retinal Disorder		Wellbutrin	SS	Glaxo Wellcome	ORAL
100MG Per day							

Date:11/21/02ISR Number: 4012737-7Report Type:Expedited (15-DaCompany Report #B0285200A  
Age: Gender:Female I/FU:I

Outcome	PT
Other	Emotional Disorder
	Feeling Hot
	Insomnia
	Lethargy
	Pain
	Tinnitus

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG			Zyban	PS	Glaxo Wellcome	ORAL
Variable dose			Esomeprazole	C		ORAL
20MG Unknown			Ethinylestradiol + Levonorgestrel	C		ORAL
1TAB per day						

Date:11/21/02ISR Number: 4013304-1Report Type:Direct Company Report #CTU 181490  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Crying		Bupropion Hcl - 100			
Other		Depression		Mg	PS		
1 TAB		Drug Effect Decreased					
EVERYDAY; 1		Feeling Abnormal					
TAB TWICE A		Libido Decreased					
DAY							

Date:11/22/02ISR Number: 4013823-8Report Type:Expedited (15-DaCompany Report #A0386899A  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Optic Neuritis		Wellbutrin	PS	Glaxo Wellcome	ORAL
				Celexa	SS		ORAL

Date:11/25/02ISR Number: 4014171-2Report Type:Periodic Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-11850310  
 Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Pancreatitis Acute	Health Professional	Sustiva Caps 600 Mg Stavudine	PS SS	Bristol-Myers Squibb Company Bristol-Myers Squibb Company	ORAL ORAL
unknown start							
date-prior to							
06-Dec-00				Videx Ec Caps	SS	Bristol-Myers Squibb Company	ORAL
at hour of							
sleep				Wellbutrin	SS		ORAL
AM + 1600 x							
30 days							

Date:11/25/02ISR Number: 4014390-5Report Type:Expedited (15-DaCompany Report #B0285487A  
Age:23 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Amnesia Anxiety Blood Creatine Phosphokinase Increased Confusional State Convulsion Dermabrasion Disorientation Ecchymosis Haematoma Incoherent

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Intentional Misuse Irritability Mydriasis	Report Source	Product	Role	Manufacturer	Route
1 WK		Suicidal Ideation		Zyban	PS	Glaxo Wellcome	ORAL
		Suicide Attempt Urine Amphetamine Positive					

Date:11/25/02ISR Number: 4014438-8Report Type:Expedited (15-DaCompany Report #WAES 0211USA01406  
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Interaction Heat Stroke		Cogentin Wellbutrin Neurontin	PS SS SS	Merck & Co., Inc	

Date:11/25/02ISR Number: 4017279-0Report Type:Expedited (15-DaCompany Report #CEL-2002-01391-ROC(0)  
Age:14 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Grand Mal Convulsion	Foreign Literature Health Professional	Methylphenidate Tablets (Unspecified) (Methylphenidate Hydrochloride)	PS		
60 MG (UNK, DAILY)				Bupropion (Bupropion)	SS		
SEE IMAGE	4 WK						

Date:11/26/02ISR Number: 4015319-6Report Type:Expedited (15-DaCompany Report #A0378902A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Congenital Anomaly 300MG Per day	Bronchitis	Health	Wellbutrin	PS	Glaxo Wellcome
	Coarctation Of The Aorta Complications Of Maternal	Professional	Albuterol Inhaler Depakote	SS SS	Glaxo Wellcome
500MG Per day	Exposure To Therapeutic Drugs		Synthroid	SS	Glaxo Wellcome
.175MG Per day	Maternal Drugs Affecting Foetus Pregnancy		Effexor Xr Trimox Niferex	SS SS SS	Glaxo Wellcome
150MG Per day	Premature Baby		Zithromax	SS	
1TAB Per day	Ventricular Septal Defect Acquired		Phenergan Vc With Codeine	SS	
250MG Per day 5 2TSP As required	DAY 8 DAY				

Date:11/26/02ISR Number: 4015329-9Report Type:Expedited (15-DaCompany Report #A0387263A  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypersensitivity		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Per day 3	DAY	Multiple Sclerosis Oral Mucosal Blistering Swelling Throat Tightness		Betaseron Ditropan Baclofen	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/26/02ISR Number: 4015340-8Report Type:Expedited (15-DaCompany Report #B0285464A  
Age:32 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Retinal Artery Thrombosis		Zyban	PS	Glaxo Wellcome	
UNKNOWN	150MG Twice					
Initial or Prolonged	Visual Disturbance					
per day	6 DAY					

Date:11/26/02ISR Number: 4015341-XReport Type:Expedited (15-DaCompany Report #B0285584A  
Age:47 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Depressed Level Of		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	31 DAY					
Initial or Prolonged	Consciousness		Humalog Insulin	C		
UNKNOWN	Dysstasia		Stagid	C		ORAL
	Malaise		Nicotine Patch	C	Glaxo Wellcome	
TRANSDERMAL						
	Muscular Weakness		Alcohol	C		ORAL
1 DAY	Paraplegia					
	Sensory Loss					

Date:11/26/02ISR Number: 4017903-2Report Type:Expedited (15-DaCompany Report #US010405  
Age:17 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Coma	Health	Gabitril	PS		
Initial or Prolonged	Road Traffic Accident	Professional	Wellbutrin	SS		
	Status Epilepticus					
	Tonic Clonic Movements					

Date:11/26/02ISR Number: 4017971-8Report Type:Expedited (15-DaCompany Report #12110755  
Age:49 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Warfarin (Warfarin			
Other		Drug Level Above Therapeutic Overdose	Health Professional	Sodium) Carbamazepine Bupropion (Bupropion Hcl)	PS SS SS		

Date:11/26/02ISR Number: 4018186-XReport Type:Expedited (15-DaCompany Report #2002066467  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Deafness Neurosensory	Health	Zoloft (Sertraline)	PS		ORAL
175 MG		Tinnitus	Professional				
(DAILY), ORAL				Bupropion	SS		ORAL
300 MG (TID),							
ORAL				Fexofenadine Hydrochloride	C		
				Fluticasone Propionate	C		
				All Other Therapeutic Products	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/27/02ISR Number: 4019168-4Report Type:Expedited (15-DaCompany Report #SKEL000128

Age:30 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Death	Cardiac Arrest	Health	Skelaxin			
Hospitalization -	Convulsion	Professional	(Metaxalone)	PS		ORAL
ORAL						
Initial or Prolonged	Disseminated	Distributor	Tylenol			
ORAL	Intravascular Coagulation		(Acetaminophen)	SS		ORAL
	Hepatic Failure		Tylenol Pm, Extra			
ORAL	Mydriasis		Strength			
	Nystagmus		(Acetaminophen D	SS		ORAL
	Posturing		Effexor			
ORAL	Pupil Fixed		(Venlafaxine)	SS		ORAL
	Pyrexia		Wellbutrin Sr			
ORAL	Renal Failure		(Bupropion)	SS		ORAL
	Rhabdomyolysis		Librium			
ORAL	Tachycardia		(Chlordiazepoxide)	SS		ORAL
			Ambien (Zolpidem)	SS		ORAL
ORAL						
			Midol			
ORAL			(Apap/Pyrilamine/Caf			
			feine)	SS		ORAL
			Pyridium			
ORAL			(Phenazopyridine)	SS		ORAL
			Aleve (Naproxen)	SS		ORAL
ORAL						
			Relafen (Nabumetone)	SS		

Date:11/29/02ISR Number: 4018288-8Report Type:Expedited (15-DaCompany Report #2002003791

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Colitis Ulcerative	Consumer	Zoloft (Sertraline)	PS		
Initial or Prolonged	Depression		Mirtazapine	SS		
Other	Flashback		Bupropion			

Gallbladder Disorder  
Gastric Ulcer  
Gastritis Erosive  
Loss Of Consciousness  
Pain  
Panic Attack  
Restless Legs Syndrome  
Serotonin Syndrome  
Suicidal Ideation  
Weight Decreased

Hydrochloride SS  
All Other  
Therapeutic Products SS  
Anti-Parkinson  
Agents SS

Date:11/29/02ISR Number: 4019863-7Report Type:Expedited (15-DaCompany Report #200213918BCC  
Age:30 YR Gender:Male I/FU:F

Outcome PT  
Death Body Temperature  
Hospitalization - Increased  
Initial or Prolonged Cardio-Respiratory Arrest  
Completed Suicide  
Convulsion  
Disseminated  
Intravascular Coagulation  
Hepatic Failure  
Nystagmus  
Posturing  
Pupil Fixed  
Renal Failure

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Rhabdomyolysis Tachycardia	Report Source	Product	Role	Manufacturer	Route
ORAL			Literature Health	Aleve (Naproxen Sodium)	PS		ORAL
			Professional Other	Tylenol Pm Effexor (Venlafaxine Hydrochloride) Wellbutrin Sr (Amfebutamone Hydrochloride) Librium Ambien (Zolpidem Tartrate) Skelaxin (Metaxalone) Pyridium (Phenazopyridine Hydrochloride) Tylenol (Paracetamol) Relafen (Nabumetone) Midol	SS SS SS SS SS SS SS SS SS SS SS SS		

Date:12/02/02ISR Number: 4017532-0Report Type:Expedited (15-DaCompany Report #A0387586A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Intentional Misuse Suicide Attempt		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:12/02/02ISR Number: 4018349-3Report Type:Direct Company Report #CTU 181873  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Agitation Heart Rate Increased Hypoaesthesia		Wellbutrin Sr 150mg (Bupropion Hydrochloride)	PS		

1 TWICE A DAY

Oedema Peripheral  
Pain  
Pruritus  
Rash

Date:12/03/02ISR Number: 4018226-8Report Type:Expedited (15-DaCompany Report #A0387654B

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Congenital Anomaly		Maternal Drugs Affecting Foetus Necrotising Colitis Pregnancy Premature Baby Pulmonary Valve Stenosis Congenital Vacuum Extractor Delivery		Bupropion	PS	Glaxo Wellcome	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/03/02ISR Number: 4018236-0Report Type:Expedited (15-DaCompany Report #B0134604A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Discomfort Insomnia Oesophageal Spasm Urticaria		Zyban Mercilon	PS C	Glaxo Wellcome	ORAL ORAL

Date:12/03/02ISR Number: 4018237-2Report Type:Expedited (15-DaCompany Report #B0259750A  
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG See Initial or Prolonged dosage text	32 DAY	Back Pain  Bronchitis Chronic C-Reactive Protein Increased Hodgkin'S Disease Neutrophil Count Increased Red Blood Cell Sedimentation Rate Increased		Zyban	PS	Glaxo Wellcome	ORAL

Date:12/03/02ISR Number: 4021391-XReport Type:Expedited (15-DaCompany Report #2002107579GB  
Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Drug Interaction Lumbar Vertebral Fracture	Foreign Literature Health  Professional  Other	Depo-Medrone (Methylp rednisolone) Suspension, Sterile	PS		
INTRA-ARTICULAR	30 MG,						
SINGLE,							
INTRA-ARTICUL							

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	160 MG (BID),		Bundle Branch Block	Health Professional	Geodon (Ziprasidone)	PS		ORAL
					Bupropion Hydrochloride	SS		ORAL
	200 MG (BID),				Lorazepam	C		

Date:12/03/02ISR Number: 4025895-5Report Type:Periodic Company Report #2002061599  
Age:54 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	160 MG (BID),		Bundle Branch Block	Health Professional	Geodon (Ziprasidone)	PS		ORAL
					Bupropion Hydrochloride	SS		ORAL
	200 MG (BID),				Lorazepam	C		

Date:12/04/02ISR Number: 4018742-9Report Type:Expedited (15-DaCompany Report #A0385736A  
Age: Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Cardiac Failure Congestive	Health Professional	Wellbutrin	PS	Glaxo Wellcome	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/04/02ISR Number: 4018748-XReport Type:Expedited (15-DaCompany Report #A0387850A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Drug Ineffective		Zyban	PS	Glaxo Wellcome	ORAL

Date:12/04/02ISR Number: 4019467-6Report Type:Direct

Company Report #CTU 182169

Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG PO BID		Bradycardia		Bupropion	PS		ORAL
Initial or Prolonged 75MG PO BID				Metoprolol	SS		ORAL

[GREATER THAN

1 YR]

Date:12/04/02ISR Number: 4019473-1Report Type:Direct

Company Report #CTU 182171

Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Convulsion		Wellbutrin	PS		
				St John'S Wart	SS		
				Colostrum	SS		

Date:12/06/02ISR Number: 4022598-8Report Type:Expedited (15-DaCompany Report #B0285143A

Age:34 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Intentional Misuse	Literature Health Professional	Diphenhydramine Hydrochloride (Formulation Unknown)			

ORAL				(Diphenhydramine	PS		ORAL
				Isotretinoin (Formulation Unknown) (Isotretinoin)	SS		ORAL
ORAL				Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	SS		

Date:12/06/02ISR Number: 4022600-3Report Type:Expedited (15-DaCompany Report #B0285160A  
Age:33 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Literature Health Professional	Diphenhydramine Hydrochloride (Formulation Unknown) (Diphenhydramine	PS		ORAL
ORAL				Risperidone (Formulation Unknown) (Risperidone)	SS		ORAL
ORAL				Wellbutrin (Formulation Unknown) (Bupropion			

Freedom Of Information (FOI) Report

ORAL Hydrochloride) SS ORAL

Date:12/09/02ISR Number: 4020258-0Report Type:Expedited (15-DaCompany Report #B0265019A  
 Age:60 YR Gender:Male I/FU:F

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day 30 DAY	Asthenia	Health	Zyban	PS	Glaxo Wellcome	ORAL
20MG per day	Convulsion	Professional	Simvastatin	C		
200MG per day	Dizziness		Seloken	C		
10MG per day	Fatigue		Renitec	C		
40MG per day	Fear		Frusemide	C	Glaxo Wellcome	
75MG per day	Gastrointestinal Disorder		Aspirin	C		
	General Physical Health Deterioration Loss Of Consciousness Malaise Memory Impairment Pancreatitis Panic Attack Peritonitis Purulence Sexual Dysfunction Sleep Disorder Swelling					

Date:12/09/02ISR Number: 4020263-4Report Type:Expedited (15-DaCompany Report #B0283974A  
 Age:51 YR Gender:Male I/FU:F

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 29 DAY	Aphonia	Health	Zyban	PS	Glaxo Wellcome	ORAL
20MG Per day	Dysphonia	Professional	Tahor	C		ORAL

Laryngitis

Date:12/09/02ISR Number: 4020264-6Report Type:Expedited (15-DaCompany Report #B0284170A  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Lung Neoplasm Malignant	Health	Zyban	PS	Glaxo Wellcome	ORAL
14 DAY			Professional				

Date:12/09/02ISR Number: 4020278-6Report Type:Expedited (15-DaCompany Report #B0287228A  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Emotional Distress		Zyban	PS	Glaxo Wellcome	ORAL
150MG Unknown	21 DAY	Nausea		Mirena	C		
INTRA-UTERINE		Throat Tightness					
		Urticaria					
		Vomiting					
		Wheezing					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/09/02ISR Number: 4020872-2Report Type:Direct  
Age:22 YR Gender:Male I/FU:I

Company Report #CTU 182333

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 1 PILL A.M. Initial or Prolonged ORAL	Convulsion		Welbutrin 100mg Sr Glaxo	PS	Glaxo	ORAL
1 PILL A.M.  ORAL			Wellbutrin 150mg Sr Glaxo	SS	Glaxo	ORAL

Date:12/09/02ISR Number: 4020922-3Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #182415

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration	Arthralgia Blood Pressure Systolic Increased Excoriation Heart Rate Increased Hypersensitivity Oedema Peripheral Respiratory Disorder Respiratory Rate Increased Serum Sickness Throat Tightness		Zyban	PS		

Date:12/10/02ISR Number: 4020896-5Report Type:Expedited (15-DaCompany Report #B0273556A  
Age:53 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Per day 12 DAY Initial or Prolonged 800UG Per day	Supraventricular  Tachycardia		Zyban  Rivotril	PS  C	Glaxo Wellcome	ORAL  ORAL

Date:12/10/02ISR Number: 4020905-3Report Type:Expedited (15-DaCompany Report #B0286752A

Age:34 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Twice	Diabetes Mellitus		Zyban	PS	Glaxo Wellcome	ORAL
	per day	Inadequate Control					
	40MG per day			Frusemide	C	Glaxo Wellcome	
				Porcine Insuline	C		
				Warfarin	C	Glaxo Wellcome	
				Insulin	C		
				Ramipril	C		
	5MG per day						

Date:12/10/02ISR Number: 4020907-7Report Type:Expedited (15-DaCompany Report #B0286803A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	10 DAY	Depersonalisation		Zyban	PS	Glaxo Wellcome	ORAL
	MON	Feeling Abnormal		Deroxat	C	Glaxo Wellcome	ORAL
		Insomnia					
		Nausea					
		Tearfulness					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/10/02ISR Number: 4021777-3Report Type:Direct  
Age:36 YR Gender:Female I/FU:I

Company Report #CTU 182455

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG 2X PER		Dizziness Dyspnoea Erythema Feeling Abnormal Flushing Hypersensitivity Hypoaesthesia Palpitations Paraesthesia Tongue Oedema		Wellbutrin 150mg Tablets Glaxo	PS	Glaxo	ORAL

Date:12/10/02ISR Number: 4024748-6Report Type:Expedited (15-DaCompany Report #HYDROCOD2002-00427  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death PO		Acute Pulmonary Oedema	Literature	Hydrocodone/Apap	PS		ORAL
PO		Completed Suicide	Health	Propoxyphene	SS		ORAL
PO		Drug Screen Positive	Professional	Bupropion	SS		ORAL
PO		Overdose		Doxepin	SS		ORAL
PO		Pulmonary Congestion		Fluoxetine	SS		ORAL
PO				Zolpidem	SS		ORAL

Date:12/10/02ISR Number: 4024802-9Report Type:Expedited (15-DaCompany Report #02P-163-0205266-00  
Age:42 YR Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Completed Suicide	Literature Health	Hydrocodone/Acetamin ophen (Vicodin)			

Professional

(Hydrocodone/Acetami

nophen)

(Hydrocodone/Acetami PS

Dextropropoxyphene SS

Bupropion SS

Date:12/11/02ISR Number: 4021400-8Report Type:Expedited (15-DaCompany Report #A0388692A

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 5 DAY	Burning Sensation		Wellbutrin	PS	Glaxo Wellcome	
Initial or Prolonged	Photosensitivity Reaction					

Date:12/11/02ISR Number: 4021401-XReport Type:Expedited (15-DaCompany Report #B0111947A

Age:39 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150MG Twice	Asthenia		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day 7 DAY	Bundle Branch Block					
	Chest Pain		Zopiclone	C		
	Dizziness		Propiomazine	C		
	Palpitations		Somadril	C		
	Pulmonary Embolism					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/11/02ISR Number: 4021402-1Report Type:Expedited (15-DaCompany Report #B0286866A  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State		Zyban	PS	Glaxo Wellcome	
150MG Unknown							
		Fear		Aspirin	C		ORAL
75MG Unknown							
		Hyperventilation		Glyceryl Trinitrate	C	Glaxo Wellcome	
SUBLINGUAL							
		Malaise		Salbutamol	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)	2PUFF Unknown						

Date:12/12/02ISR Number: 4021689-5Report Type:Expedited (15-DaCompany Report #A0388553A  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Drug Interaction		Wellbutrin	PS	Glaxo Wellcome	
150MG Twice							
		Pneumonia					
per day							
				Biaxin	C		ORAL
500MG Twice							
per day							
				Prednisone	C		ORAL
				Avandia	C	Glaxo Wellcome	ORAL
4MG Per day							
SUBCUTANEOUS				Humulin R	C		
SUBCUTANEOUS				Humulin N	C		
				Lorcet 10	C		ORAL
				Xanax	C		ORAL
				Diflucan	C		ORAL
200MG Per day							
				Lithium	C	Glaxo Wellcome	ORAL
900MG Per day							
				Paxil	C	Glaxo Wellcome	ORAL
40MG Per day							

600MG Per day	Neurontin	C		ORAL
30MG Per day	Prevacid	C		ORAL
RESPIRATORY (INHALATION)	Albuterol	C	Glaxo Wellcome	
RESPIRATORY (INHALATION)	Atrovent	C		
RESPIRATORY (INHALATION)	Flovent	C	Glaxo Wellcome	

Date:12/12/02ISR Number: 4021701-3Report Type:Expedited (15-DaCompany Report #B0287227A  
Age:52 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150MG Per day	Arterial Spasm	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
Other UNKNOWN	Drug Interaction 30ML per day 1 DAY		Iopamidol	SS		
75MG per day	Electrocardiogram St		Aspirin	C		ORAL
10MG per day	Segment Elevation		Felodipine	C		ORAL
20MG per day	Nausea		Atorvastatin	C		ORAL
20MG per day	Vomiting		Rabeprazole	C		ORAL

Date:12/12/02ISR Number: 4021702-5Report Type:Expedited (15-DaCompany Report #B0287325A  
Age:50 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1 MON	Amnesia		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	Anxiety Confusional State Depression					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/12/02ISR Number: 4021724-4Report Type:Expedited (15-DaCompany Report #B0260318A  
 Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2TABS per day 3 DAY Initial or Prolonged		Angina Unstable	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
		Chest Pain					
		Coronary Artery Occlusion					
		Coronary Artery Stenosis					
		Diabetes Mellitus					
		Rash Erythematous					
		Rash Pruritic					

Date:12/12/02ISR Number: 4021725-6Report Type:Expedited (15-DaCompany Report #B0281421A  
 Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Per day 6 DAY Other		Dysstasia	Health	Zyban	PS	Glaxo Wellcome	ORAL
		Muscle Rupture	Professional				
		Myalgia					

Date:12/12/02ISR Number: 4021730-XReport Type:Expedited (15-DaCompany Report #B0287403A  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG per day		Galactorrhoea		Zyntabac	PS	Glaxo Wellcome	ORAL

Date:12/12/02ISR Number: 4021731-1Report Type:Expedited (15-DaCompany Report #D0039257A  
 Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 4TAB per day 2 DAY Hospitalization - Initial or Prolonged		Blood Bicarbonate	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
		Decreased		Metformin	SS		ORAL
		Blood Glucose Increased					

Other

Blood Ph Decreased  
Blood Pressure Increased  
Circulatory Collapse  
Communication Disorder  
Convulsion  
Disturbance In Attention  
General Physical Health  
Deterioration  
Lactic Acidosis  
Overdose  
Refusal Of Treatment By  
Patient  
Sinus Tachycardia  
White Blood Cell Count  
Increased

Date:12/12/02ISR Number: 4025484-2Report Type:Expedited (15-DaCompany Report #02P-163-0205719-00  
Age:38 YR Gender:Male I/FU:I

Outcome PT  
Hospitalization - Arthralgia  
Initial or Prolonged Difficulty In Walking  
Lymphadenopathy  
Myalgia  
Oral Intake Reduced

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Weight Decreased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
6 TABLET, 1 IN 1 D, PER ORAL		Consumer	Kaletra Soft Gelatin Capsules (Kaletra) (Lopinavir/ Ritonavir)(Lopinavir / Ritonavir)	PS		ORAL
150 MG , 2 IN 1 D, PER ORAL			Bupropion Hydrochloride	SS		ORAL
			Tenofovir	C		
			Lamivudine	C		
			Lexapro	C		

Date:12/13/02ISR Number: 4022654-4Report Type:Expedited (15-DaCompany Report #A0388881A  
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 5 DAY Initial or Prolonged		Asthenia Choreoathetosis Coordination Abnormal		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:12/13/02ISR Number: 4022655-6Report Type:Expedited (15-DaCompany Report #A0388982A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 400MG Per day YR		Blood Pressure Increased Hepatic Neoplasm		Wellbutrin	PS	Glaxo Wellcome	ORAL
				Effexor	C		
				Birth Control	C		
				Atenolol	C		



Date:12/13/02ISR Number: 4022657-XReport Type:Expedited (15-DaCompany Report #A0389005A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Breast Cancer Female		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:12/13/02ISR Number: 4027164-6Report Type:Expedited (15-DaCompany Report #02P-163-0205470-00  
 Age:19 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardio-Respiratory Arrest Completed Suicide	Literature Health Professional	Valproic Acid (Depakene) (Valproic Acid) (Valproic Acid)	PS		ORAL
PER ORAL				Bupropion	SS		ORAL
PER ORAL				Olanzapine	SS		ORAL
PER ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/13/02ISR Number: 4027237-8Report Type:Expedited (15-DaCompany Report #S02-USA-03225-01

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Optic Neuritis	Health Professional	Celexa (Citalopram Hydrobromide)	PS		
				Wellbutrin Sr (Bupropion Hydrochloride)	SS		

Date:12/17/02ISR Number: 4026091-8Report Type:Direct

Company Report #CTU 182820

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Conversion Disorder		Wellbutrin 200mg	PS		ORAL
200MG TWICE A							
		Crying					
DAY ORAL		Memory Impairment					

Date:12/18/02ISR Number: 4025589-6Report Type:Direct

Company Report #CTU 182981

Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Loss Of Consciousness		Bupropion	PS		
300MG QD							
Initial or Prolonged		Petit Mal Epilepsy					
CHRONIC		Road Traffic Accident		Amitriptyline	SS		ORAL
75MG PO QHS							
CHRONIC				Labetolol	C		
				Altace	C		
				Levoxyl	C		
				Norvasc	C		
				Wellbutrin	C		

Date:12/18/02ISR Number: 4029418-6Report Type:Expedited (15-DaCompany Report #A0387790A  
Age:14 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State	Foreign	Wellbutrin			
		Drug Interaction	Literature	Unspecified Tablet			
		Grand Mal Convulsion	Health	(Bupropion			
		Somnolence	Professional	Hydrochloride)	PS		ORAL
PER DAY/ORAL				Methylphenidate			
				(Formulation Unknown			
				) (Methylphenidate)	SS		
60 MG/PER DAY							

Date:12/18/02ISR Number: 4029817-2Report Type:Expedited (15-DaCompany Report #B0287094A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Condition Aggravated	Foreign	Zyban Tablet-Zyban			
Initial or Prolonged		Drug Dependence	Literature	(Bupropion			
		Myocardial Infarction	Health	Hydrochloride)	PS		
		Tobacco Abuse	Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/19/02ISR Number: 4025667-1Report Type:Expedited (15-DaCompany Report #A0386329A  
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice		Grand Mal Convulsion	Professional				
per day		Tic		Klonopin	C		ORAL
1MG Three							
times per day				Lorcet	C		

Date:12/19/02ISR Number: 4025708-1Report Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-12135786  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Condition Aggravated	Health	Aripiprazole	PS	Otsuka	
Initial or Prolonged		Schizoaffective Disorder	Professional			Pharmaceutical Company, Ltd.	ORAL
				Wellbutrin	SS		
				Prozac	SS		
				Topamax	C		
				Klonopin	C		

Date:12/19/02ISR Number: 4026267-XReport Type:Direct Company Report #CTU 183081  
Age:28 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
1T PO QD X3D		Agitation		Zyban 180mg Tbsr	PS		ORAL
THEN 1T PO		Insomnia					
BID				Nicorette	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 50MG PO QID		Chills Coma		Tramadol 50mg Qid Started 12/9	PS		ORAL
Other SUBCUTANEOUS	6 MG X	Confusional State SQ X		Imitrex 6mg Sq X 1	SS		
20 MG PO QD		Delirium		Paxil 20mg	SS		ORAL
150 MG PO BID		Hypotension		Zyban 150 Mg	SS		ORAL
TOPICAL	21G TOP QD	Labile Blood Pressure		Nicotine Patch 21	SS		
		Muscle Spasms Pyrexia Restlessness Tremor					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Respiratory Failure	Health Professional Company Representative	Geodon (Ziprasidone) Zoloft (Sertraline) Wellbutrin (Bupropion Hydrochloride)	PS SS  SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/23/02ISR Number: 4027471-7Report Type:Expedited (15-DaCompany Report #A0375990A  
Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day			Health	Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG See dosage text	3	MON	Professional				
			Confusional State	Zyban	SS	Glaxo Wellcome	ORAL
			Convulsion				
			Embolism	Zocor	C		
			Fracture	Stool Softener	C		
			Headache				

Date:12/23/02ISR Number: 4027475-4Report Type:Expedited (15-DaCompany Report #A0389501A  
Age:15 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 5 MON			Haemothorax	Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:12/23/02ISR Number: 4027476-6Report Type:Expedited (15-DaCompany Report #A0389718A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Renal Failure	Wellbutrin	PS	Glaxo Wellcome	ORAL
				Trilafon	C	Glaxo Wellcome	
				Zoloft	C		
				Unknown	C		

Date:12/23/02ISR Number: 4027488-2Report Type:Expedited (15-DaCompany Report #B0288015A  
Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other 150MG Twice		Dizziness		Zyban	PS	Glaxo Wellcome	ORAL
per day	7	DAY	Exophthalmos				
			Face Oedema	Ethinylloestradiol +			
			Malaise	Levonorgestrel	C		ORAL
			Migraine				
			Visual Disturbance				

Date:12/24/02ISR Number: 4033557-3Report Type:Expedited (15-DaCompany Report #B0287475A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Arthralgia	Foreign	Zyban Tablet-Zyban			
Intervention to		Influenza Like Illness		(Bupropion			
Prevent Permanent		Myalgia		Hydrochloride)	PS		ORAL
150 MG/ ORAL							
Impairment/Damage							

Date:12/24/02ISR Number: 4033559-7Report Type:Expedited (15-DaCompany Report #B0287549A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Suicidal Ideation	Foreign	Zyban Tablet-Zyban			
Intervention to		Suicide Attempt		(Bupropion			
Prevent Permanent				Hydrochloride)	PS		ORAL
ORAL							
Impairment/Damage				Paroxetine			
				Hydrochloride	C		
				Alprazolam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/24/02ISR Number: 4033566-4Report Type:Expedited (15-DaCompany Report #B0126002A  
Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG/AS DIRECTED/ORAL	Angioneurotic Oedema Toxic Skin Eruption	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:12/24/02ISR Number: 4033568-8Report Type:Expedited (15-DaCompany Report #B0134062A  
Age:24 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG/AS DIRECTED/ORAL	Cerebrovascular Stenosis Convulsion	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
			Diazepam (Diazepam) Ethanol	SS C		

Date:12/24/02ISR Number: 4033757-2Report Type:Expedited (15-DaCompany Report #S02-USA-03223-01  
Age:19 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death	Completed Suicide Intentional Misuse	Health Professional Company Representative	Celexa (Citalopram Hydrobromide) Celexa (Citalopram Hyrobromide) Lamictal (Lamotrigine) Wellbutrin (Bupropion Hydrochloride)	PS SS SS SS		



Date:12/24/02ISR Number: 4034046-2Report Type:Expedited (15-DaCompany Report #B0287207A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Arthralgia	Foreign Health	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
Intervention to Prevent Permanent ORAL Impairment/Damage		Myalgia Pruritus Purpura Rash Pruritic	Professional				

Date:12/24/02ISR Number: 4034295-3Report Type:Expedited (15-DaCompany Report #A0388161A  
Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse	Health Professional	Lamictal Unspecified Tablet (Lamotrigine)	PS		ORAL
ORAL			Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	SS		ORAL
ORAL				Citalopram Hydrobromide (Citalopram Hydrobromide)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/24/02ISR Number: 4034575-1Report Type:Expedited (15-DaCompany Report #B0287627A  
 Age:38 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150 MG / SEE DOSAGE TEXT / ORAL	Hyperhidrosis Insomnia Irritability Nephrolithiasis Renal Colic	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:12/24/02ISR Number: 4036242-7Report Type:Periodic Company Report #A0100574A  
 Age:46 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Other Required 100 MG / Intervention to TWICE PER DAY Prevent Permanent / ORAL Impairment/Damage	Dyskinesia Tardive Dyskinesia	Health Professional Company Representative	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL

Date:12/24/02ISR Number: 4036247-6Report Type:Periodic Company Report #A0383258A  
 Age:39 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other ORAL 10 MG / PER	Convulsion	Health Professional Other	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride) Lexapro (Formulation Unknown) (Lexapro)	PS SS		ORAL ORAL

DAY / ORAL

Fluoxetine  
Hydrochloride  
(Formulation  
Unknown) (Fluoxetine  
Hydrochloride) SS

ORAL

80 MG / PER

DAY / ORAL

Fluoxetine  
Hydrochloride  
(Formulation  
Unknown) (Fluoxetine  
Hydrochloride) SS

ORAL

40 MG / PER

DAY / ORAL

Date:12/24/02ISR Number: 4036254-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #A0379765A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hypertension	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/24/02ISR Number: 4036259-2Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #A0376644A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Required ORAL Intervention to Prevent Permanent Impairment/Damage	Grand Mal Convulsion	Health Professional	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL

Date:12/24/02ISR Number: 4036277-4Report Type:Periodic  
 Age:25 YR Gender:Female I/FU:I

Company Report #A0372838A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Convulsion	Consumer	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL

Date:12/24/02ISR Number: 4036278-6Report Type:Periodic  
 Age:39 YR Gender:Male I/FU:I

Company Report #A0369927A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100 MG / Initial or Prolonged TWICE PER DAY / ORAL ORAL SEE DOSAGE TEXT / ORAL	Grand Mal Convulsion	Health Professional Other	Wellbutrin  Ethanol Halls  Gabapentin Oxcarbazepine	PS  SS SS  C C		ORAL  ORAL  ORAL

Date:12/24/02ISR Number: 4036279-8Report Type:Periodic  
Age:61 YR Gender:Male I/FU:I

Company Report #A0174498A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Required 100 MG / Intervention to TWICE PER DAY Prevent Permanent / ORAL Impairment/Damage	Grand Mal Convulsion	Health Professional Company Representative	Wellbutrin Tablet (Bupropion Hydrochloride)  Simvastatin Doxazosin Mesylate Aspirin Meclozine Hydrochloride	PS  C C C C		ORAL

Date:12/24/02ISR Number: 4036280-4Report Type:Periodic  
Age:19 YR Gender:Male I/FU:I

Company Report #A0169662A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged  150 MG / SEE  DOSAGE TEXT /  ORAL	Convulsion Overdose Suicide Attempt	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

DAY

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/27/02ISR Number: 4034649-5Report Type:Expedited (15-DaCompany Report #A0389722A

Age:5 MON Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Study	Wellbutrin Sr			
Other		Drug Exposure Via Breast Milk	Health Professional	Tablet-Controlled Release (Bupropion Hydrochloride)	PS		
TRANSMAMMARY	TRANSMAMMARY	Foetus		Fluvoxamine Meleate (Fluvoxamine Maleate)	SS		
TRANSMAMMARY	TRANSMAMMARY						

Date:12/27/02ISR Number: 4035526-6Report Type:Expedited (15-DaCompany Report #B0287652A

Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Systolic Increased	Foreign	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
Other		Flushing					
ORAL		Malaise Nausea Tremor					

Date:12/30/02ISR Number: 4032147-6Report Type:Expedited (15-DaCompany Report #A0389949A

Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
Other		Loss Of Consciousness		Birth Control Pill	C		
450MG per day	2 WK	Tremor		Flexeril	C		
10MG per day				Losec	C		ORAL
20MG per day							

Date:12/30/02ISR Number: 4032148-8Report Type:Expedited (15-DaCompany Report #A0390002A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice		Difficulty In Walking					
per day		Dysarthria					
		Vision Blurred					

Date:12/30/02ISR Number: 4032161-0Report Type:Expedited (15-DaCompany Report #B0288380A  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	6 DAY	Haematuria		Bisoprolol	C		
		Leukocyturia		Hydrochlorothiazide	C		
		Malaise		Amlodipine	C		
		Proteinuria		Atorvastatin	C		
		Renal Failure Acute					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/30/02ISR Number: 4032162-2Report Type:Expedited (15-DaCompany Report #B0288386A  
Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	9TAB per day	Headache		Quomem	PS	Glaxo Wellcome	ORAL
Initial or Prolonged UNKNOWN		Hyperhidrosis		Alprazolam	C		
UNKNOWN		Nausea		Amitriptyline	C		
UNKNOWN		Overdose		Clonidine	C		
		Palpitations					
		Vomiting					

Date:12/30/02ISR Number: 4036494-3Report Type:Expedited (15-DaCompany Report #2002070646  
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	160 MG, ORAL	Fall	Health	Geodon (Ziprasidone)	PS		ORAL
ORAL		Respiratory Failure	Professional	Zoloft (Sertraline)	SS		ORAL
		Toxicologic Test Abnormal	Company Representative	Wellbutrin (Bupropion Hydrochloride)	SS		
				Obetrol	C		
				Risperidone	C		

Date:12/30/02ISR Number: 4036588-2Report Type:Expedited (15-DaCompany Report #B0286165A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Coordination Abnormal	Foreign Health	Zyban Tablet-Zyban (Bupropion Hydrochloride)			
SEE DOSAGE		Eye Movement Disorder	Professional		PS		ORAL
TEXT/ORAL		Impaired Driving Ability					
		Insomnia					



Speech Disorder  
Vertigo

Date:12/31/02ISR Number: 4036699-1Report Type:Expedited (15-DaCompany Report #A0390002A  
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia Difficulty In Walking Dysarthria	Foreign Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG TWICE PER DAY ORAL		Vision Blurred					

Date:12/31/02ISR Number: 4036700-5Report Type:Expedited (15-DaCompany Report #A0389949A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Loss Of Consciousness Tremor	Foreign	Zyban Tablet- Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL	2 WK			Oral Contraceptive Cyclobenzaprine Hcl Omeprazole	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/31/02ISR Number: 4036743-1Report Type:Expedited (15-DaCompany Report #B0288380A  
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent 150 MG / PER Impairment/Damage DAY / ORAL		Asthenia Renal Failure Acute	Foreign Health Professional Company Representative	Zyban Tablet-Zyban (Bupropion Hydrochloride)  Bisoprolol Hydrochlorothiazide Amlodipine Atorvastatin Calcium	PS  C C C C	Zyban	ORAL

Date:12/31/02ISR Number: 4036744-3Report Type:Expedited (15-DaCompany Report #B0288386A  
 Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Headache Hyperhidrosis Nausea  Overdose Palpitations Vomiting	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)  Alprazolam Amitriptyline Clonidine	PS  C C C	Zyban	ORAL

Date:01/01/03ISR Number: 4033979-0Report Type:Direct Company Report #CTU 183686  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG DAILY ORAL		Rash Erythematous Rash Macular		Wellbutrin Sr 150 Mg Glaxosmithkline	PS	Glaxosmithkline	ORAL

Date:01/02/03ISR Number: 4033703-1Report Type:Expedited (15-DaCompany Report #A0390229A  
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Cognitive Disorder Drug Dependence		Zyban	PS	Glaxo Wellcome	ORAL

Date:01/02/03ISR Number: 4033708-0Report Type:Expedited (15-DaCompany Report #B0267268A  
Age:22 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Exophthalmos		Quomem	PS	Glaxo Wellcome	ORAL
150MG		Eye Pain					
Variable dose	15 DAY	Ocular Vascular Disorder Somnolence Swelling Vision Blurred					

Date:01/03/03ISR Number: 4034662-8Report Type:Expedited (15-DaCompany Report #A0390178A  
Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Wellbutrin	PS	Glaxo Wellcome	
UNKNOWN		Intentional Misuse		Lamictal	SS	Glaxo Wellcome	
UNKNOWN				Celexa	SS		
UNKNOWN							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/03/03ISR Number: 4034681-1Report Type:Expedited (15-DaCompany Report #B0288569A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 19 DAY		Coordination Abnormal		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Dizziness Motor Dysfunction Muscular Weakness Pallor		Levothyrox	C	Glaxo Wellcome	

Date:01/06/03ISR Number: 4039508-XReport Type:Expedited (15-DaCompany Report #US010405  
Age:17 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Coma Complex Partial Seizures Epileptic Aura Grand Mal Convulsion Parosmia Road Traffic Accident Status Epilepticus	Health Professional	Gabitril Wellbutrin	PS SS		

Date:01/07/03ISR Number: 4037697-4Report Type:Expedited (15-DaCompany Report #A0386665A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Asthenia Blindness Unilateral		Wellbutrin	PS	Glaxo Wellcome	ORAL
60MG Per day		Drug Interaction		Metadate Cd	C		ORAL
150MG Per day		Hypoaesthesia		Effexor Xr	C		ORAL
20MG Per day		Vision Blurred		Geodon	C		ORAL

Date:01/08/03ISR Number: 4038453-3Report Type:Expedited (15-DaCompany Report #A0390432A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Coma Intentional Misuse	Health Professional	Wellbutrin Alcohol	PS SS	Glaxo Wellcome	

Date:01/09/03ISR Number: 4039389-4Report Type:Expedited (15-DaCompany Report #B0267268A  
Age:22 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability 150MG Variable dose 15 DAY	Exophthalmos Eye Pain Ocular Vascular Disorder Somnolence Swelling Vasodilatation Vision Blurred		Quomem	PS	Glaxo Wellcome	ORAL

Date:01/10/03ISR Number: 4040007-XReport Type:Expedited (15-DaCompany Report #A0391302A  
Age:18 YR Gender:Male I/FU:I

Outcome	PT
Other	Aggression Libido Increased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sexual Abuse

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day			Wellbutrin	PS	Glaxo Wellcome	

Date:01/10/03ISR Number: 4040008-1Report Type:Expedited (15-DaCompany Report #A0391477A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2 YR		Abortion Spontaneous		Wellbutrin	PS	Glaxo Wellcome	ORAL
Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus							

Date:01/10/03ISR Number: 4040012-3Report Type:Expedited (15-DaCompany Report #A0391683A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 225MG Per day		Abortion Spontaneous		Bupropion	PS	Glaxo Wellcome	ORAL
Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus Pregnancy							

Date:01/10/03ISR Number: 4040013-5Report Type:Expedited (15-DaCompany Report #A0391700A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death	Completed Suicide	Wellbutrin	PS	Glaxo Wellcome
	Intentional Misuse	Benadryl	SS	Glaxo Wellcome
	Photosensitivity Reaction			
	Somnolence			
	Sunburn			

Date:01/10/03ISR Number: 4041455-4Report Type:Expedited (15-DaCompany Report #200212-1919 (0)  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Somnolence Sunburn	Consumer	Benadryl - Formulation Unspecified (Diphenhydramine)	PS		ORAL
PER ORAL				Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
PER ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/13/03ISR Number: 4040999-9Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #CTU 184264

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 125 MG PO QHS		Condition Aggravated		6-Mercaptopurine	PS		ORAL
Initial or Prolonged 30 MG PO QWK		Hypotension		Methotrexate	SS		ORAL
200 MG PO BID		Pyrexia		Wellbutrin	SS		ORAL
250 MG PO BID		Sepsis		Depakote	SS		ORAL
0.6 MG QHS		Sinusitis		Clonidine	SS		
INTRAVENOUS	1 GM IV EVERY			Primaxin	SS		
8 HOURS							

Date:01/13/03ISR Number: 4041054-4Report Type:Expedited (15-DaCompany Report #A0391727A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 75MG Twice per day		Anxiety Disorder		Bupropion	PS	Glaxo Wellcome	
5MG Per day		Arthropathy					
		Attention		Paxil	C	Glaxo Wellcome	
		Deficit/Hyperactivity Disorder					
		Autism Spectrum Disorder					
		Breech Presentation					
		Congenital Anomaly					
		Expressive Language Disorder					
		Maternal Drugs Affecting Foetus					

Date:01/13/03ISR Number: 4041055-6Report Type:Expedited (15-DaCompany Report #A0391739A  
 Age: Gender:Female I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 400MG Per day			Communication Disorder	Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged			Convulsion	Topomax	C		
Other			Difficulty In Walking	Prozac	C		
			Loss Of Consciousness	Desyrel	C		

Date:01/13/03ISR Number: 4041060-XReport Type:Expedited (15-DaCompany Report #B0287627A  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See			Hyperhidrosis	Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged dosage text			Insomnia				
			Irritability				
			Nephrolithiasis				
			Renal Colic				

Date:01/13/03ISR Number: 4042347-7Report Type:Periodic Company Report #2002-05-2190  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Depression	Claritin-D 12 Hour (Loratadine 5mg / Pseudoephedrine Extended Release Tablet	PS		ORAL
ORAL			Health Professional Company Representative	Paxil	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Wellbutrin  
(Bupropion) SS

Date:01/13/03ISR Number: 4042370-2Report Type:Periodic  
Age:13 YR Gender:Male I/FU:I

Company Report #2002063780

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Convulsion	Health Professional	Diflucan Tablets (Fluconazole)	PS		ORAL
250 MG  (DAILY), ORAL				Bupropion Hydrochloride	SS		ORAL

Date:01/15/03ISR Number: 4042096-5Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 184528

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 1 TWICE DAILY Intervention to ORAL Prevent Permanent Impairment/Damage		Tinnitus		Wellbutrin Sr 150mg	PS		ORAL

Date:01/15/03ISR Number: 4042128-4Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 184548

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG BID  ORAL		Dysgeusia		Welbutrin Sr 150 Mg	PS		ORAL
				Nasonex	C		

Date:01/15/03ISR Number: 4042195-8Report Type:Expedited (15-DaCompany Report #B0289056A  
 Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	3 DAY	Abdominal Pain Anaemia	Consumer	Zyban Ercefuryl	PS SS	Glaxo Wellcome Glaxo Wellcome	ORAL ORAL
	3 DAY	Haemolytic Uraemic Syndrome		Visceralgine Vogalene	SS SS		ORAL ORAL
	3 DAY	Haptoglobin Decreased		Mushrooms	C		ORAL
	1 DAY	Hyperbilirubinaemia Influenza Like Illness Jaundice Purpura Renal Failure Scan Abdomen Abnormal Thrombocytopenia Thrombotic Microangiopathy Vomiting					

Date:01/15/03ISR Number: 4042196-XReport Type:Expedited (15-DaCompany Report #B0289066A  
 Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Twice	per day	Chest Pain Pain		Zyntabac	PS	Glaxo Wellcome	ORAL
	27 DAY						

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/15/03ISR Number: 4042303-9Report Type:Direct  
Age:59 YR Gender:Female I/FU:I

Company Report #CTU 184544

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG PO BID	Rash		Wellbutrin 150mg	PS		ORAL
		Urticaria		Premarin	C		
				Synthroid	C		
				Aciphex	C		
				Celebrex	C		

Date:01/16/03ISR Number: 4042655-XReport Type:Expedited (15-DaCompany Report #A0388161A  
Age:19 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Lamictal	PS	Glaxo Wellcome	ORAL
		Intentional Misuse		Wellbutrin	SS	Glaxo Wellcome	ORAL
				Celexa	SS		

Date:01/16/03ISR Number: 4042657-3Report Type:Expedited (15-DaCompany Report #A0390178A  
Age:19 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS	Glaxo Wellcome	
UNKNOWN		Intentional Misuse		Lamictal	SS	Glaxo Wellcome	
UNKNOWN				Celexa	SS		

Date:01/16/03ISR Number: 4042670-6Report Type:Expedited (15-DaCompany Report #D0010841A  
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day 9 DAY	Amnesia		Zyban	PS	Glaxo Wellcome	ORAL
Hospitalization -		Cardiac Arrest					

Initial or Prolonged Electroencephalogram  
Other Abnormal  
Grand Mal Convulsion

Date:01/17/03ISR Number: 4043214-5Report Type:Expedited (15-DaCompany Report #A0392519A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blindness		Wellbutrin	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day				Adipex-P	C	Glaxo Wellcome	

Date:01/17/03ISR Number: 4043215-7Report Type:Expedited (15-DaCompany Report #A0392586A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Phaeochromocytoma		Wellbutrin	PS	Glaxo Wellcome	ORAL

Date:01/17/03ISR Number: 4043226-1Report Type:Expedited (15-DaCompany Report #B0289128A  
Age:31 YR Gender:Male I/FU:I

Outcome PT  
Hospitalization - Blood Creatine  
Initial or Prolonged Phosphokinase Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
40TABS Single		Dizziness Dysarthria Intentional Misuse	Consumer	Zyban	PS	Glaxo Wellcome	ORAL
dose	1 DAY	Oxygen Saturation Decreased					
1PACK Single		Sinus Tachycardia		Zopiclone	SS		ORAL
dose	1 DAY	Somnolence					
1PACK Single		Vomiting		Valium	SS		ORAL
dose	1 DAY						

Date:01/20/03ISR Number: 4043555-1Report Type:Expedited (15-DaCompany Report #A0380824A  
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day		Dysphagia		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged 1TAB Per day		Hypersensitivity		Birth Control Pill	C		
Other		Oesophageal Stenosis		Vitamin	C		

Date:01/20/03ISR Number: 4043558-7Report Type:Expedited (15-DaCompany Report #A0392609A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Ileus Paralytic		Wellbutrin	PS	Glaxo Wellcome	
Initial or Prolonged per day	5 MON	Urinary Retention					
40MG Per day				Paxil	C	Glaxo Wellcome	ORAL
.5MG As				Klonopin	C		ORAL

required

Date:01/20/03ISR Number: 4043561-7Report Type:Expedited (15-DaCompany Report #B0260713A  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Anxiety		Zyban	PS	Glaxo Wellcome	ORAL
14 DAY		Excitability					
		Insomnia					
		Lymphocytic Lymphoma					
		Pruritus Generalised					
		Rash Erythematous					
		Vertigo					

Date:01/20/03ISR Number: 4043562-9Report Type:Expedited (15-DaCompany Report #B0271161A  
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Mean Cell Volume		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	24 DAY	Increased		Miniphase	SS		ORAL
1TAB Per day	YR	Pain In Extremity		Lexomil	SS		ORAL
.5TAB Twice		Peripheral Sensory					
per day	YR	Neuropathy		Di-Antalvic	SS		ORAL
2UNIT Monthly	YR	Spinal Osteoarthritis					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/20/03ISR Number: 4043563-0Report Type:Expedited (15-DaCompany Report #B0273874A  
 Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300MG per day 31 DAY	Aspartate		Zyban	PS	Glaxo Wellcome	ORAL
Life-Threatening	26 DAY	Aminotransferase		Lectil	SS		ORAL
Hospitalization -	10 DAY	Increased		Tanganil	SS		ORAL
Initial or Prolonged	10MG Per day 7 DAY	Blood Creatine		Stilnox	SS		ORAL
Other		Phosphokinase Increased Blood Creatinine Increased Brain Oedema Cardiac Enzymes Increased Cardio-Respiratory Arrest Cervical Vertebral Fracture Coma Corneal Reflex Decreased Electrocardiogram T Wave Amplitude Increased Fall Hyperkalaemia Malaise Mydriasis Pallor Sinus Tachycardia Ventricular Fibrillation					

Date:01/20/03ISR Number: 4043568-XReport Type:Expedited (15-DaCompany Report #B0289266A  
 Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	Cardiomegaly		Zyban	PS	Glaxo Wellcome	
UNKNOWN		Coronary Artery Atherosclerosis Coronary Artery Disease Coronary Artery Stenosis Fatigue					



Haemorrhage

Date:01/20/03ISR Number: 4043570-8Report Type:Expedited (15-DaCompany Report #B0289510A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Retinal Oedema		Zyban	PS	Glaxo Wellcome	ORAL
150MG Per day	4 DAY						

Date:01/21/03ISR Number: 4045566-9Report Type:Expedited (15-DaCompany Report #L02-USA-03743-04  
Age:31 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Literature Health Professional	Citalopram (Citalopram Hydrobromide) Bupropion Amphetamine/Dextroph etamine	PS SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/22/03ISR Number: 4044515-7Report Type:Expedited (15-DaCompany Report #B0289403A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypothyroidism		Zyban	PS	Glaxo Wellcome	ORAL
150MG Twice							
per day	21 DAY	Urticaria					
				Dextropropoxyphene + Paracetamol	C		ORAL

Date:01/22/03ISR Number: 4044549-2Report Type:Direct Company Report #CTU 184990  
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Depressed Level Of		Zyprexa	PS		ORAL
10MG BID ORAL							
Hospitalization -		Consciousness		Wellbutrin	SS		ORAL
150MG BID							
Initial or Prolonged		Neuroleptic Malignant					
ORAL		Syndrome					

Date:01/23/03ISR Number: 4044995-7Report Type:Expedited (15-DaCompany Report #A0381384A  
Age:16 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser	Health	Wellbutrin	PS	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)	900MG	Euphoric Mood	Professional				
dose	Single	Grand Mal Convulsion					
		Intentional Misuse		Ethanol	C		
		Medication Error		Cannabis	C		

Date:01/23/03ISR Number: 4047156-0Report Type:Expedited (15-DaCompany Report #A0393335A  
Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Coma Completed Suicide Convulsion Overdose Tachycardia	Literature Health Professional	Wellbutrin (Unspecified Tablet (Bupropion Hydrochloride)	PS		

Date:01/23/03ISR Number: 4047161-4Report Type:Expedited (15-DaCompany Report #A0393333A  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Electrolytes Abnormal Cardiac Arrest Coma Completed Suicide Convulsion Hyperpyrexia Liver Function Test Abnormal Overdose Renal Failure Respiratory Arrest	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/23/03ISR Number: 4047223-1Report Type:Expedited (15-DaCompany Report #L03-USA-00135-08

Age:49 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Carbamazepine	PS		
		Intentional Misuse	Health	Warfarin	SS		
			Professional	Bupropion	SS		

Date:01/24/03ISR Number: 4045780-2Report Type:Expedited (15-DaCompany Report #A0393342A

Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG Twice		Dehydration	Health	Bupropion	PS	Glaxo Wellcome	ORAL
Initial or Prolonged per day	9 DAY	Dysarthria	Professional				
10MG At night		Emotional Distress		Ambien	SS		ORAL
5MG As		Insomnia		Oxycodone	SS		ORAL
required		Oral Intake Reduced					
30MG Three				Oxycontin	SS		ORAL
times per day				Blinded Trial Medication	C	Glaxo Wellcome	
65 DAY				Neurontin	C		
				Tenormin	C		
				Paxil	C	Glaxo Wellcome	
				Pamelor	C		

Date:01/24/03ISR Number: 4045781-4Report Type:Expedited (15-DaCompany Report #A0393594A

Age:10 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anorexia		Wellbutrin	PS	Glaxo Wellcome	ORAL
4 YR							

Growth Retardation  
Malnutrition

Depakote  
Lithium  
Synthroid

C  
C  
C  
Glaxo Wellcome  
Glaxo Wellcome

Date:01/24/03ISR Number: 4045782-6Report Type:Expedited (15-DaCompany Report #B0100993A  
Age:57 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150MG See Hospitalization - dosage text 4 DAY Initial or Prolonged RESPIRATORY Other (INHALATION) 50MCG Twice per day RESPIRATORY (INHALATION) 2PUFF Four times per day RESPIRATORY (INHALATION) 2PUFF Twice per day	Asthma Chronic Obstructive Airways Disease Exacerbated	Health Professional	Zyban  Salmeterol  Salbutamol  Becotide	PS  C  C  C	Glaxo Wellcome  Glaxo Wellcome  Glaxo Wellcome  Glaxo Wellcome	ORAL

Date:01/24/03ISR Number: 4045783-8Report Type:Expedited (15-DaCompany Report #B0107035A  
Age:18 YR Gender:Male I/FU:F

Outcome PT  
Death Aggression  
Alcohol Poisoning  
Anorexia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration		Report Source	Product	Role	Manufacturer	Route
3	WK	Anxiety Completed Suicide Depression	Health	Zyban	PS	Glaxo Wellcome	ORAL
		Disturbance In Attention	Professional	Chloroquine	C		
		Drug Abuser Feeling Abnormal Head Injury Headache Hypochondriasis Indifference Injury Asphyxiation Irritability Loss Of Consciousness Mania Mood Swings Palpitations Personality Change Vision Blurred					

Date:01/24/03ISR Number: 4048253-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #A001-002-005407

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1 IN 1 D, PER Initial or Prolonged ORAL		Confusional State	Consumer	Aricept (Donepezil)	PS		ORAL
		Fall					
25 MG, 1 IN 1 D, PER ORAL		Injury Salivary Hypersecretion		Zoloft (Sertraline Hydrochloride)	SS		ORAL
		Weight Decreased					
ORAL				Wellbutrin (Bupropion)	SS		ORAL

Date:01/27/03ISR Number: 4046864-5Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-12093845

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
50 mg at night		Drug Interaction		Trazodone Hcl Tabs	PS	Apothecon	ORAL
		Feeling Abnormal					
		Sedation		Wellbutrin Sr	I		ORAL

Date:01/27/03ISR Number: 4047032-3Report Type:Expedited (15-DaCompany Report #A0393180A  
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Dyspnoea		Zyban	PS	Glaxo Wellcome	ORAL

Date:01/27/03ISR Number: 4047033-5Report Type:Expedited (15-DaCompany Report #A0393978A  
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 200MG per day		Abortion Spontaneous		Bupropion	PS	Glaxo Wellcome	
		Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus Pregnancy					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/27/03ISR Number: 4047034-7Report Type:Expedited (15-DaCompany Report #B0087348A

Age:53 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice Initial or Prolonged per day 8 DAY	Confusional State		Bupropion	PS	Glaxo Wellcome	ORAL
TRANSDERMAL	Depressed Level Of Consciousness Grand Mal Convulsion 1 DAY		Premique Nicotine	SS C	Glaxo Wellcome	
2UNIT Four times per day	Malaise Parosmia Periorbital Oedema Vision Blurred		Omeprazole Paracetamol	C C	Glaxo Wellcome	

Date:01/27/03ISR Number: 4047036-0Report Type:Expedited (15-DaCompany Report #B0285584A

Age:47 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Per day 29 DAY	Arrhythmia		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged UNKNOWN	Blood Alcohol Increased		Humalog Insulin	C		
TRANSDERMAL	Blood Cholesterol Increased 46 DAY		Stagid Nicotine Patch	C C	Glaxo Wellcome	ORAL
1 DAY	Chest Pain		Alcohol	C		ORAL
	Confusional State Depressed Level Of Consciousness Fall Hypertension Hypertriglyceridaemia Lipids Abnormal Malaise Muscular Weakness Paresis Sensory Loss Transient Ischaemic					



Date:01/27/03ISR Number: 4047042-6Report Type:Expedited (15-DaCompany Report #B0290194A  
 Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Exanthem		Zyban	PS	Glaxo Wellcome	
UNKNOWN	150MG	Twice					
per day	16	DAY					
				Fluticasone Propionate	C	Glaxo Wellcome	
RESPIRATORY							
(INHALATION)	.5MG	Twice					
per day							

Date:01/27/03ISR Number: 4047946-4Report Type:Expedited (15-DaCompany Report #325095  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Literature	Accutane			
Life-Threatening		Coagulopathy	Health	(Isotretinoin)	PS		
		Completed Suicide	Professional	Bupropion (Bupropion			
		Haemorrhage		Hydrochloride)	SS		
		Multi-Organ Failure		Diphenhydramine			
				(Diphenhydramine			
				Hydrochloride)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/27/03ISR Number: 4048687-XReport Type:Expedited (15-DaCompany Report #S02-USA-03223-01

Age:19 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose	Health Professional Company Representative	Celexa (Citalopram Hydrobromide)	PS		
	60 MG QD PO			Celexa (Citalopram Hydrobromide)	SS		ORAL
				Lamictal (Lamotrigine)	SS		
				Wellbutrin (Bupropion Hydrochloride)	SS		
	300 MG QD						

Date:01/28/03ISR Number: 4047979-8Report Type:Expedited (15-DaCompany Report #A0388161A

Age:19 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse		Lamictal	PS	Glaxo Wellcome	ORAL
				Wellbutrin	SS	Glaxo Wellcome	ORAL
				Celexa	SS		

Date:01/28/03ISR Number: 4047982-8Report Type:Expedited (15-DaCompany Report #A0392609A

Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Ileus Paralytic		Wellbutrin	PS	Glaxo Wellcome	
150MG Twice Initial or Prolonged per day	5 MON	Urinary Retention		Paxil	C	Glaxo Wellcome	ORAL
40MG Per day				Klonopin	C		ORAL
.5MG As							

required

Date:01/28/03ISR Number: 4047983-XReport Type:Expedited (15-DaCompany Report #A0393413A  
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day		Convulsion		Zyban	PS	Glaxo Wellcome	ORAL
Initial or Prolonged		Drug Interaction		Unspecified Medications	SS		

Date:01/28/03ISR Number: 4048549-8Report Type:Direct Company Report #CTU 185460  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG DAILY		Rash Erythematous Rash Macular		Wellbutrin Sr 150mg Glaxosmithkline	PS	Glaxosmithkline	ORAL
ORAL							

Date:01/28/03ISR Number: 4048845-4Report Type:Expedited (15-DaCompany Report #A03200300069  
Age:57 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Blood Potassium Abnormal Cerebral Ischaemia Dehydration Dysarthria Electrocardiogram Qrs

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Complex Prolonged Electrocardiogram T Wave Abnormal Electroencephalogram	Report Source	Product	Role	Manufacturer	Route
10 MG HS -		Abnormal Emotional Distress Insomnia	Health Professional	Ambien - Zolpidem Tartrate - Tablet - 10 Mg	PS		ORAL
ORAL		Oligodipsia					
NI UNK -		Oral Intake Reduced		Investigational Drug - Not Provided - Unknown - Unit Dose: Unknown	SS		
UNKNOWN							
30 MG TID -				Oxycontin - Oxycodone Hydrochloride - Unknown - Unit Dose: Unknown	SS		ORAL
ORAL							
5 MG PRN -				Oxycodone - Unknown - Unit Dose: Unknown	SS		ORAL
ORAL							
100 MG BID -				Wellbutrin - Bupropion Hydrochloride - Unknown - Unit Dose: Unknown	SS		ORAL
ORAL							
				Gabapentin	C		
				Atenolol	C		
				Paroxetine			
				Hydrochloride	C		
				Nortriptyline			
				Hydrochloride	C		

Date:01/28/03ISR Number: 4049367-7Report Type:Expedited (15-DaCompany Report #A0392905A  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest Completed Suicide Overdose Respiratory Arrest	Literature Health Professional	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		

Date:01/29/03ISR Number: 4048725-4Report Type:Expedited (15-DaCompany Report #B0290356A  
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day	20 DAY	Abdominal Pain Upper Dyspepsia Joint Swelling Urticaria		Zyban	PS	Glaxo Wellcome	ORAL

Date:01/31/03ISR Number: 4049592-5Report Type:Expedited (15-DaCompany Report #A0393425A  
Age:17 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Complex Partial Seizures Epileptic Aura

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Road Traffic Accident Status Epilepticus Tonic Clonic Movements	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Health Professional	Wellbutrin Gabitril	PS SS	Glaxo Wellcome	ORAL

Date:01/31/03ISR Number: 4049980-7Report Type:Expedited (15-DaCompany Report #WAES 0211USA01406  
Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Circulatory Collapse Drug Interaction Drug Level Increased Heat Stroke		Cogentin Wellbutrin Neurontin	PS SS SS	Merck & Co., Inc	

Date:01/31/03ISR Number: 4050970-9Report Type:Expedited (15-DaCompany Report #USA-2002-0002070  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Anoxic Encephalopathy Apallic Syndrome Circulatory Collapse Depression	Consumer Health Professional Other	Oxycontin Tablets(Oxycodone Hydrochloride) Cr Tablet			
Hospitalization - Initial or Prolonged Other					PS		
INTRAVENOUS	INTRAVENOUS	Drug Toxicity Loss Of Consciousness Multi-Organ Failure Overdose Polysubstance Abuse Respiratory Arrest Skin Disorder		Trazodone (Trazodone) Propoxyphene (Dextrop ropoxyphene) Unknown Acetaminophen (Parace tamol) Unknown Bupropion (Amfebutamo ne)	SS SS SS SS		

Date:02/03/03ISR Number: 4050409-3Report Type:Direct Company Report #CTU 185772  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Chest Pain		Zyban 150 Mg Began			

Erythema Multiforme 1/9 PS  
 ZYBAN 150 MG  
 BID  
 Nicotine Patch Began 1/9/03 SS  
 NICOTINE  
 PATCH

Date:02/03/03ISR Number: 4050686-9Report Type:Direct Company Report #CTU 183825  
 Age:17 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Contusion		Bupropion Sr 150mg	PS		ORAL
150MG BID		Purpura					
ORAL							

Date:02/03/03ISR Number: 4051499-4Report Type:Expedited (15-DaCompany Report #2003002684  
 Age:84 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Literature Health Professional	Doxazosin (Doxazosin) Digoxin (Digoxin)	PS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Bupropion  
(Bupropion) SS

Date:02/04/03ISR Number: 4052245-0Report Type:Expedited (15-DaCompany Report #B0290660A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Foreign Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL							

Date:02/04/03ISR Number: 4052642-3Report Type:Expedited (15-DaCompany Report #HQWYE070029JAN03  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Electroencephalogram Abnormal Grand Mal Convulsion Headache Malaise Parosmia	Health Professional Other	Premique (Conjugated Estrogens/Medroxyprogesterone Acetate, Tablet, 5 Mg)	PS		ORAL
150MG DAILY	8 DAY	Periorbital Oedema Vision Blurred		Zyban (Amfebutaome Hydrochloride)	SS		
				Nicotine (Nicotine)	C		
				Omeprazole (Omeprazole)	C		
				Paracetamol (Paracetamol)	C		

Date:02/04/03ISR Number: 4052662-9Report Type:Expedited (15-DaCompany Report #B0290662A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Suicide Attempt	Foreign Consumer	Bupropion Hydrochloride Tablet-Zyban (Bupropion			



ORAL

Hydrochloride)

PS

ORAL

Date:02/05/03ISR Number: 4051209-0Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0263321A

Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	ORAL
19 DAY		Agitation		Zithromax	C		ORAL
250MG Twice		Bipolar Disorder					
per day	3	DAY					
200MG Twice		Crying		Surgam	C		ORAL
per day	3	DAY					
3 DAY		Insomnia					
		Micturition Urgency		Doliprane	C	Glaxosmithkline	ORAL
		Obsessive-Compulsive Disorder					
		Suicidal Ideation					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/05/03ISR Number: 4051210-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0282944A  
 Age:37 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Per day 32 DAY Initial or Prolonged	Aggression Anxiety Delirium Delusional Disorder, Persecutory Type Depression Suicide Attempt		Zyban	PS	Glaxosmithkline	ORAL

Date:02/05/03ISR Number: 4051211-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0283322A  
 Age:43 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 10 DAY Initial or Prolonged	Dyspnoea Nephritis		Zyban	PS	Glaxosmithkline	ORAL

Date:02/05/03ISR Number: 4051217-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0290894A  
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 81 DAY Initial or Prolonged	Brain Neoplasm Convulsion		Zyban	PS	Glaxosmithkline	ORAL

Date:02/05/03ISR Number: 4051769-XReport Type:Direct Company Report #CTU 186083  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose 150 MG PO 1 BID	Anxiety Insomnia		Wellbutrin Sr	PS		ORAL

Date:02/06/03ISR Number: 4051792-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0395439A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly			Congenital Anomaly	Wellbutrin	PS	Glaxosmithkline	
			Maternal Drugs Affecting Foetus	Unknown Medication	C		

Date:02/06/03ISR Number: 4051793-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0395460A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Renal Disorder	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/06/03ISR Number: 4051794-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0395488A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG Twice Initial or Prolonged per day			Dehydration	Wellbutrin	PS	Glaxosmithkline	ORAL
			Drug Interaction	Phenergan With Codeine	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/07/03ISR Number: 4053840-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #2002107546US

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Dyspnoea Urticaria	Health Professional	Celebrex(Celecoxib)C apsule	PS		
				Neurontin(Gabapentin )	SS		
				Topamax(Topiramate)	SS		
				Zantac(Ranitidine Hydrochloride)	SS		
				Wellbutrin(Amfebutam one Hydrochloride)	SS		
				Paxil(Paroxetine Hydrochloride)	SS		
				Klonopin(Clonazepam)	SS		
				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		
				Progesterone	C		
				Salbutamol(Salbutamol)	C		
				Pentosan Polysulfate(Pentosan Polysulfate)	C		
				Hydroxyzine(Hydroxyzine)	C		
				Amitriptyline	C		

Date:02/07/03ISR Number: 4055166-2Report Type:Expedited (15-DaCompany Report #B0290194A  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage TWICE PER DAY		Aggression Dizziness Exanthem Hyperhidrosis Nausea Skin Exfoliation	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
/ ORAL				Fluticasone			

Sleep Disorder  
Vomiting

Propionate  
Nicotine

C  
C

Date:02/10/03ISR Number: 4053176-2Report Type:Expedited (15-DaCompany Report #NZ-GLAXOSMITHKLINE-B0289919A  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Intolerance		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	41 DAY	Hypertension					

Date:02/10/03ISR Number: 4055356-9Report Type:Expedited (15-DaCompany Report #B0290662A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Suicide Attempt	Foreign Consumer	Bupropion Hydrochloride Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/10/03ISR Number: 4055467-8Report Type:Expedited (15-DaCompany Report #A0394895A  
Age:16 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Neuroleptic Malignant Syndrome	Health Professional Company Representative	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride) Aripiprazole (Formulation Unknown) (Aripiprazole)	PS  SS		

Date:02/11/03ISR Number: 4053975-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0388881A  
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Asthenia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 5 DAY							
Initial or Prolonged		Choreoathetosis		Prograf	C		
Other		Coordination Abnormal		Neurontin	C		
				Synthroid	C	Glaxosmithkline	
				Fosamax	C		
				Vasotec	C		
				Florinef	C		
				Hydrocortisone	C	Glaxosmithkline	

Date:02/11/03ISR Number: 4053983-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0288380A  
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day 6 DAY		Asthenia		Zyban	PS	Glaxosmithkline	ORAL
		Haematuria		Bisoprolol	C		
		Laboratory Test Interference		Hydrochlorothiazide	C		
		Leukocyturia		Amlodipine	C		
		Malaise		Atorvastatin	C		
		Proteinuria					

Date:02/11/03ISR Number: 4053985-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0290826A  
Age:49 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300MG per day Initial or Prolonged	Dehydration Dermatitis Exfoliative Ectropion Malaise Mean Cell Volume Increased Psoriasis Rash Papular Skin Hypertrophy Staphylococcal Infection		Zyban	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/11/03ISR Number: 4053991-5Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0040415A  
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged dosage text 1 DAY		Coma		Zyban	PS	Glaxosmithkline	ORAL
500MG See dosage text 1 DAY		Intentional Misuse Suicide Attempt		Paracetamol	SS	Glaxosmithkline	ORAL

Date:02/11/03ISR Number: 4056013-5Report Type:Direct Company Report #CTU 186548  
 Age:22 WK Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other TRANSMAMMARY	DAILY BF	Convulsion Drug Exposure Via Breast Milk		Luvox (Taken By Mother) (Fluvoxamine Maleate)	PS		
TRANSMAMMARY	300MG DAILY BF			Wellbutrin (Taken By Mother)(Amfebutamone Hydrochloride)	SS		

Date:02/11/03ISR Number: 4059250-9Report Type:Direct Company Report #CTU 187041  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Urticaria		Bupropion 150mg Sr	PS		

Date:02/12/03ISR Number: 4054733-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0392917A  
 Age:50 YR Gender:Female I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Supraventricular Tachycardia		Wellbutrin Trazodone	PS SS	Glaxosmithkline	ORAL

50MG Per day

Date:02/12/03ISR Number: 4054736-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0395212A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Grand Mal Convulsion Impaired Work Ability		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/12/03ISR Number: 4057640-1Report Type:Periodic Company Report #A0385046A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Chromatopsia Hallucination	Consumer Other	Paxil (Formulation Unknown) (Paroxetine Hydrochloride)	PS		ORAL
ORAL				Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	SS		
				Esomeprazole	C		
				Diphenhydramine Hcl	C		
				Citalopram			

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Hydrobromide	C
Metaxalone	C
Biaxin	C
Ziac	C
Tenoretic	C
Excedrin Migraine	C
Arthrotec	C
Perphenazine	C
Famotidine	C
Benztropine Mesylate	C
Decongestant	C
Ziprasidone Hci	C
Oxcarbazepine	C
Paracetamol	C
Hydrocodone	C
Alka-Seltzer	C
Cold Medication	C

Date:02/13/03ISR Number: 4055486-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0396170A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 9900MG Single Initial or Prolonged dose			Intentional Misuse	Wellbutrin	PS	Glaxosmithkline	
			Suicide Attempt				

Date:02/13/03ISR Number: 4055489-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0396270A  
 Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Overdose	Professional	Amitriptyline	SS		

Date:02/14/03ISR Number: 4056052-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0380141A  
 Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other 150MG Twice per day	14	DAY	Chills Face Oedema Mouth Ulceration Pharyngeal Oedema	Health Professional	Zyban  Equate Nicotine Patch	PS  SS	Glaxosmithkline  Glaxosmithkline	ORAL
INTRADERMAL			Pyrexia Sinus Congestion Tongue Oedema Tongue Ulceration Urticaria Viral Infection		Clarinx	C		

Date:02/14/03ISR Number: 4056053-6Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0390002A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day		Asthenia  Difficulty In Walking  Dysarthria Vision Blurred		Zyban	PS	Glaxosmithkline	ORAL

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Date:02/14/03ISR Number: 4056061-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0396296B

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day Initial or Prolonged			Coarctation Of The Aorta	Bupropion	PS	Glaxosmithkline	
			Maternal Drugs Affecting Foetus				

Date:02/14/03ISR Number: 4056063-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0396676A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other			Convulsion Drug Interaction	Wellbutrin Provigil Ambien Seroquel	PS SS SS SS	Glaxosmithkline	ORAL

Date:02/14/03ISR Number: 4056072-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0291519A

Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Unknown 5 DAY			Wheezing	Zyban	PS	Glaxosmithkline	ORAL
UNKNOWN	2MG per day			Oestradiol	C		
UNKNOWN	50UG Twice			Salmeterol	C	Glaxosmithkline	
per day				Salbutamol	C	Glaxosmithkline	
UNKNOWN	100UG As						
required				Beclomethasone	C	Glaxosmithkline	
RESPIRATORY (INHALATION)	100UG Twice						
per day							

Date:02/14/03ISR Number: 4056075-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0291822A  
 Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Per day	5 DAY	Alcohol Withdrawal	Consumer	Zyban	PS	Glaxosmithkline	ORAL
200MG Per day	5 DAY	Syndrome		Disulfiram	SS		ORAL
40MG Per day		Coccydynia		Atorvastatin	C		ORAL
				Alcohol	C		
UNKNOWN							

Date:02/14/03ISR Number: 4056077-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0291848A  
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Multiple Sclerosis		Zyban	PS	Glaxosmithkline	ORAL
150MG							
Initial or Prolonged							
Variable dose	36 DAY						
3UNIT per day				Thyroxine	C	Glaxosmithkline	
1UNIT per day				Simvastatin	C		
				Nortriptyline	C		
				Lactulose	C		
				Frusemide	C	Glaxosmithkline	
1UNIT per day				Aspirin	C		
1UNIT per day				Diltiazem	C	Glaxosmithkline	
3UNIT per day				Cephalexin	C	Glaxosmithkline	
2UNIT per day				Tolterodine	C		

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Date:02/14/03ISR Number: 4056079-2Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0291940A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebral Haemorrhage		Bupropion Hydrochloride	PS	Glaxosmithkline	
UNKNOWN	150MG	Unknown 2 MON					

Date:02/14/03ISR Number: 4056674-0Report Type:Direct Company Report #CTU 186681

Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Loss Of Libido		Zoloft 200mg	PS		ORAL
Other		Sexual Dysfunction					
200MG/DAY							
ORAL				Wellbutrin Sr 100mg	SS		ORAL
100MG/DAY							
ORAL							

Date:02/18/03ISR Number: 4056391-7Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0290894A

Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Asthenia		Zyban	PS	Glaxosmithkline	ORAL
81 DAY							
Initial or Prolonged		Blood Glucose Abnormal		Aspirin	C		
UNKNOWN	1TAB per day						
UNKNOWN		Brain Neoplasm		Antibiotics	C		
		Convulsion		Alcohol	C		
		Nausea					
		Somnolence					
		Ventricular Tachycardia					
		Vomiting					

Date:02/18/03ISR Number: 4056396-6Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0291987A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pleuritic Pain		Zyban	PS	Glaxosmithkline	ORAL
300MG per day	6 DAY						

Date:02/18/03ISR Number: 4056397-8Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-B0292143A  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Abasia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
Initial or Prolonged		Generalised Oedema					
per day	22 DAY						
Disability		Immobile					
Other		Pain					
		Pyrexia					
		Rash					
		Weight Decreased					

Date:02/19/03ISR Number: 4056867-2Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0039257A  
Age:55 YR Gender:Male I/FU:F

Outcome	PT
Life-Threatening	Blood Glucose Abnormal
Hospitalization -	Blood Pressure Systolic
Initial or Prolonged	Increased
Other	Circulatory Collapse
	Communication Disorder
	Convulsion

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
4TAB per day	2 DAY	Disturbance In Attention General Physical Health Deterioration		Zyban	PS	Glaxosmithkline	ORAL
		Lactic Acidosis		Metformin	SS		ORAL
		Overdose Sinus Tachycardia White Blood Cell Count Increased					

Date:02/19/03ISR Number: 4057591-2Report Type:Direct Company Report #CTU 186846  
 Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Anxiety Discomfort Dizziness Extrasystoles Heart Rate Increased Insomnia Ventricular Extrasystoles		Wellbutrin 150 Mg	PS		

Date:02/19/03ISR Number: 4059931-7Report Type:Periodic Company Report #A03200201075  
 Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion Hypotension Loss Of Consciousness	Health Professional	Ambien Wellbutrin - Slow Release	PS SS		ORAL
400 MG DAILY				Ethanol	SS		
ORAL	2 YR			Venlafaxine Hydrochloride Adderall	C C		



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 50 MG QD ORAL		Drug Interaction	Consumer	Provigil	PS		ORAL
Initial or Prolonged 40 MG QD ORAL		Headache		Celexa	SS		ORAL
				Wellbutrin	SS		ORAL
100 MG QD							
ORAL				Synthroid	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 500MG PO TID		Blood Glucose Decreased		Amoxicillin 500mg	PS		ORAL
150MG PO BID		Dyspnoea		Zyban 150mg	SS		ORAL
		Rash		Metformin	C		
				Actos	C		
				Xenical	C		
				Zocor	C		
				Lisinopril	C		
				Hctz	C		
				Oxybutin	C		

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Mvi C  
 Percocet C  
 Darvocet C  
 Fish Oil C  
 Calcium C

Date:02/20/03ISR Number: 4058166-1Report Type:Direct  
 Age:39 YR Gender:Male I/FU:I

Company Report #CTU 187006

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tremor		Wellbutrin (Bupropion) 150 Mg Sr Tablet	PS		ORAL
300 MG/DAY							
ORAL							

Valproic Acid C  
 Hydroxyzine C  
 Trazodone C  
 Paxil C

Date:02/20/03ISR Number: 4058315-5Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #CTU 186926

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion		Sertraline 100mg	PS		ORAL
200MG DAILY							
Initial or Prolonged							
ORAL							
75MG BID ORAL							

Bupropion 75mg SS ORAL  
 Carbamazepine C  
 Fentanyl C  
 Furosemide C  
 Insulin C  
 Metolazone C  
 Propoxyphene/Apap C  
 Simvastatin C  
 Trazosone C  
 Valsartan C  
 Warfarin C

Date:02/21/03ISR Number: 4058516-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0397012A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300MG per day							
			Blighted Ovum	Bupropion	PS	Glaxosmithkline	ORAL
			Maternal Drugs Affecting Foetus Pregnancy Vaginal Haemorrhage				

Date:02/21/03ISR Number: 4058518-XReport Type:Expedited (15-DaCompany Report #NZ-GLAXOSMITHKLINE-B0284138A  
Age:28 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Balance Disorder Derealisation Difficulty In Walking Discomfort Dyspnoea Emotional Distress

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MGD per day	16 DAY	Erythema Multiforme Face Oedema Fatigue Feeling Abnormal Memory Impairment Oedema Peripheral Panic Reaction Rash Papular Rash Pruritic Tenderness Tinnitus		Zyban	PS	Glaxosmithkline	ORAL

Date:02/21/03ISR Number: 4058519-1Report Type:Expedited (15-DaCompany Report #NZ-GLAXOSMITHKLINE-B0284417A  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG per day Initial or Prolonged	9 DAY	Anaphylactic Reaction Neutrophilia		Zyban	PS	Glaxosmithkline	ORAL

Date:02/21/03ISR Number: 4058525-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0292446A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day TRANSDERMAL		Chills Diplopia Dizziness Headache Musculoskeletal Stiffness Tremor		Zyban Oestradiol	PS C	Glaxosmithkline	ORAL

Date:02/21/03ISR Number: 4059246-7Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 187072

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Urticaria Papular		Bupropion 150mg Sr	PS		

Date:02/21/03ISR Number: 4059247-9Report Type:Direct Company Report #CTU 187074  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Palpitations		Bupropion 150mg Sr	PS		

Date:02/24/03ISR Number: 4059456-9Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0396990A  
Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Fall		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Motor Dysfunction Paralysis Psychomotor Skills Impaired					

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Date:02/25/03ISR Number: 4059899-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0397019A  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 24G Single Initial or Prolonged dose		Brain Damage	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
		Convulsion	Professional				
		Electrocardiogram Qt Prolonged Hypoxia Overdose		Alcohol	SS		ORAL

Date:02/25/03ISR Number: 4059901-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0397394A  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Per day		Abortion Spontaneous		Bupropion	PS	Glaxosmithkline	ORAL
		Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus					

Date:02/25/03ISR Number: 4059902-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0397823A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Brain Damage Convulsion		Wellbutrin	PS	Glaxosmithkline	

Date:02/25/03ISR Number: 4059905-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0292325A  
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Anxiety  
 150MG See Zyban PS Glaxosmithkline  
 Initial or Prolonged Depression  
 dosage text  
 Neuralgia  
 Paraesthesia  
 Pyrexia  
 Sense Of Oppression

Date:02/25/03ISR Number: 4059909-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0292690A

Age:33 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Creatinine	Consumer	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Increased					
per day	9 DAY						
UNKNOWN		Drug Interaction		Tacrolimus	SS		
UNKNOWN		Drug Level Decreased		Insulin	C		

Date:02/25/03ISR Number: 4059916-0Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0038746A

Age:55 YR Gender:Male I/FU:F

Outcome	PT
Life-Threatening	Blood Pressure Systolic
Hospitalization -	Increased
Initial or Prolonged	Circulatory Collapse
Other	Communication Disorder

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2	DAY	Convulsion Diabetes Mellitus Disturbance In Attention	Health	Zyban	PS	Glaxosmithkline	ORAL
850MG	Per day	General Physical Health Deterioration	Professional	Metformin	SS		ORAL
		Lactic Acidosis Po2 Decreased Prescribed Overdose Refusal Of Treatment By Patient Sinus Tachycardia White Blood Cell Count Increased					

Date:02/26/03ISR Number: 4060635-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0292855A  
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG		Agitation		Zyban	PS	Glaxosmithkline	ORAL
Variable dose	12	DAY	Hyperventilation					
INTRAMUSCULAR	150MG	See	Syncope		Medroxyprogesterone	C		
dosage text			Tremor					

Date:02/26/03ISR Number: 4060636-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0292859A  
Age:24 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG per day	12 DAY	Asthma		Zyban	PS	Glaxosmithkline	ORAL
RESPIRATORY					Salmeterol + Fluticasone	C	Glaxosmithkline	
(INHALATION)	50UG	Unknown						



Salbutamol

C

Glaxosmithkline

RESPIRATORY

(INHALATION) 100UG Unknown

Date:02/26/03ISR Number: 4060668-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0397636A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
UNKNOWN		12 YR		Wellbutrin	PS	Glaxosmithkline	
		Abdominal Distension					
		Difficulty In Walking					
		Disorientation					
		Dizziness					
		Drug Withdrawal Syndrome					
		Dysstasia					
		Migraine					
		Musculoskeletal Stiffness					
		Tinnitus					

Date:02/26/03ISR Number: 4060674-4Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0397797B

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
150MG Twice							
Initial or Prolonged							
per day		Neonatal		Bupropion	PS	Glaxosmithkline	
		Maternal Drugs Affecting					
		Foetus					
		Tachypnoea					

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Date:02/27/03ISR Number: 4061861-1Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0397804A  
 Age:26 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Deafness		Zyban	PS	Glaxosmithkline	ORAL
Other		Drug Interaction					
150MG Twice		Ear Pain		Anesthetic	SS		
per day		Hearing Impaired		Tylenol #3	SS		
		Hypoacusis					
		Tinnitus					

Date:02/27/03ISR Number: 4061865-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0107035A  
 Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression		Zyban	PS	Glaxosmithkline	ORAL
Death		Agitation		Chloroquine	C		
3 WK		Alcoholism					
		Anorexia					
		Anxiety					
		Completed Suicide					
		Depression					
		Disturbance In Attention					
		Drug Abuser					
		Feeling Abnormal					
		Head Injury					
		Headache					
		Hypochondriasis					
		Irritability					
		Loss Of Consciousness					
		Mania					
		Mood Swings					
		Obsessive-Compulsive					
		Personality Disorder					
		Palpitations					
		Personality Change					
		Restlessness					
		Vision Blurred					

Date:02/27/03ISR Number: 4068402-3Report Type:Expedited (15-DaCompany Report #200310607BCC  
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Consumer	Aleve (Naproxen			
440 MG, ONCE,		Difficulty In Walking	Other	Sodium)	PS		ORAL
ORAL		Pruritus					
				Wellbutrin			
				(Amfebutamone			
				Hydrochloride)	SS		

Date:02/27/03ISR Number: 4068453-9Report Type:Expedited (15-DaCompany Report #HQWYE535320FEB03  
Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Confabulation
Initial or Prolonged	Confusional State
	Convulsion
	Delirium
	Disorientation

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Dose	Duration	Drug Interaction Hallucination, Visual Memory Impairment	Report Source	Product	Role	Manufacturer	Route
37.5 MG OR 75 MG DAILY, ORAL; 375 MG 1X PER 1 DAY, ORAL			Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		ORAL
200 MG 2X PER 1 DAY, ORAL				Wellbutrin - Slow Release (Amfebutamone Hydrochloride, )	SS		ORAL
				Zyprexa (Olanzapine)	C		

Date:02/28/03ISR Number: 4062501-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0393969B  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day Initial or Prolonged		Caesarean Section		Bupropion	PS	Glaxosmithkline	
		Maternal Drugs Affecting Foetus		Terbutaline	C		
		Neonatal Respiratory		Zofran	C	Glaxosmithkline	
		Distress Syndrome		Ambien	C		
		Premature Baby		Tylenol	C	Glaxosmithkline	
				Tylenol Pm	C		

Date:02/28/03ISR Number: 4062503-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0395488A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	11 DAY	Cardiac Arrest	Other	Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -	11 DAY	Drug Toxicity		Prozac	SS		
Initial or Prolonged	11 DAY	Lobar Pneumonia		Ultram	SS		
per day				Phenergan With Codeine	SS		
UNKNOWN							

Date:02/28/03ISR Number: 4062511-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0397981A  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	11 DAY	Cardiac Arrest	Other	Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -	11 DAY	Drug Toxicity		Prozac	SS		
Initial or Prolonged	11 DAY	Lobar Pneumonia		Ultram	SS		
per day				Phenergan	SS	Glaxosmithkline	

Date:02/28/03ISR Number: 4062515-8Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0398318A  
Age:40 YR Gender:I/FU:I

Outcome	PT
Other	Convulsion Depressed Level Of Consciousness Electrocardiogram Qrs Complex Prolonged

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
7000MG Single dose		Hypoxia Opisthotonus Overdose  Tachycardia	Consumer	Bupropion  Plaquenil	PS  SS	Glaxosmithkline	ORAL

Date:02/28/03ISR Number: 4062516-XReport Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0398320A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Electrocardiogram Qt Prolonged Grand Mal Convulsion Intentional Misuse	Consumer	Bupropion Citalopram Methotrimeprazine	PS SS SS	Glaxosmithkline	

Date:02/28/03ISR Number: 4062517-1Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0398322A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Potassium Decreased Convulsion Electrocardiogram Qrs Complex Prolonged Hallucination Pyrexia Serotonin Syndrome	Consumer	Bupropion Celexa Clonazepam	PS SS SS	Glaxosmithkline	

Date:02/28/03ISR Number: 4062518-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0282937A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged RESPIRATORY		Blood Pressure Increased Fall		Zyban Flixotide	PS C	Glaxosmithkline Glaxosmithkline	ORAL

(INHALATION) 2PUFF Twice  
 Flushing  
 Malaise  
 per day  
 Paraesthesia Serevent C Glaxosmithkline  
 RESPIRATORY  
 (INHALATION) 2PUFF Twice  
 Syncope  
 Tetany  
 per day  
 Trismus  
 Visual Disturbance

Date:02/28/03ISR Number: 4062522-5Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0292806A  
 Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Zyban	PS	Glaxosmithkline	ORAL
31 DAY		Complications Of Maternal Exposure To Therapeutic Drugs Drug Exposure During Pregnancy					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/03ISR Number: 4062523-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0292808A  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day	32 DAY	Abortion Spontaneous	Zyban	PS	Glaxosmithkline	ORAL
			Complications Of Maternal Exposure To Therapeutic Drugs				
			Maternal Drugs Affecting Foetus				

Date:02/28/03ISR Number: 4067181-3Report Type:Expedited (15-DaCompany Report #02P-163-0204399-00  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage			Convulsion	Semisodium Valproate (Depakote) (Divalproex Sodium) (Divalproex Sodium)	PS		ORAL
ORAL				Bupropion Hydrochloride	SS		ORAL
ORAL							

Date:03/03/03ISR Number: 4064786-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0398359A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Abortion Spontaneous	Wellbutrin	PS	Glaxosmithkline	ORAL
			Complications Of Maternal Exposure To Therapeutic Drugs	Seroquel	C		
			Drug Exposure During Pregnancy				

Date:03/03/03ISR Number: 4069171-3Report Type:Expedited (15-DaCompany Report #2003002619  
 Age:36 YR Gender:Unknown I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature	Sinequan (Cap)			
Other		Completed Suicide	Health	(Doxepin)	PS		
		Intentional Misuse	Professional	Bupropion	SS		

Date:03/04/03ISR Number: 4066053-8Report Type:Direct Company Report #CTU 187869  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Growth Retardation		Wellbutrin 100 Mg Sr	PS		ORAL
100 MG SR QD		Weight Gain Poor					
ORAL				Wellbutrin	SS		ORAL
100 MG SR BID							
ORAL				Depakote	C		
				Synthroid	C		
				Remeron	C		
				Lithium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/04/03ISR Number: 4066218-5Report Type:Direct  
Age: Gender: I/FU:I

Company Report #USP 55553

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin Sr	PS	Glaxosmithkline	
				Wellbutrin Sr	SS	Glaxosmithkline	

Date:03/04/03ISR Number: 4070209-8Report Type:Expedited (15-DaCompany Report #2003EU005404  
Age:33 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Creatinine Increased	Foreign Health Professional	Prograf (Tacrolimus) Formulation Unknown	PS		
		Drug Interaction	Other	Zyban(Amfebutamone Hydrochloride)	SS		ORAL
	150 MG, BID,						
	ORAL						

Date:03/05/03ISR Number: 4066228-8Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0393180A  
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG Twice		Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day Disability	6 DAY	Dyspnoea					

Date:03/05/03ISR Number: 4066230-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0398610A  
Age:17 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin	PS	Glaxosmithkline	
300MG per day		Drug Exposure During Pregnancy		Seroquel	C		

Maternal Drugs Affecting  
Foetus

Date:03/05/03ISR Number: 4066242-2Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0010662A  
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day 9 DAY	Amnesia		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization -		Cardiac Arrest					
Initial or Prolonged		Grand Mal Convulsion					
Other							

Date:03/05/03ISR Number: 4066243-4Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0010841A  
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day 9 DAY	Amnesia		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization -		Cardiac Arrest					
Initial or Prolonged		Grand Mal Convulsion					
Other							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/05/03ISR Number: 4068206-1Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #CTU 187921

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG DAILY		Dysarthria Grand Mal Convulsion Mental Impairment		Bupropion Sr 300mg Daily (Glaxosmithkline)	PS	Glaxosmithkline	
				Sertraline	C		
				Lamictal	C		
				Eskalith Cr	C		
				Seroquel	C		

Date:03/05/03ISR Number: 4071300-2Report Type:Expedited (15-DaCompany Report #US010627  
 Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 50 MG QD ORAL		Drug Interaction	Health	Provigil	PS		ORAL
40 MG QD ORAL		Headache	Professional	Celexa	SS		ORAL
100 MG QD ORAL		Migraine		Wellbutrin	SS		ORAL
				Synthroid	C		

Date:03/05/03ISR Number: 4072173-4Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #2003005856

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL Initial or Prolonged Other		Convulsion	Health Professional Company Representative	Geodon (Ziprasidone)	PS		ORAL
				Bupropion Hydrochloride	SS		
				Olanzapine	SS		
				All Other Non-Therapeutic Products	SS		

Date:03/06/03ISR Number: 4066862-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0398921A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Death	Intentional Misuse	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -	Somnolence	Professional	Heroin	SS		
Initial or Prolonged	Tachycardia					

Date:03/06/03ISR Number: 4066866-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0292305A  
Age:35 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Blood Alcohol Increased		Zyban	PS	Glaxosmithkline	ORAL
3000MG per						
Initial or Prolonged	Blood Pressure Systolic					
day	Increased					
	Chills					
	Confusional State					
	Drug Screen Positive					
	Intentional Misuse					
	Mydriasis					
	Respiratory Acidosis					
	Somnolence					
	Tachycardia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/06/03ISR Number: 4068853-7Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #CTU 188030

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300MG DAILY	Grand Mal Convulsion		Bupropriion Sr 300 Mg Glaxosmithkline	PS	Glaxosmithkline	ORAL
ORAL			Sertraline	C		
			Lamictal	C		
			Eskalith	C		
			Seroquel	C		

Date:03/07/03ISR Number: 4069513-9Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #CTU 188184

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 MG DAILY Initial or Prolonged	Convulsion Grand Mal Convulsion		Wellbutrin 150 Mg	PS		ORAL
ORAL			Synthroid	C		
			Lipitor	C		
			Norvasc	C		
			Phoslo	C		
			Reglan	C		
			Ibuprofen/Tylenol	C		

Date:03/07/03ISR Number: 4069585-1Report Type:Direct  
 Age:39 YR Gender:Female I/FU:I

Company Report #CTU 188131

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300 MG Initial or Prolonged MORNING 200	Convulsion Vertigo		Wellbutrin	PS		
MG 1:PM						

Date:03/07/03ISR Number: 4069592-9Report Type:Direct  
Age:29 YR Gender:Male I/FU:I

Company Report #CTU 188136

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening WELLBUTRIN 4 Hospitalization - DAYS Initial or Prolonged PAXIL 2 WEEKS Required Intervention to Prevent Permanent Impairment/Damage	Bipolar Ii Disorder		Wellbutrin  Paxil  Zyprexa Depakote Effexor	PS  SS  C C C		

Date:03/10/03ISR Number: 4068763-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0294006A  
Age:57 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150MG Hospitalization - Variable dose 10 DAY Initial or Prolonged 75MG Unknown  40MG Unknown  TOPICAL	Blood Creatine  Phosphokinase Increased  Left Ventricular Failure		Zyban  Aspirin  Simvastatin  Cinchocaine + Hydrocortisone	PS  C  C  C	Glaxosmithkline  Glaxosmithkline	ORAL  ORAL  ORAL  ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/10/03ISR Number: 4069959-9Report Type:Direct  
Age:50 YR Gender:Male I/FU:I

Company Report #CTU 188304

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness Nausea		Wellbutrin 150 Mg Catalytica Pharmaceuticals	PS	Catalytica Pharmaceuticals	

AURICULAR

(OTIC) 1 TWICE DAILY

AURICULAR

(OTIC)

Date:03/10/03ISR Number: 4070131-7Report Type:Direct  
Age:54 YR Gender:Female I/FU:I

Company Report #CTU 188263

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - BUPROPION 150 Initial or Prolonged MG BID		Agitation Formication Hallucination, Tactile Restlessness		Bupropion  Ambien Metoprolol Hctz Losartan Nexium Asa Simvastatin	PS  C C C C C C		

Date:03/10/03ISR Number: 4070273-6Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 188305

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other Required 150 MG TWICE		Ear Infection Gastrooesophageal Reflux		Wellbutrin S.R. 150 Mg Glaxo Wellcome	PS	Glaxo Wellcome	ORAL



Intervention to Disease  
DAILY ORAL  
Prevent Permanent Hiccups  
Impairment/Damage Pharyngitis  
Vomiting

Date:03/10/03ISR Number: 4070349-3Report Type:Direct Company Report #CTU 188354  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Zyban 150 Tbsr	PS	Tbsr	
ONCE PO QD X							
3 D THEN BID							

Date:03/10/03ISR Number: 4070358-4Report Type:Direct Company Report #CTU 188359  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort		Zyban 150 Mg Tbsr	PS	Tbsr	ORAL
ONCE PO QDSX							
3D THEN BID							

Date:03/11/03ISR Number: 4069296-2Report Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-12135786  
Age:45 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Condition Aggravated
Initial or Prolonged	Insomnia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Schizoaffective Disorder

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
			Aripiprazole	PS	Otsuka Pharmaceutical Company, Ltd.	ORAL
			Wellbutrin	SS		
			Prozac	SS		
			Topamax	C		
			Klonopin	C		

Date:03/11/03ISR Number: 4069473-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0395534A  
Age:19 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice	Initial or Prolonged per day 2 YR	Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Blood Pressure Systolic Increased		Sam E	C		
		Convulsion		Zoloft	C		
		Disturbance In Attention					
		Dizziness					
		Drug Screen Positive					
		Fatigue					
		Hallucination					
		Headache					
		Lethargy					
		Malaise					
		Medication Error					
		Pyrexia					
		Status Epilepticus					
		Tachycardia					
		Tongue Biting					
		Urinary Incontinence					
		Vision Blurred					

Date:03/11/03ISR Number: 4069476-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0398543A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Level Increased		Wellbutrin Risperdal	PS C	Glaxosmithkline	ORAL

Date:03/11/03ISR Number: 4069477-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0398716A

Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice per day	Death	Health Professional	Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN				Celexa	SS		
				Risperdal	C		
				Buspar	C		
				Synthroid	C	Glaxosmithkline	
				Ferrous Sulfate	C		
				Kdur	C	Glaxosmithkline	
				Lasix	C	Glaxosmithkline	
				Folic Acid	C		
				Mvi	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/11/03ISR Number: 4069479-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0399473A  
Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
		Intentional Misuse					
		Thermal Burn					

Date:03/11/03ISR Number: 4069480-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0399478A  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 250MG Per day 1 YR		Blood Creatine		Wellbutrin	PS	Glaxosmithkline	ORAL
		Phosphokinase Increased		Kaletra	C		
				Sustiva	C		
				Valcyte	C		
				Diflucan	C		
				Bactrim	C	Glaxosmithkline	
				Epivir	C	Glaxosmithkline	
				Ziagen	C	Glaxosmithkline	
				Elavil	C	Glaxosmithkline	
				Trazodone	C		
				Ativan	C		
				Percocet	C		
				Allegra	C		
				Neurontin	C		
				Albuterol	C	Glaxosmithkline	

Date:03/11/03ISR Number: 4069489-4Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0293848A  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN per day	150MG Twice	Alcohol Poisoning	Consumer	Zyban	PS	Glaxosmithkline	
		Drug Interaction					

UNKNOWN

Ethanol

SS

Date:03/11/03ISR Number: 4070395-XReport Type:Direct  
Age:26 YR Gender:Female I/FU:I

Company Report #CTU 188428

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Wellbutrin Sr 150mg	PS		ORAL
2 DAY ORAL		Dermatitis Atopic Irritability Obsessive Thoughts Pharmaceutical Product Complaint Psychiatric Symptom Rash Pruritic Reaction To Medical Agent Preservatives Suicidal Ideation					

Date:03/11/03ISR Number: 4072830-XReport Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 188491

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pharmaceutical Product Complaint		Generic Wellbutrin 75mg	PS		
75MG TWO BID							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/12/03ISR Number: 4070155-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0386663A

Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eye Disorder		Paxil	PS	Glaxosmithkline	ORAL
60MG Per day							
		Libido Decreased		Wellbutrin	SS	Glaxosmithkline	ORAL
100MG Per day							

Date:03/12/03ISR Number: 4070159-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0399790A

Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Intentional Misuse		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2 YR						

Date:03/12/03ISR Number: 4070165-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0294437A

Age:75 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardiac Failure		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	6 DAY						
		Congestive		Digoxin	C	Glaxosmithkline	
UNKNOWN	250MCG Per	Myocardial Infarction					
day							
				Aspirin	C	Glaxosmithkline	
UNKNOWN	75MG Per day						

Date:03/12/03ISR Number: 4070168-8Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0294671A

Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Aphasia		Zyban	PS	Glaxosmithkline	
43 DAY							

Other  
 Brain Scan Abnormal  
 Cerebral Arteritis  
 Grand Mal Convulsion  
 Headache  
 Mental Status Changes

Date:03/13/03ISR Number: 4071164-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0294329A  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG See dosage text	13 DAY	Blood Glucose Increased Convulsion Eye Disorder Hypokalaemia Malaise		Zyban  Tea Stablon	PS  SS C	Glaxosmithkline	ORAL  ORAL ORAL
2TAB per day							

Date:03/13/03ISR Number: 4076436-8Report Type:Expedited (15-DaCompany Report #HQWYE808303MAR03  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death		Cardiac Arrest Drug Interaction Drug Toxicity Lobar Pneumonia	Consumer	Phenergan (Promethazine Hydrochloride, Unspec) Prozac (Fluoxetine Hydrochloride, ) Ultram (Tramadol Hydrochloride, ) Wellbutrin	PS  SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Amfebutamone  
Hydrochloride, ) SS

Date:03/14/03ISR Number: 4072141-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0294420A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Periorbital Oedema		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY			Aspirin	C	Glaxosmithkline	ORAL
75MG Per day				Simvastatin	C		ORAL
20MG Per day				Atenolol	C		ORAL
50MG Per day				Ramipril	C		ORAL
1.25MG Per							
day				Nicotine	C	Glaxosmithkline	
UNKNOWN	4MG Four						
times per day							

Date:03/14/03ISR Number: 4072142-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0294422A  
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Homicidal Ideation		Zyban	PS	Glaxosmithkline	ORAL
150MG		Paranoia					
Variable dose	14 DAY						

Date:03/14/03ISR Number: 4072144-8Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0294464A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Depression	Consumer	Zyban	PS	Glaxosmithkline	
UNKNOWN	150MG Twice						



Hospitalization - Disease Recurrence  
 per day 80 DAY  
 Initial or Prolonged Suicide Attempt  
 UNKNOWN 20MG Unknown 1 DAY  
 Suspiciousness

Temazepam SS

Date:03/14/03ISR Number: 4072244-2Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0399254A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Paxil	PS	Glaxosmithkline	ORAL
Other		Anaphylactic Shock		Wellbutrin	SS	Glaxosmithkline	

Date:03/14/03ISR Number: 4074353-0Report Type:Direct Company Report #CTU 188768  
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Agitation		Bupropion	PS		
150MG BID	3 WK			Ambien	C		
Initial or Prolonged		Hallucination, Tactile Restlessness		Metoprolol	C		
				Hctz	C		
				Losartin	C		
				Nexium	C		
				Asa	C		
				Simvastatin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/17/03ISR Number: 4072846-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0398921A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged		Somnolence Tachycardia	Professional	Heroin	SS		

Date:03/17/03ISR Number: 4072850-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0400040A

Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Antepartum Haemorrhage Caesarean Section Complications Of Maternal Exposure To Therapeutic Drugs Gestational Diabetes Maternal Drugs Affecting Foetus Pregnancy Premature Separation Of Placenta		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:03/17/03ISR Number: 4072867-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0294481A

Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG per day 19 DAY	Abnormal Behaviour	Consumer	Zyban	PS	Glaxosmithkline	ORAL
		Aggression Anger Drug Interaction Judgement Impaired Laceration Mood Swings Thinking Abnormal		Alcohol	SS		ORAL

Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Eye Pain					
per day		Tremor		Medroxyprogesterone	C		
INTRAMUSCULAR		Vomiting					

Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eczema		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Rash					
per day	20 DAY			Sildenafil	C		ORAL
100MG As							
required							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/17/03ISR Number: 4074720-5Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #CTU 188835

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination		Wellbutrin	PS		ORAL
200-400 MG							
PILL ORAL							
				Lithium	C		
				Zyprexa	C		

Date:03/17/03ISR Number: 4074728-XReport Type:Direct  
 Age:16 YR Gender:Female I/FU:I

Company Report #CTU 188866

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Dysphonia		Bupropion Sr 400 Mg Gsk	PS	Gsk	
200 MG BID							
		Memory Impairment Syncope		Sudal	C		

Date:03/17/03ISR Number: 4074818-1Report Type:Direct  
 Age:60 YR Gender:Male I/FU:I

Company Report #CTU 188853

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 100 MG PO BID		Asthenia		Amantadine	PS		ORAL
Initial or Prolonged		Balance Disorder Fall Hallucinations, Mixed Serotonin Syndrome		Bupropion Olanzapine	SS C	100 Mg Po Bid	ORAL

Date:03/17/03ISR Number: 4077295-XReport Type:Expedited (15-DaCompany Report #2003009863  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Congenital Anomaly 120 MG (BID)	Complications Of Maternal Exposure To Therapeutic Drugs Congenital Anomaly Maternal Drugs Affecting Foetus Pregnancy	Health Professional Company Representative	Geodon (Ziprasidone) Bupropion Hydrochloride	PS SS
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Date:03/19/03ISR Number: 4074224-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0398921A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged		Somnolence Tachycardia	Professional	Heroin Cocaine Valium	SS SS SS		

Date:03/19/03ISR Number: 4074229-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0092202A  
Age:32 YR Gender:Female I/FU:F

Outcome	PT
Other	Aggression Anger Anhedonia Anorexia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day			Zyban	PS	Glaxosmithkline	ORAL
		Crying Depression Diarrhoea Disturbance In Attention Fatigue Feeling Abnormal Feeling Hot Headache Homicidal Ideation Irritability Lethargy Mood Altered Night Sweats Personality Change Screaming Suicidal Ideation Tinnitus Tremor				

Date:03/19/03ISR Number: 4076591-XReport Type:Direct  
Age:30 YR Gender:Female I/FU:I

Company Report #CTU 189083

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG BID./ 4-5 DAYS		Hypoaesthesia Pruritus Rash		Wellbutrin (Bupropion) 150 Mg Sr	PS		
40 MG DAILY				Citalopram (Celexa) 40 Mg	SS		

Date:03/19/03ISR Number: 4078378-0Report Type:Expedited (15-DaCompany Report #2003006006  
Age:58 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Death 40 MG  (DAILY), ORAL	Cardiomegaly  Coronary Artery  Atherosclerosis Mitral Valve Incompetence	Health  Professional  Company Representative	Geodon (Ziprasidone)   Bupropion Hydrochloride	PS   SS	ORAL   ORAL
450 MG (TID),  ORAL	Pulmonary Valve  Incompetence		Mirtazapine Lithium Carbonate Methadone Desloratadine Pseudoephedrine Hydrochloride Chlorphenamine	C C C C C C C	

Date:03/20/03ISR Number: 4074886-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0400724A  
Age:79 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1 DAY		Bronchial Obstruction		Wellbutrin	PS	Glaxosmithkline	ORAL
		Convulsion Dyspnoea					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/20/03ISR Number: 4074887-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0400777A  
 Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Eye Swelling		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	15 DAY	Face Oedema Oedema Peripheral Pruritus Skin Discolouration Urticaria		Yasmin	C		

Date:03/20/03ISR Number: 4074889-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0107035A  
 Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 3 WK		Abnormal Behaviour	Health	Zyban	PS	Glaxosmithkline	ORAL
37.5MG per day		Aggression Agitation Alcoholism Anxiety Completed Suicide Decreased Appetite Depression Dissociation Disturbance In Attention Drug Abuser Fear Feeling Abnormal Head Injury Headache Hypochondriasis Irritability Loss Of Consciousness Mania Mood Swings Obsessive-Compulsive Personality Disorder Palpitations	Professional	Chloroquine	C		



Personality Change  
Restlessness  
Vision Blurred

Date:03/20/03ISR Number: 4074898-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0294557A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG See dosage text	15	Anxiety Bulimia Nervosa Crying		Zyban	PS	Glaxosmithkline	ORAL
TRANSDERMAL		32 DAY Depression Fatigue Hyperphagia Insomnia Suicidal Ideation		Nicopatch	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/20/03ISR Number: 4074900-9Report Type:Expedited (15-DaCompany Report #CZ-GLAXOSMITHKLINE-B0294593A

Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	166 DAY	Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN		Drug Toxicity		Antidepressants	C		
UNKNOWN		Hypothermia		Carbamazepine	C		
UNKNOWN				Citalopram	C		
UNKNOWN	25MG per day			Methotrimeprazine	C		
UNKNOWN	1600MG per			Piracetam	C		
UNKNOWN	404 DAY			Promethazine	C		
UNKNOWN		42 DAY		Norethisterone Acetate + Oestradiol	C	Glaxosmithkline	

Date:03/20/03ISR Number: 4074901-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0295005A

Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice	Initial or Prolonged per day 10 DAY	Psychotic Disorder		Zyban	PS	Glaxosmithkline	ORAL

Date:03/20/03ISR Number: 4074902-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0295110A

Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG See		Arthralgia		Zyban	PS	Glaxosmithkline	ORAL

dosage text	18	DAY	Erythema			
3	DAY		Gastroenteritis	Ercefuryl	C	Glaxosmithkline
3	DAY		Pruritus	Debridat	C	
3	DAY		Urticaria	Ultra Levure	C	

Date:03/20/03ISR Number: 4080630-XReport Type:Expedited (15-DaCompany Report #L02-USA-03743-04  
 Age:31 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aspiration Cardiac Arrest Completed Suicide	Literature Health Professional	Citalopram (Citalopram Hydrobromide)	PS		ORAL
900 MG ONCE		Convulsion					
PO		Intentional Misuse		Bupropion	SS		ORAL
3000 MG ONCE		Somnolence					
PO				Amphetamine / Dextroamphetamine 30 Mg	SS		ORAL
10 UNITS ONCE							
PO				Lisinopril Trazodone	SS SS		

Date:03/21/03ISR Number: 4075751-1Report Type:Expedited (15-DaCompany Report #IE-GLAXOSMITHKLINE-B0293852A  
 Age:30 YR Gender:Male I/FU:F

Outcome	PT
Other	Abnormal Behaviour Agitation Confusional State Drug Interaction Memory Impairment

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Paranoia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG			Zyban	PS	Glaxosmithkline	ORAL
Variable dose	8 DAY					
UNKNOWN	2UNIT Unknown		Alcohol	SS		

Date:03/24/03ISR Number: 4076732-4Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0040676A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 18G per day Initial or Prolonged				Zyban	PS	Glaxosmithkline	ORAL
		Convulsion					
		Intentional Misuse Suicide Attempt					

Date:03/24/03ISR Number: 4082407-8Report Type:Expedited (15-DaCompany Report #US010627  
 Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 50 MG QD ORAL			Health	Provigil	PS		ORAL
40 MG QD ORAL			Professional	Celexa	SS		ORAL
100 MG QD ORAL				Welbutrin	SS		ORAL
				Synthroid	C		

Date:03/25/03ISR Number: 4077698-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0107035A  
 Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death	Abnormal Behaviour	Health	Zyban	PS	Glaxosmithkline	ORAL
3 WK	Aggression	Professional	Chloroquine	C		
37.5MG per	Agitation					
day	Alcoholism					
	Anxiety					
	Apathy					
	Cardiac Disorder					
	Completed Suicide					
	Decreased Appetite					
	Depression					
	Dissociation					
	Drug Abuser					
	Fear					
	Head Injury					
	Headache					
	Hypochondriasis					
	Irritability					
	Loss Of Consciousness					
	Mania					
	Mood Swings					
	Obsessive-Compulsive					
	Personality Disorder					
	Palpitations					
	Personality Change					
	Restlessness					
	Treatment Noncompliance					
	Vision Blurred					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/25/03ISR Number: 4080663-3Report Type:Direct  
Age:55 YR Gender:Male I/FU:I

Company Report #CTU 189456

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Aggression		Bupropion Sr	PS		
Initial or Prolonged	Agitation		Trazodone	SS		
	Blood Creatine		Olanzapine	SS		
	Phosphokinase Increased		Ziprasidone	C		
	Depression		Lithium	C		
	Drug Ineffective		Beclomethasone	C		
	Electrocardiogram Qt		Fexofenadine	C		
	Prolonged		Folic Acid	C		
	Medication Error		Naproxen	C		
	Mental Status Changes		Propranolol	C		
	White Blood Cell Count		Rabeprazole	C		
	Increased		Simvastatin	C		
			Thiamine	C		
			Trazodone	C		
			Ziprasidone	C		
			Lithium	C		
			Olanzapine	C		
			Bupropion	C		

Date:03/26/03ISR Number: 4083442-6Report Type:Expedited (15-DaCompany Report #200310607BCC  
Age:26 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Arthralgia	Consumer	Aleve (Naproxen			
440 MG, ONCE,	Difficulty In Walking	Other	Sodium)	PS		ORAL
ORAL	Pain In Extremity					
	Pruritus Generalised		Wellbutrin			
	Tendonitis		(Amfebutamone			
			Hydrochloride)	SS		

Date:03/27/03ISR Number: 4079588-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0401597A  
Age:5 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					

Other 150MG Twice			Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3	WK	Oedema Oliguria Rash				

Date:03/27/03ISR Number: 4079589-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0401801A  
 Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
Other UNKNOWN		Overdose					
		Suicide Attempt		Paxil	SS	Glaxosmithkline	
UNKNOWN				Xanax	SS		
UNKNOWN							

Date:03/27/03ISR Number: 4082800-3Report Type:Direct Company Report #USP 55741  
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr 100mg (Bupropion)	PS	Glaxo Wellcome	
Other		Medication Error		Wellbutrin Sr 150mg			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Bupropion)

SS

Glaxo Wellcome

Date:03/28/03ISR Number: 4080292-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0395460A

Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Creatine		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Phosphokinase Increased					
per day		Renal Disorder		Vioxx	C		ORAL

Date:03/28/03ISR Number: 4083683-8Report Type:Direct Company Report #CTU 189714

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia		Wellbutrin Sr 150mg			
Required		Chest Discomfort		Glaxo Smith Kline	PS	Glaxo Smith Kline	ORAL
SEE IMAGE							
Intervention to		Myalgia					
Prevent Permanent		Rash					
Impairment/Damage		Urticaria					

Date:03/31/03ISR Number: 4080832-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0264861A

Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Adrenal Adenoma		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	6 DAY	Headache		Alcohol	C		ORAL
		Hypertension					
		Malaise					
		Vision Blurred					

Date:03/31/03ISR Number: 4080836-XReport Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0294464A

Age:42 YR Gender:Male I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Depression	Consumer	Zyban	PS	Glaxosmithkline	
UNKNOWN	150MG	Twice					
Hospitalization -		Somnolence					
per day	80	DAY					
Initial or Prolonged		Suicide Attempt		Temazepam	SS		
UNKNOWN	20MG	Unknown					
		Suspiciousness					

Date:03/31/03ISR Number: 4080847-4Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0040676A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Aspiration	Health	Zyban	PS	Glaxosmithkline	ORAL
18G per day							
Hospitalization -		Coma	Professional				
Initial or Prolonged		Epilepsy					
		Intentional Misuse					
		Suicide Attempt					

Date:04/02/03ISR Number: 4083494-3Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0292587A  
Age:75 YR Gender:Male I/FU:F

Outcome	Duration	PT
Disability		Diarrhoea
		Dissociation
		Hallucination

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Poisoning Tremor Vomiting	Report Source	Product	Role	Manufacturer	Route
1UNIT Twice per day	8 DAY			Zyban	PS	Glaxosmithkline	ORAL

Date:04/02/03ISR Number: 4083502-XReport Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0295743A  
Age: Gender:Female I/FU:I

Outcome Dose Other	Duration	PT Drug Interaction Epilepsy	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	19 DAY			Zyban	PS	Glaxosmithkline	ORAL
25MG per day				Trimipramine	SS		ORAL
5MG Per day				Methotrimeprazine	SS		ORAL
300MG Per day				Moclobemide	SS		ORAL
2MG per day				Alprazolam	C		ORAL

Date:04/02/03ISR Number: 4083504-3Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0295748A  
Age:32 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT Agitation Clonic Convulsion Overdose Suicide Attempt	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 40TAB Initial or Prolonged cumulative dose	1 DAY			Zyntabac	PS	Glaxosmithkline	ORAL

Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphasia		Zyban	PS	Glaxosmithkline	ORAL
43 DAY		Cerebral Arteritis Cerebral Haemorrhage Convulsion Nausea Personality Change Vomiting					

Date:04/02/03ISR Number: 4086302-XReport Type:Expedited (15-DaCompany Report #B0294984A

Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Coagulopathy Hepatic Failure Renal Impairment	Foreign Literature Health	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		
10 DAY		Sepsis	Professional	Carbimazole (Formulation Unknown) (Carbimazole) Propranolol Hydrochloride	SS  C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/02/03ISR Number: 4087906-0Report Type:Expedited (15-DaCompany Report #B0294985A

Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaemia Aspartate Aminotransferase	Foreign Literature Health	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL		Increased Asthenia Blood Lactate Dehydrogenase Increased Haematuria Haemolytic Uraemic Syndrome Headache Jaundice Menometrorrhagia Petechiae Red Blood Cell Schistocytes Present Red Blood Cell Sedimentation Rate Increased Thrombotic Thrombocytopenic Purpura Uterine Leiomyoma	Professional				

Date:04/03/03ISR Number: 4084091-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0391256A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour Conversion Disorder Convulsion Drug Abuser Feeling Abnormal Mania Medication Error	Health Professional	Wellbutrin Zoloft	PS SS	Glaxosmithkline	NASAL NASAL

Date:04/03/03ISR Number: 4084093-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0402194A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams	Consumer	Bupropion	PS	Glaxosmithkline	ORAL
		Convulsion		Tegretol	SS		ORAL
		Drug Interaction		Nicoderm	C	Glaxosmithkline	OTHER
		Dyskinesia					
		Epistaxis					
		Head Injury					
		Laceration					
		Local Swelling					

Date:04/03/03ISR Number: 4084095-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0402428A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG Twice Initial or Prolonged per day		Stevens-Johnson Syndrome		Wellbutrin	PS	Glaxosmithkline	ORAL
				Paxil	SS	Glaxosmithkline	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/03/03ISR Number: 4087135-0Report Type:Direct  
Age:22 YR Gender:Male I/FU:I

Company Report #CTU 190119

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent ORAL Impairment/Damage	150 MG BID	Diabetes Mellitus		Wellbutrin Sr 150 Mg Glaxosmithkline	PS	Glaxosmithkline	ORAL
				Insulin Pump	C		

Date:04/04/03ISR Number: 4085023-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0393425A  
Age:17 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	20 DAY	Complex Partial Seizures		Wellbutrin	PS	Glaxosmithkline	ORAL
	20 DAY	Epileptic Aura		Gabitril	SS		ORAL
		Road Traffic Accident Status Epilepticus Tonic Clonic Movements					

Date:04/04/03ISR Number: 4085025-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0402014A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Abortion Spontaneous		Wellbutrin	PS	Glaxosmithkline	
		Drug Exposure During Pregnancy Maternal Drugs Affecting Foetus					

Date:04/04/03ISR Number: 4085027-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0402277A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG Per day		Drug Interaction		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged 50MG Per day		Headache		Provigil	SS		ORAL
Disability 40MG Per day		Migraine		Celexa	SS		ORAL
				Synthroid	C	Glaxosmithkline	

15 YR

Date:04/04/03ISR Number: 4085028-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0402520A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hepatic Cirrhosis Hepatitis C		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:04/04/03ISR Number: 4085030-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0402594A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Level Above Therapeutic		Wellbutrin	PS	Glaxosmithkline	ORAL
				Paxil	SS	Glaxosmithkline	ORAL
				Benadryl	C	Glaxosmithkline	
				Doxylamine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/04/03ISR Number: 4085032-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0402679A  
 Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 200MG Twice		Choreoathetosis		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	MON	Drug Interaction					
75MG Twice		Therapeutic Agent		Effexor Xr	SS		
per day		Toxicity					
40MG Per day				Pepcid	C		
150MCG Per day				Synthroid	C	Glaxosmithkline	
1.5MG At night				Klonopin	C		
2TAB Three times per day				Lorcet 10/650	C		
25MG Per day				Vioxx	C		
25MG Per day				Hctz	C		

Date:04/04/03ISR Number: 4085035-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0402911A  
 Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion Encephalopathy Intentional Misuse Suicide Attempt Tachycardia		Wellbutrin	PS	Glaxosmithkline	



Date:04/04/03ISR Number: 4085132-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0402860A  
Age:84 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Bradycardia		Digoxin	PS	Glaxosmithkline	ORAL
Initial or Prolonged	Cardiac Failure		Bupropion			
	Congestive		Hydrochloride	SS	Glaxosmithkline	ORAL
	Drug Toxicity		Insulin	C		
	Pulmonary Oedema		Levothyroxine Sodium	C	Glaxosmithkline	
	Ventricular Extrasystoles		Linezolid	C		
			Metoclopramide Hcl	C	Glaxosmithkline	
			Zolpidem Tartrate	C		

Date:04/04/03ISR Number: 4088056-XReport Type:Direct Company Report #CTU 190286  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Other	Rash		Wellbutrin Er	PS		
100MG BID			Ibuprofen	C		

Date:04/07/03ISR Number: 4085773-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0403090A  
Age:39 YR Gender:Female I/FU:F

Outcome	PT
Other	Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus Pregnancy

Freedom Of Information (FOI) Report

Stillbirth

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
			Wellbutrin	PS	Glaxosmithkline	
			Paxil	SS	Glaxosmithkline	ORAL

Date:04/07/03ISR Number: 4085775-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0403504A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Dysphemia					
300MG Per day		Energy Increased Speech Disorder		Zoloft	C		

Date:04/07/03ISR Number: 4085776-8Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0099812A  
 Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	
Disability		Alopecia					
150MG Twice		Arthralgia					
Other		Asthenia		Quinapril	C		
per day		Balance Disorder		Trisequens	C	Glaxosmithkline	
		Blood Pressure Increased		Ketoconazole	C		
		Circulatory Collapse		Alcohol	C		
		Dehydration					
		Difficulty In Walking					
		Disturbance In Attention					
		Dizziness					
		Drug Withdrawal Syndrome					
		Dry Skin					
		Feeling Abnormal					
		Food Allergy					
		Headache					
		Hypersensitivity					
		Loss Of Libido					
		Malaise					
		Memory Impairment					

Pain In Extremity  
Pain Of Skin  
Pallor  
Palpitations  
Paraesthesia  
Pharyngeal Oedema  
Sneezing  
Tongue Dry  
Tongue Oedema

Date:04/07/03ISR Number: 4085780-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0295862A

Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged dosage text	34 DAY	Oedema Peripheral Psoriasis		Zyban  Elisor Questran Puva Daivonex Diprosone	PS  C C C C C	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/07/03ISR Number: 4085781-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0296281A  
 Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other	300MG per day	16 DAY	Urticaria	Zyban	PS	Glaxosmithkline	ORAL
	100MG per day			Sildenafil	C		ORAL

Date:04/08/03ISR Number: 4086290-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0296280A  
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other	150MG Unknown	15 DAY	Blood Creatine	Zyban	PS	Glaxosmithkline	ORAL
			Phosphokinase Increased				
			Capillary Disorder				
			Dermatomyositis				
			Erythema				
			Myalgia				

Date:04/08/03ISR Number: 4086294-3Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0038185A  
 Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death	150MG Unknown	WK	Myocardial Infarction	Zyban	PS	Glaxosmithkline	ORAL
			Renal Embolism				

Date:04/08/03ISR Number: 4090982-2Report Type:Direct Company Report #CTU 190441  
 Age:86 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -	100 MG (2)		Convulsion	Wellbutrin	PS		
Initial or Prolonged	DAILY	6 MON					

Date:04/09/03ISR Number: 4091030-0Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 190498

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	100 MG TID	Convulsion		Tramadol 50 Mg	PS		ORAL
Hospitalization - ORAL	Initial or Prolonged 100 MG TID			Bupropion 100 Mg	SS		ORAL
ORAL							

Date:04/10/03ISR Number: 4088144-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0404175A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Twice per day	Abdominal Distension Abdominal Pain Upper		Lotronex Wellbutrin	PS SS	Glaxosmithkline Glaxosmithkline	ORAL ORAL
		Bladder Cancer					
		Flatulence		Tenormin	C		
		Irritable Bowel Syndrome		Klonopin	C		
		Tinnitus		Claritin	C		
				Serzone	C		
				Estratest	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/10/03ISR Number: 4088150-3Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0296475A  
Age:39 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 7500MG Single Initial or Prolonged dose	Overdose		Zyban	PS	Glaxosmithkline	ORAL
UNKNOWN	Supraventricular Extrasystoles		Alcohol	C		
	Ventricular Extrasystoles					

Date:04/10/03ISR Number: 4088153-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0296761A  
Age:33 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Unknown 15 DAY Initial or Prolonged	Intestinal Obstruction		Zyban	PS	Glaxosmithkline	ORAL
			Hivid	C		
			Retrovir	C	Glaxosmithkline	

Date:04/10/03ISR Number: 4093521-5Report Type:Expedited (15-DaCompany Report #B0294984A  
Age:41 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death Hospitalization - Initial or Prolonged 10 DAY	Activated Partial Thromboplastin Time Prolonged	Foreign Literature Health	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		
5 YR	Cholestasis Coagulopathy	Professional	Carbimazole(Carbimazole)	SS		
	Dialysis		Propranolol			
	Disseminated		Hydrochloride	C		
	Intravascular Coagulation		Paracetamol	C		
	Encephalopathy		Ethanol	C		
	Haemoglobin Decreased					
	Hepatic Failure					
	Hepatic Necrosis					
	Hepatorenal Syndrome					

Hypokalaemia  
Hyponatraemia  
Hypotension  
Intra-Abdominal  
Haemorrhage  
Metabolic Acidosis  
Mouth Haemorrhage  
Oliguria  
Prothrombin Time  
Prolonged  
Pulmonary Haemorrhage  
Pulmonary Oedema  
Rectal Haemorrhage  
Renal Impairment  
Sepsis

Date:04/10/03ISR Number: 4093835-9Report Type:Expedited (15-DaCompany Report #A208038  
Age:17 YR Gender:Male I/FU:F

Outcome  
Life-Threatening  
Hospitalization -  
Initial or Prolonged  
Required  
Intervention to

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Prevent Permanent Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2400 MG, ORAL		Acidosis Coma	Literature Health	Ziprasidone (Ziprasidone)	PS		ORAL
ORAL		Electrocardiogram Qrs Complex Prolonged Electrocardiogram Qt Corrected Interval	Professional	Wellbutrin (Bupropion) (Bupropion Hydrohloride)	SS		ORAL
7.5 MG, ORAL		Prolonged Electrocardiogram Qt		Klonopin (Clonazepam0	SS		ORAL
2 MG, ORAL		Prolonged Hyperventilation		Ativan (Lorazepam) (Lorazepam)	SS		ORAL
		Intentional Misuse Lethargy Somnolence					

Date:04/11/03ISR Number: 4088863-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0404161A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Carcinoid Syndrome Pheochromocytoma		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:04/11/03ISR Number: 4088871-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0295110A

Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG See Other dosage text	18 DAY	Abdominal Pain	Health	Zyban	PS	Glaxosmithkline	ORAL
3TAB per day	4 DAY	Arthralgia	Professional				
4 DAY		Blood Immunoglobulin E Increased		Ercefuryl Debridat	C C	Glaxosmithkline	ORAL ORAL



Increased  
 Diarrhoea  
 Erythema  
 Gastroenteritis  
 Pruritus  
 Urticaria  
 Vomiting  
 White Blood Cell Count  
 Increased

Date:04/11/03ISR Number: 4088878-5Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0296637A  
 Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aptyalism		Zyban	PS	Glaxosmithkline	ORAL
Other		Electrocardiogram Qt Prolonged		Norethisterone + Ethinylloestradiol	C		ORAL
		Intentional Misuse Somnolence Speech Disorder Tachycardia Therapeutic Agent Toxicity Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/11/03ISR Number: 4088880-3Report Type:Expedited (15-DaCompany Report #CH-GLAXOSMITHKLIN-B0296745A  
 Age:49 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150MG Per day	Duration 8 DAY	Depression Consumer	Zyban	PS	Glaxosmithkline	ORAL
40MG per day		Disease Recurrence	Citalopram	SS		ORAL
.5MG Twenty times per day		Drug Interaction	Flupentixol + Melitracen Hydrochloride	SS		ORAL
.1MG per day			Thyroxine Sodium	C	Glaxosmithkline	ORAL

Date:04/11/03ISR Number: 4088885-2Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLIN-D0040836A  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Duration	Ageusia Anosmia Myalgia	Zyban	PS	Glaxosmithkline	ORAL

Date:04/11/03ISR Number: 4091130-5Report Type:Direct Company Report #CTU 190714  
 Age:43 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to 1Q AM Prevent Permanent Impairment/Damage	Duration	Deafness Unilateral	Wellburtin 100mg Glaxosmithkline	PS	Glaxosmithkline	
			Alprazolam	C		

Date:04/11/03ISR Number: 4091756-9Report Type:Direct Company Report #CTU 190755  
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Cardio-Respiratory Arrest		Wellbutrin 100mg Po			
Intervention to		Grand Mal Convulsion		7am 150mg Po 830hs	PS		ORAL
150MG PO 7PM							
Prevent Permanent							
; 150MG PO							
Impairment/Damage							
830PM							

Date:04/14/03ISR Number: 4092094-0Report Type:Direct Company Report #CTU 190836  
 Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin Sr 150 Mg			
Other				Glaxo Smith Kline	PS	Glaxo Smith Kline	ORAL
150 MG BID							
ORAL							

Date:04/14/03ISR Number: 4092097-6Report Type:Direct Company Report #CTU 190838  
 Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Hypersensitivity		Wellbutrin Sr 150 Mg			
Initial or Prolonged				Glaxo Smith Kline	PS	Glaxo Smith Kline	ORAL
150 MG BID							
ORAL							
				Zoloft	C		
				Seroquel	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/14/03ISR Number: 4092136-2Report Type:Direct  
Age:41 YR Gender:Female I/FU:I

Company Report #CTU 190772

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin 150mg			
				Glaxo Smith Kline	PS	Glaxo Smith Kline	ORAL
150MG BID							
ORAL							

Date:04/14/03ISR Number: 4092138-6Report Type:Direct  
Age:21 YR Gender:Female I/FU:I

Company Report #CTU 190775

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dyspnoea		Wellbutrin Sr 160mg			
Initial or Prolonged		Hypersensitivity		Glaxo Smith Kline	PS	Glaxo Smith Kline	ORAL
150MG BID							
		Urticaria					
ORAL							

Date:04/14/03ISR Number: 4092327-0Report Type:Direct  
Age:31 YR Gender:Female I/FU:I

Company Report #CTU 190805

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Arthralgia		Wellbutrin	PS		
Initial or Prolonged		Face Oedema					
Other		Joint Swelling					
		Oedema Peripheral					
		Pruritus					
		Tremor					

Date:04/14/03ISR Number: 4095197-XReport Type:Expedited (15-DaCompany Report #A0403475A  
Age:17 YR Gender:Male I/FU:I

Company Report #A0403475A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Acidosis	Literature	Wellbutrin Sr			
Initial or Prolonged		Arrhythmia	Health	Tablet-Controlled			

SEE DOSAGE	Drug Toxicity Electrocardiogram Qrs	Professional	Release (Bupropion Hydrochloride)	PS	ORAL
TEXT/ORAL	Complex Prolonged				
2400 MG/ SINGLE DOSE/ ORAL	Electrocardiogram Qt Prolonged Lethargy Overdose  Somnolence		Ziprasidone Hcl (Formulation Unknown)(Ziprasidone Hcl)	SS	ORAL
7.5 MG / SINGLE DOSE / ORAL			Clonazepam (Formulation Unknown)	SS	ORAL
2 MG / SINGLE DOSE /			Lorazepam (Formulation Unknown) (Lorazepam)	SS	

Date:04/15/03ISR Number: 4090707-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0385531A  
Age:38 YR Gender:Female I/FU:F

Outcome PT  
Hospitalization - Chest Discomfort  
Initial or Prolonged Heart Rate Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hyperhidrosis  
Ventricular Tachycardia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
400MG Per day			Wellbutrin	PS	Glaxosmithkline	ORAL
.5MG At night			Effexor Xr	C		
			Xanax	C		ORAL
75MG As			Atarax	C		ORAL
required						

Date:04/15/03ISR Number: 4090711-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0392586A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Laboratory Test Interference Pheochromocytoma		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:04/15/03ISR Number: 4090713-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0400724A  
Age:79 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bronchial Obstruction	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
1 DAY		Convulsion Dyspnoea Mucous Membrane Disorder	Professional				

Date:04/15/03ISR Number: 4090714-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0404165A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							

Drug Exposure During  
Pregnancy  
Maternal Drugs Affecting  
Foetus

Date:04/15/03ISR Number: 4090715-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0404554A  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300MG per day		Drug Exposure During		Wellbutrin	PS	Glaxosmithkline	ORAL
		Pregnancy					
		Pregnancy					
		Pregnancy Induced					
		Hypertension					
		Proteinuria					

Date:04/15/03ISR Number: 4090742-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0403696A  
Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
		Deafness	Consumer	Wellbutrin Sr	PS	Glaxosmithkline	
		Dysphonia		Paxil	SS	Glaxosmithkline	ORAL
		Speech Disorder		Remeron	SS		
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/18/03ISR Number: 4093145-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0380858A

Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dacryostenosis Acquired		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Single							
dose	2	DAY	Energy Increased				
			Face Oedema	Sudafed	C	Glaxosmithkline	
			Flushing				
			Headache				
			Pain In Jaw				
			Pruritus				
			Rash Erythematous				

Date:04/18/03ISR Number: 4093147-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0397981A

Age:58 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Acute Respiratory	Other	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day			Distress Syndrome				
			Bronchospasm	Prozac	SS		ORAL
40MG Per day							
			Cardiac Arrest	Ultram	SS		ORAL
50MG As							
required	13	DAY	Cardiac Disorder				
			Cardiomegaly	Phenergan With			
			Drug Toxicity	Codeine	SS		ORAL
1TSP As							
required	12	DAY	Fall				
			Lobar Pneumonia	Chlorpheniramine	SS		
			Lung Consolidation	Ultracet	SS		
			Lung Disorder	Vioxx	C		
			Musculoskeletal Pain	Reglan	C	Glaxosmithkline	
			Neck Pain	Theodur	C		
300MG Twice							
per day			Pernicious Anaemia				
			Pleural Disorder	Prednisone	C		



Pneumonia  
Renal Disorder  
Thyroiditis

Flovent C  
Singulair C  
Premarin C  
Prevacid C  
Rhinocort C  
Klonopin C  
Vioxx C  
Combivent C

Glaxosmithkline

RESPIRATORY

(INHALATION) 2PUFF Three  
times per day

Neurontin C  
Baclofen C  
Vitamin B12 C  
Darvocet C

Glaxosmithkline

1TAB Three  
times per day

Levaquin C

Date:04/18/03ISR Number: 4093158-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0297338A  
Age:40 YR Gender:Female I/FU:I

Outcome PT  
Other Ageusia  
Confusional State  
Depression  
Excitability  
Hyposmia  
Insomnia  
Mood Altered

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

				Muscle Twitching					
				Somnolence					
					Report Source	Product	Role	Manufacturer	Route
Dose	Duration					Zyban	PS	Glaxosmithkline	ORAL
150MG See									
dosage text	29	DAY							

Date:04/18/03ISR Number: 4093159-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0297365A  
 Age:27 YR Gender:Female I/FU:F

					Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	PT				Zyban	PS	Glaxosmithkline	ORAL
Dose				Abdominal Pain Upper					
Other	150MG Unknown	4	DAY			Eugynon	C		ORAL
1Z Per day				Anorexia					
				Depression					
				Disturbance In Attention					
				Fatigue					
				Headache					
				Irritability					
				Nausea					

Date:04/18/03ISR Number: 4093161-8Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0038838A  
 Age:36 YR Gender:Male I/FU:F

					Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	PT				Zyban	PS	Glaxosmithkline	ORAL
Dose				Aggression					
Disability	150MG Per day	6	WK			Seroxat	SS	Glaxosmithkline	
Other	UNKNOWN			Bipolar Disorder					
				92	DAY				
				Crying					
				Diarrhoea					
				Disturbance In Attention					
				Gastrointestinal Disorder					
				Hyperhidrosis					
				Hypomania					
				Illusion					
				Migraine					
				Nausea					

Nervousness  
 Sleep Disorder  
 Suicidal Ideation  
 Tachycardia  
 Vertigo  
 Weight Decreased

Date:04/22/03ISR Number: 4094756-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0405398A

Age:43 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 450MG Three Initial or Prolonged times per day	Lethargy		Wellbutrin	PS	Glaxosmithkline	ORAL
	Prescribed Overdose					
	Sensation Of Heaviness		Trazodone	C		
	Tremor		Zyprexa	C		
	Visual Disturbance		Zolpidem	C		
			Colace	C		
			Iron	C		
			Vicodin	C		
			Motrin	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/22/03ISR Number: 4094792-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0405398A  
 Age:43 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 450MG Three Initial or Prolonged times per day	Lethargy	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
	Prescribed Overdose	Professional				
	Sensation Of Heaviness		Trazodone	C		
	Tremor		Zyprexa	C		
	Visual Disturbance		Zolpidem	C		
			Colace	C		
			Iron	C		
			Vicodin	C		
			Motrin	C	Glaxosmithkline	

Date:04/23/03ISR Number: 4095787-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0145759A  
 Age:49 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 100MG Per day 2 YR	Dizziness Postural		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
	Hypertension		Norvasc	C		
	Lethargy		Premarin	C		
	Myocardial Ischaemia		Prevacid	C		
	Somnolence		Multivitamin	C		
	Ventricular Hypertrophy					

Date:04/23/03ISR Number: 4095791-6Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0405380A  
 Age: Gender: I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Difficulty In Walking		Wellbutrin	PS	Glaxosmithkline	ORAL
	Myalgia					
	Rhabdomyolysis					

Date:04/23/03ISR Number: 4102151-8Report Type:Expedited (15-DaCompany Report #2003016733  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Murder	Consumer	Zoloft (Sertraline) Bupropion Hydrochloride (Bupropion Hydrochloride)	PS  SS		

Date:04/24/03ISR Number: 4100490-8Report Type:Direct  
Age:27 YR Gender:Female I/FU:I

Company Report #CTU 191554

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 50 MG DAY Intervention to ORAL Prevent Permanent Impairment/Damage 150 MG DAY  OTHER		Aggression  Agitation  Depression Irritability  Mood Altered  Sexual Dysfunction		Zoloft 50 Mg Pfizer   Wellbutrin 150 Mg Glaxo	PS  SS	Pfizer  Glaxo	ORAL  OTHER

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/25/03ISR Number: 4097721-XReport Type:Direct  
Age:31 YR Gender:Male I/FU:I

Company Report #USP 51576

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Wellbutrin Sr(Bupropion)	PS	Wellcome	

Date:04/25/03ISR Number: 4101237-1Report Type:Direct  
Age:20 YR Gender:Male I/FU:I

Company Report #USP 55812

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin Sr 150	PS	Glaxo	

Date:04/25/03ISR Number: 4117226-7Report Type:Periodic  
Age:19 YR Gender:Female I/FU:I

Company Report #NSADSS2002032542

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Literature	Ultram (50 Mg Tablet) (Tramadol Hydrochloride)	PS		ORAL
ORAL		Respiratory Depression	Health				
		Therapeutic Response	Professional				
		Increased		Metoclopramide(Metoclopramide)	SS		ORAL
ORAL				Bupropion (Amfebutamone)	SS		ORAL

Date:04/28/03ISR Number: 4101197-3Report Type:Direct  
Age:41 YR Gender:Female I/FU:I

Company Report #CTU 191710

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Balance Disorder		Paxil 20 Mg 1x A			
Other		Bedridden		Day ?	PS		
		Decreased Activity		Wellbutrin Sr 100 Mg			
		Difficulty In Walking		1x A Day	SS		
		Eye Movement Disorder					

Headache  
Medication Error  
Migraine  
Movement Disorder  
Nausea  
Pharmaceutical Product  
Complaint  
Photophobia  
Suicidal Ideation  
Vertigo  
Vomiting

Date:04/28/03ISR Number: 4101671-XReport Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 191740

Outcome PT  
Other Aggression  
Confusional State  
Disorientation  
Dizziness  
Nightmare  
Sexual Dysfunction  
Social Avoidant Behaviour  
Weight Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Weight Loss Poor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
ZOLOFT 1 X			Zoloft 25 Mg	PS		ORAL
PER ORAL						
WELLBUTRIN 1			Wellbutrin 100 Mg	SS		ORAL
X PER ORAL						

Date:04/29/03ISR Number: 4099833-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0406259A  
 Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
Hospitalization -		Facial Bones Fracture					
per day							
Initial or Prolonged							
Disability							
Other							

Date:04/29/03ISR Number: 4099835-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0406391A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Arthralgia		Bupropion	PS	Glaxosmithkline	ORAL
		Asthenia					
		Convulsion					
		Dizziness					
		Dyspnoea					
		Hyperaesthesia					
		Hypersensitivity					
		Hypoxia					
		Immune System Disorder					
		Muscle Spasms					
		Myalgia					
		Nightmare					



Throat Tightness  
Tremor  
Vestibular Disorder

Date:04/29/03ISR Number: 4099845-XReport Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0297272A  
Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day 5 DAY	Cardiomyopathy		Zyntabac	PS	Glaxosmithkline	ORAL
		Erythema Multiforme					
		Hypertension					

Date:04/30/03ISR Number: 4100926-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0406365A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Anxiety		Wellbutrin	PS	Glaxosmithkline	
		Chest Pain					
		Hypotension		Vicodin	C		
		Lung Disorder					
		Myocardial Infarction					
		Palpitations					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/30/03ISR Number: 4103722-5Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #CTU 191975

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Celexa 40mg 40mg	PS		ORAL
11 ORAL							
		Drug Ineffective		Wellbutrin Sr 150 Mg	SS		ORAL
150 2 ORAL							
		Feeling Abnormal					
		Loss Of Libido					
		Neurological Symptom					
		Oral Intake Reduced					
		Weight Increased					

Date:04/30/03ISR Number: 4105818-0Report Type:Expedited (15-DaCompany Report #A0372195A  
 Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Carbon Monoxide	Literature	Wellbutrin			
Initial or Prolonged		Blood Chloride	Health	Unspecified Tablet			
Other		Blood Creatinine	Professional	(Bupropion			
		Blood Glucose	Other	Hydrochloride)	PS		ORAL
4500 MG/							
SINGLE		Blood Potassium					
		Blood Sodium					
DOSE/ORAL		Blood Urea		Topiramate Tablet			
		Depressed Level Of		(Topiramate)	SS		ORAL
3000		Consciousness					
MG/SINGLE		Hypertension					
DOSE/ORAL		Metabolic Acidosis		Glimepiride			
		Overdose		(Formulation			
		Tachycardia		Unknown)			
20 MG/SINGLE				(Glimepiride)	SS		
DOSE				Fluoxetine			
				Hydrochloride			

600 MG/SINGLE

DOSE

(Formulation  
Unknown),  
(Fluoxetine SS

200 MG/SINGLE

DOSE

Enalapril Maleate  
(Formulation  
Unknown) (Enalapril  
Maleate) SS

Date:05/01/03ISR Number: 4105194-3Report Type:Direct  
Age:45 YR Gender:Male I/FU:I

Company Report #CTU 192111

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG BID PO		Hyperreflexia		Wellbutrin Sr 150mg	PS		ORAL
Other Required Intervention to Prevent Permanent Impairment/Damage		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/02/03ISR Number: 4107055-2Report Type:Expedited (15-DaCompany Report #2003017811  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	20 MG	Anaphylactic Shock	Foreign Health Professional	Atorvastatin (Atorvastatin)	PS		ORAL
(DAILY), ORAL				Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		ORAL
300 MG							
(DAILY), ORAL							

Date:05/05/03ISR Number: 4103016-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0397981A  
 Age:58 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice	Acute Respiratory Distress Syndrome	Other	Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Asthma		Prozac	SS		ORAL
40MG Per day		Cardiac Arrest		Ultram	SS		ORAL
50MG As		Cyanosis					
required	13 DAY	Drug Toxicity Fall		Phenergan With Codeine	SS		ORAL
1TSP As		Lobar Pneumonia					
required	12 DAY	Lung Consolidation		Chlorpheniramine	SS		
				Ultracet	SS		
				Vioxx	C		
				Reglan	C	Glaxosmithkline	
				Theodur	C		
300MG Twice							

per day

Prednisone	C	
Flovent	C	Glaxosmithkline
Singulair	C	
Premarin	C	
Prevacid	C	
Rhinocort	C	
Klonopin	C	
Vioxx	C	
Combivent	C	

RESPIRATORY

(INHALATION) 2PUFF Three

times per day

Neurontin	C	
Baclofen	C	
Vitamin B12	C	Glaxosmithkline
Darvocet	C	

1TAB Three

times per day

Levaquin	C	
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Date:05/05/03ISR Number: 4103022-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0406924A

Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Colitis		Wellbutrin	PS	Glaxosmithkline	ORAL
4 MON		Diarrhoea		Klonopin	C		
		Intestinal Obstruction		Trileptal	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/05/03ISR Number: 4105686-7Report Type:Direct  
 Age:74 YR Gender:Female I/FU:I

Company Report #CTU 192292

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required	150 MG BID PO	Condition Aggravated		Bupropion	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage		Tremor					

Date:05/05/03ISR Number: 4107962-0Report Type:Expedited (15-DaCompany Report #HQWYE808303MAR03  
 Age:58 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Exposure	Health	Phenergan			
Other		Accidental Overdose	Professional	(Promethazine Hydrochloride, Unspec)	PS		
		Cardiac Arrest		Prozac (Fluoxetine Hydrochloride, )	SS		
		Cardiomegaly		Ultram (Tramadol Hydrochloride, )	SS		
		Drug Interaction		Wellbutrin (Amfebutamone Hydrochloride, )	SS		
		Drug Toxicity					
		Lobar Pneumonia					
		Lung Disorder					
		Medication Error					
		Oedema					
		Renal Disorder					
		Toxicologic Test Abnormal					

Date:05/06/03ISR Number: 4103911-XReport Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0298177A  
 Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 12 DAY		Anxiety		Zyntabac	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Extrasystoles		Verapamil	C		ORAL
		Ventricular Extrasystoles					

Date:05/06/03ISR Number: 4103913-3Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0298247A  
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence		Bupropion			
		Euphoric Mood		Hydrochloride	PS	Glaxosmithkline	
UNKNOWN	150MG	As					
		Libido Increased					
directed							
		Personality Change		Cannabis	C		
UNKNOWN							

Date:05/07/03ISR Number: 4104935-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0407308A  
Age:15 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG per day							
Initial or Prolonged		Status Epilepticus					

Date:05/07/03ISR Number: 4104938-4Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0293561A  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT
Hospitalization -		Adverse Drug Reaction
Initial or Prolonged		Arthralgia
		Chest Discomfort

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG		Chest Pain Dizziness Dyspnoea					
		Hypoaesthesia		Zyban	PS	Glaxosmithkline	ORAL
Variable dose	7 DAY	Mediastinal Disorder					
UNKNOWN		Pallor Palpitations		Diclofenac + Misoprostol	C		
		Paraesthesia					

Date:05/07/03ISR Number: 4104940-2Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0298038A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Unknown	3 WK	Blood Pressure Decreased	Consumer	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged UNKNOWN		Burning Sensation 1 DAY		Anesthetic	SS		
		Face Oedema Flushing Rash Urticaria					

Date:05/07/03ISR Number: 4104942-6Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0298053A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN	150MG Unknown	Aggression		Zyban	PS	Glaxosmithkline	
		Arrhythmia Cardiac Disorder Insomnia Nausea Sensation Of Foreign Body					



Date:05/07/03ISR Number: 4104946-3Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0298454A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening UNKNOWN		Chest Discomfort		Zyban	PS	Glaxosmithkline	
Hospitalization - Initial or Prolonged		Mental Impairment Neoplasm Personality Change					

Date:05/07/03ISR Number: 4107641-XReport Type:Direct Company Report #CTU 192460

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hypersensitivity		Wellbutrin Sr 150 Mg Gsk	PS	Gsk	ORAL
150 MG QD							
ORAL							

Date:05/07/03ISR Number: 4109692-8Report Type:Expedited (15-DaCompany Report #HQWYE808303MAR03

Age:58 YR Gender:Female I/FU:F

Outcome	PT
Death	Accidental Exposure Cardiac Arrest Cardiomegaly Drug Interaction

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Screen Positive Drug Toxicity Lobar Pneumonia	Report Source	Product	Role	Manufacturer	Route
		Lung Disorder Medication Error Renal Disorder Respiratory Arrest	Consumer	Phenergan (Promethazine Hydrochloride, Unspec)	PS		
				Prozac (Fluoxetine Hydrochloride, )	SS		
				Ultram (Tramadol Hydrochloride, )	SS		
				Wellbutrin (Amfebutamone Hydrochloride, )	SS		

Date:05/08/03ISR Number: 4110115-3Report Type:Expedited (15-DaCompany Report #2003UW05734  
Age:73 YR Gender:Female I/FU:I

Outcome Dose Required 25 MG DAILY Intervention to PO Prevent Permanent 300 MG Impairment/Damage 0.25 MG PRN 150 MG BID	Duration	PT Retinal Haemorrhage	Report Source Health Professional	Product	Role	Manufacturer	Route
				Seroquel	PS		ORAL
				Eskalith	SS		
				Xanax	SS		
				Wellbutrin	SS		

Date:05/08/03ISR Number: 4110196-7Report Type:Expedited (15-DaCompany Report #200310607BCC  
Age:26 YR Gender:Female I/FU:F

Outcome Dose Other 440 MG, ONCE, ORAL	Duration	PT Arthralgia Difficulty In Walking Pain In Extremity Pruritus Generalised	Report Source Consumer Health Professional Other	Product	Role	Manufacturer	Route
				Aleve (Naproxen Sodium)	PS		ORAL
				Wellbutrin			

Tendonitis

(Amfebutamone  
Hydrochloride)

SS

Date:05/09/03ISR Number: 4106574-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0298813A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged dosage text		Diplopia		Zyban	PS	Glaxosmithkline	ORAL
		Ophthalmoplegia					
		Visual Disturbance					

Date:05/12/03ISR Number: 4110853-2Report Type:Direct Company Report #CTU 192718  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Alopecia		Wellbutrin Sr 150mg Glaxo Smith Kline	PS	Glaxo Smith Kline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/12/03ISR Number: 4111111-2Report Type:Direct  
Age: Gender: I/FU:I

Company Report #USP 51755

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Welbutrin 75mg	PS	Glaxo Wellcome	
75 MG				Welbutrin Sr 150	SS	Glaxowellcome	

Date:05/12/03ISR Number: 4111153-7Report Type:Direct  
Age: Gender: I/FU:I

Company Report #USP 51611

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Zyban	PS	Glaxo Wellcome	
				Buspar	SS	Bristol-Myers Squibb	

Date:05/13/03ISR Number: 4108378-3Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0408081A  
Age: Gender:Female I/FU:F

Company Report #CA-GLAXOSMITHKLINE-A0408081A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Creatinine Increased		Bupropion	PS	Glaxosmithkline	ORAL

Date:05/13/03ISR Number: 4108380-1Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0291940A  
Age:56 YR Gender:Male I/FU:F

Company Report #NL-GLAXOSMITHKLINE-B0291940A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cerebral Haemorrhage Nicotine Dependence	Consumer	Bupropion Hydrochloride	PS	Glaxosmithkline	
UNKNOWN	150MG	Unknown 2 MON Restlessness					

Date:05/13/03ISR Number: 4111365-2Report Type:Expedited (15-DaCompany Report #8002289  
Age:36 YR Gender: I/FU:I

Company Report #8002289

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide Intentional Misuse	Literature Health	Hydrocodone W/Acetaminophen	PS		ORAL
PO				Professional	Bupropion (Long-Acting)	SS		ORAL
PO					Carisoprodol	SS		ORAL

Date:05/13/03ISR Number: 4111367-6Report Type:Expedited (15-DaCompany Report #8002291  
Age:44 YR Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Cardio-Respiratory Arrest Overdose	Literature Health	Hydrocodone W/Acetaminophen	PS		ORAL
PO				Professional	Fluoxetine	SS		ORAL
PO					Bupropion	SS		ORAL

Date:05/13/03ISR Number: 4111793-5Report Type:Expedited (15-DaCompany Report #HQWYE446302APR03  
Age:68 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Other	Abnormal Behaviour Accidental Overdose Choreoathetosis Confusional State

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Interaction	Report Source	Product	Role	Manufacturer	Route
		Euphoric Mood Gait Disturbance					
		Hallucination Medication Error Mental Status Changes Movement Disorder Nausea	Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)			ORAL
SEE IMAGE					PS		
5 MG 1X PER 1 DAY, ORAL		Poisoning Self-Medication		Ditropan (Oxybutynin, )	SS		ORAL
		Systemic Lupus					
1.5 MG 1X PER 1 DAY		Erythematosis		Klonopin (Clonazepam, )	SS		
150 MCG, ORAL				Levothyroxine (Levothyroxine, )	SS		ORAL
150 MCG, ORAL; 0.1 MG 1X PER 1 DAY, ORAL				Synthroid (Levothyroxine Sodium, )	SS		ORAL
200 MG 2X PER 1 DAY	3 MON			Wellbutrin (Amfebutamone Hydrochloride, )	SS		
				Pepcid (Famotidine)	C		
				Hydrochlorothiazide	C		
				Elavil (Amitriptyline Hydrochloride)	C		
				Vioxx (Rofecoxib)	C		
				Lortab (Hydrocodone Bitartrate/Paracetam			

ol)	C
Valdecoxib	C
Flexeril	
(Cyclobenzaprine	
Hydrochloride)	C
Enalapril	C
Tylenol	
(Paracetamol)	C

Date:05/14/03ISR Number: 4109214-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0298537A  
 Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hyperhidrosis		Zyban	PS	Glaxosmithkline	ORAL
		Urinary Tract Infection					

Date:05/14/03ISR Number: 4109300-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0298799A  
 Age:55 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Alanine Aminotransferase
Initial or Prolonged	Increased
	Gamma-Glutamyltransferase
	Increased
	Lipase Increased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pancreatitis Acute  
Pancreatolithiasis

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Consumer	Zyban	PS	Glaxosmithkline	ORAL
			Acid			
			Acetylsalicylique	SS	Glaxosmithkline	ORAL
			Maalox	SS		ORAL
			Carbocisteine	SS	Glaxosmithkline	ORAL
1UNIT per day	3 DAY		Augmentin	SS	Glaxosmithkline	ORAL
2UNIT per day	5 DAY		Myolastan	C		ORAL
3UNIT per day	8 DAY					

Date:05/15/03ISR Number: 4109955-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0377262A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphasia		Wellbutrin	PS	Glaxosmithkline	ORAL
2 YR		Balance Disorder		Estrace	C		ORAL
3MG Per day		Constipation		Advil	C	Glaxosmithkline	ORAL
		Dry Mouth		Calcium/Vitamin D	C		ORAL
1000MG Per		Dysgeusia					
day		Insomnia		Fish Oil	C		ORAL
		Libido Increased		Aspirin	C	Glaxosmithkline	
		Memory Impairment					
		Nausea					
		Pharyngolaryngeal Pain					
		Therapeutic Response					
		Unexpected					
		Tinnitus					

Date:05/15/03ISR Number: 4109957-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0395424A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Other 150MG Twice  per day	Abnormal Behaviour  Anger  Bipolar Disorder Paranoia Rash	Wellbutrin	PS	Glaxosmithkline	ORAL
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Date:05/15/03ISR Number: 4109958-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0402679A  
 Age:67 YR Gender:Female I/FU:F

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 200MG Twice Initial or Prolonged per day	Blood Urea Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
75MG Twice  per day	Choreoathetosis MON Confusional State		Effexor Xr	SS		
40MG Per day	Drug Interaction Hallucination, Visual		Pepcid	C		
150MCG Per day	Medication Error Poisoning		Synthroid	C	Glaxosmithkline	
1.5MG At night	White Blood Cell Count Increased		Klonopin	C		
2TAB Three times per day			Lorcet 10/650	C		
25MG Per day			Vioxx	C		
25MG Per day			Hctz	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/15/03ISR Number: 4109978-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0299369A  
 Age:26 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Unknown	15 DAY	Eyelid Oedema	Zyban	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged Other			Hypersensitivity Local Swelling Obstructive Airways Disorder				

Date:05/15/03ISR Number: 4109979-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0299399A  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG per day	8 DAY	Facial Palsy	Zyban	PS	Glaxosmithkline	ORAL
160MG per day				Gliclazide	C		ORAL
500MG Twice per day				Metformin	C		ORAL

Date:05/15/03ISR Number: 4112818-3Report Type:Expedited (15-DaCompany Report #B0298148A  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Abdominal Pain Alanine Aminotransferase Increased Aspartate Aminotransferase Increased Cardiolipin Antibody Positive Fibrosis Hypersensitivity Red Blood Cell Sedimentation Rate	Literature Health Professional		Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS

Increased  
Subarachnoid Haemorrhage

Date:05/15/03ISR Number: 4113339-4Report Type:Periodic Company Report #327494  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tinnitus	Health Professional	Pegasys (Peg-Interferon Alfa 2a)	PS		
SUBCUTANEOUS	1 PER WEEK;						
SUBCUTANEOUS							
10 MG DAILY;				Lexapro (Escitalopram)	SS		ORAL
ORAL							
2 PER DAY;				Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
ORAL							
1 PER DAY;				Ambien (Zolpidem Tartrate)	SS		ORAL
ORAL							
1 PER; PRN				Phenergan (Promethazine Hydrochloride)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/16/03ISR Number: 4114135-4Report Type:Direct  
Age:77 YR Gender:Male I/FU:I

Company Report #CTU 193195

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Mental Status Changes		Duragesic	PS		
Initial or Prolonged			Vicodin	SS		
			Cardura	SS		
			Wellbutrin	SS		
			Neurontin	SS		
			Tequin	SS		

Date:05/20/03ISR Number: 4112078-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0408639A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Liver Function Test		Bupropion	PS	Glaxosmithkline	ORAL
1 YR						
Initial or Prolonged	Abnormal					
	Viral Infection					

Date:05/20/03ISR Number: 4112089-8Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0299690A  
Age:24 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Death		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/03ISR Number: 4113005-5Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0293848A  
Age:41 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Alcohol Poisoning		Zyban	PS	Glaxosmithkline	
UNKNOWN	150MG Twice					
	Drug Interaction					
per day						
	Loss Of Consciousness		Ethanol	SS		
UNKNOWN						
	Memory Impairment					

Date:05/21/03ISR Number: 4113006-7Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0295743A

Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
		Drug Interaction					
per day	19	DAY					
		Epilepsy		Trimipramine	SS		ORAL
25MG per day							
		Headache		Methotrimeprazine	SS		ORAL
5MG Per day							
		Loss Of Consciousness		Moclobemide	SS		ORAL
300MG Per day							
		Muscle Spasms		Alprazolam	C		ORAL
2MG per day							
		Salivary Hypersecretion					
		Tongue Biting					

Date:05/21/03ISR Number: 4115507-4Report Type:Expedited (15-DaCompany Report #B0299573A

Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Hypokalaemia	Foreign	Zyban Tablet-Zyban			
Initial or Prolonged		Muscular Weakness		(Bupropion			
		Tremor		Hydrochloride)	PS		
				Irbesartan	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/03ISR Number: 4115551-7Report Type:Expedited (15-DaCompany Report #B0299299A  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Pressure Increased Chest Pain Dilatation Ventricular	Foreign Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL	4	DAY					

Date:05/21/03ISR Number: 4115553-0Report Type:Expedited (15-DaCompany Report #B0299244A  
 Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to Prevent Permanent ORAL Impairment/Damage		Depression Malaise Suicide Attempt	Foreign	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
				Pravastatin Sodium	C		

Date:05/21/03ISR Number: 4115578-5Report Type:Expedited (15-DaCompany Report #B0289510A  
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to Prevent Permanent 150 MG/PER Impairment/Damage DAY/ORAL		Retinal Oedema Visual Field Defect	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:05/21/03ISR Number: 4115582-7Report Type:Expedited (15-DaCompany Report #B0288569A  
 Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to Prevent Permanent		Coordination Abnormal Dizziness Motor Dysfunction	Foreign Health Professional	Wellbutrin Tablet-Zyban (Bupropion			

Impairment/Damage Muscular Weakness Hydrochloride) PS ORAL  
 SEE DOSAGE  
 Pallor  
 TEXT / ORAL  
 Thyroxine Sodium C  
 Date:05/21/03ISR Number: 4115598-0Report Type:Expedited (15-DaCompany Report #A0402277A  
 Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction Headache Migraine	Health Professional Other	Wellbutrin (Bupropion Hydrochloride) (Formulation Unknown)	PS		ORAL
100 MG/ PER							
DAY/ ORAL				Modafinil (Modafinil) (Formulation Unknown)	SS		ORAL
50 MG/ PER							
DAY/ ORAL				Citalopram Hydrobromide (Citalopram Hydrobromide) (Formulation	SS		ORAL
40 MG/ PER							
DAY/ ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Thyroxine Sodium C

Date:05/21/03ISR Number: 4115785-1Report Type:Expedited (15-DaCompany Report #B0287207A  
 Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent ORAL Impairment/Damage		Arthralgia Myalgia Purpura Serum Sickness Urticaria	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:05/22/03ISR Number: 4113844-0Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0409059A  
 Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Per day 9 DAY		Drug Interaction Grand Mal Convulsion	Consumer	Bupropion Alcohol Nexium	PS SS C	Glaxosmithkline	ORAL

Date:05/22/03ISR Number: 4113849-XReport Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0295775A  
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 43 DAY		Aphasia Cerebral Arteritis Cerebral Haemorrhage Convulsion Headache Nausea Personality Change Vomiting	Consumer	Zyban	PS	Glaxosmithkline	ORAL



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aptyalism	Consumer	Zyban	PS	Glaxosmithkline	ORAL
		Electrocardiogram Qt		Norethisterone +			
		Prolonged		Ethinylloestradiol	C		ORAL
		Intentional Misuse					
		Poisoning					
		Somnolence					
		Speech Disorder					
		Tachycardia					
		Tremor					

Date:05/22/03ISR Number: 4117383-2Report Type:Direct Company Report #USP 51006  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Buspar	PS	Bristol Myers Squibb	
				Wellbutrin	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/22/03ISR Number: 4160216-9Report Type:Periodic  
Age:59 YR Gender:Male I/FU:I

Company Report #SU-2003-000485

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
20 MG QD PO		Pruritus	Consumer	Benicar	PS		ORAL
		Rash		Wellbutrin	SS		
		Skin Odour Abnormal		Prevacid	C		
				Claritin	C		
				Effexor - Slow			
				Release	C		
				Nabumetone	C		

Date:05/23/03ISR Number: 4114683-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0300222A  
Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	10 DAY			Lansoprazole	C		ORAL
30MG Per day		Crohn'S Disease		Azathioprine	C	Glaxosmithkline	ORAL
50MG Three							
times per day				Hydroxocobalamin	C	Glaxosmithkline	
INTRAMUSCULAR	3MG Monthly			Codeine	C		ORAL
30MG As							
required							

Date:05/23/03ISR Number: 4118608-XReport Type:Direct  
Age:12 YR Gender:Male I/FU:I

Company Report #USP 55845

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
TABLET		Medication Error		Voltaren	PS		
TABLET				Wellbutrin	SS		

Date:05/28/03ISR Number: 4117117-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0392586A  
Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Laboratory Test	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Interference	Professional				
per day							
.05MG Per day				Levothyroxine	C	Glaxosmithkline	

Date:05/28/03ISR Number: 4117118-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0407308A  
Age:15 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
300MG per day							
Initial or Prolonged		Overdose	Professional				
		Status Epilepticus					

Date:05/28/03ISR Number: 4117120-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0409371A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Drug Interaction		Bupropion	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Nausea		Strattera	SS		
		Vomiting					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/03ISR Number: 4117121-3Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0038838A  
 Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day 6 WK	Aggression	Other	Zyban	PS	Glaxosmithkline	ORAL
Other UNKNOWN		Anxiety		Seroxat	SS	Glaxosmithkline	
		92 DAY		Sortis	C		ORAL
		Bipolar Disorder		Diclo	C	Glaxosmithkline	
INTRAMUSCULAR		Crying					
		Depression		Dexa-Phlogont	C		
INTRAMUSCULAR							
		Diarrhoea		Perenterol	C		ORAL
		Disturbance In Attention		Mutaflor	C		ORAL
		Gastrointestinal Disorder					
		Hyperhidrosis					
		Hypomania					
		Illusion					
		Migraine					
		Nausea					
		Nervousness					
		Sleep Disorder					
		Suicidal Ideation					
		Tachycardia					
		Vertigo					
		Weight Decreased					

Date:05/29/03ISR Number: 4118197-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0409518A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Gastrointestinal		Bupropion	PS	Glaxosmithkline	ORAL
		Haemorrhage		Ibuprofen	SS	Glaxosmithkline	
		Intentional Misuse					

Date:05/29/03ISR Number: 4118198-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0409537A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Congenital Anomaly	Cleft Lip	Bupropion	PS	Glaxosmithkline
	Maternal Drugs Affecting	Neurontin	C	
	Foetus	Maxalt	C	

Date:05/29/03ISR Number: 4118204-4Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0297338A  
 Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Ageusia	Health	Zyban	PS	Glaxosmithkline	ORAL
Other dosage text	36 DAY	Aggression	Professional				
		Agitation		Oligosol(S)	C		ORAL
		Akathisia					
		Asthenia					
		Confusional State					
		Depression					
		Hyposmia					
		Insomnia					
		Mood Altered					
		Muscle Twitching					
		Somnolence					
		Suicidal Ideation					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/03ISR Number: 4119394-XReport Type:Direct  
Age:19 YR Gender:Female I/FU:I

Company Report #CTU 194258

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100 MG SR		Grand Mal Convulsion		Bupropion Sr	PS		
BID		Loss Of Consciousness					
				Sertraline	C		
				Zolpidem	C		
				Alcohol	C		

Date:05/29/03ISR Number: 4119395-1Report Type:Direct  
Age:5 YR Gender:Male I/FU:I

Company Report #CTU 194260

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100 MG SR Q		Convulsion		Bupropion Sr	PS		
AM							
				Guanfacine	C		

Date:05/29/03ISR Number: 4119454-3Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 194224

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tremor		Wellbutrin 150 Mg X			
300 MG PO QD				R	PS		ORAL

Date:05/29/03ISR Number: 4121166-7Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-12252938

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Akathisia		Abilify Tabs 15 Mg	PS	Otsuka	
		Anxiety				Pharmaceutical	

10 mg/day Headache Company, Ltd. ORAL  
from Hypomania  
18-Mar-2003 Panic Disorder

until  
07-Apr-2003, Abilify Tabs 15 Mg SS Otsuka  
10 mg/day Pharmaceutical Company, Ltd. ORAL

from  
18-Mar-2003  
until  
07-Apr-2003, Wellbutrin Sr SS  
Vitamin E C  
Lamictal C  
Trazodone Hcl C Apothecon  
Inderal La C  
Premarin C  
Vitamin C C  
Valtrex C

Date:05/30/03ISR Number: 4119376-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0397636A  
Age:57 YR Gender:Female I/FU:F

Outcome PT  
Other Abdominal Distension  
Difficulty In Walking

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Disorientation Dizziness Drug Withdrawal Syndrome	Report Source	Product	Role	Manufacturer	Route
UNKNOWN	12 YR	Dysstasia		Wellbutrin	PS	Glaxosmithkline	
		Migraine		Maxalt	C		
		Musculoskeletal Stiffness		Botox	C		
		Tinnitus		Luvox	C		
				Eskalith	C	Glaxosmithkline	
				Premarin	C		

Date:05/30/03ISR Number: 4119380-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0410077A  
Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	8 MON	Death		Bupropion	PS	Glaxosmithkline	

Date:05/30/03ISR Number: 4119390-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0300655A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 3.7G per day	1 DAY	Agitation		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged 6.25MG per	1 DAY	Balance Disorder		Xanax	C		ORAL
day	1 DAY	Chorea					
		Coordination Abnormal					
		Dysarthria					
		Hallucinations, Mixed					
		Suicide Attempt					
		Tremor					

Date:05/30/03ISR Number: 4119391-4Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0300707A  
Age:30 YR Gender:Female I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	300MG per day 16 DAY	Angioneurotic Oedema		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged		Blood Fibrinogen Decreased		Esberiven	C		ORAL
		C-Reactive Protein		Orthosiphon	C		ORAL
UNKNOWN		Increased		Lithium Oligosol	C		
		Dermographism					
		Dyspnoea					
		Generalised Oedema					
		Leukocytosis					
		Lung Consolidation					
		Macroglossia					
		Pharyngolaryngeal Pain					
		Pruritus					
		Pyrexia					
		Respiratory Disorder					

Date:05/30/03ISR Number: 4119515-9Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0399254A  
Age:31 YR Gender:Female I/FU:F

Outcome	PT
Disability	Abdominal Distension
Other	Anaphylactic Shock
	Erythema
	Hallucination, Auditory

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Date:06/02/03ISR Number: 4120339-7Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0410089A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Convulsion		Bupropion	PS	Glaxosmithkline	ORAL
2 MON							
Other		Disability					
		Fall					
		Head Injury					

Date:06/02/03ISR Number: 4120684-5Report Type:Direct Company Report #CTU 194444  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS		ORAL
300 MG BID							
ORAL							

Date:06/02/03ISR Number: 4122361-3Report Type:Expedited (15-DaCompany Report #USA-2003-0005834  
Age:46 YR Gender:Male I/FU:I

Outcome	PT
Death	Coma
	Coronary Artery Disease
	Cryptogenic Organizing
	Pneumonia
	Hepatic Haemorrhage

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
		Hepatic Necrosis Hilar Lymphadenopathy Hypertensive Cardiomegaly					
80 MG, Q8H		Pneumonia Aspiration Prescribed Overdose Pulmonary Haemorrhage	Health Professional Other	Oxycontin Tablets(Oxycodone Hydrochloride) Cr Tablet	PS		
				Hydrocodone Bitartrate (Similar To Ind 59,175)(Hydrocodone Bitartrate) Unknown	SS		
				Trazodone(Trazodone)	SS		
				Acetaminophen(Parace tamol)	SS		
				Diazepam(Diazepam)	SS		
				Wellbutrin(Amfebutam one Hydrochloride)	SS		

Date:06/02/03ISR Number: 4140932-5Report Type:Periodic Company Report #2003005856  
Age:25 YR Gender:Male I/FU:F

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
80 MG, BID, ORAL		PT Convulsion	Health Professional	Geodon (Ziprasidone)	PS		ORAL
300 MG BID			Company Representative	Bupropion Hydrochloride	SS		
				Non-Therapeutic Products	SS		
				Alprazolam	C		
				Zolpidem Tartrate	C		

Date:06/03/03ISR Number: 4122045-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0410328A  
Age:34 YR Gender:Female I/FU:I

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
		PT					

Other 150MG Per day Antepartum Haemorrhage Bupropion PS Glaxosmithkline ORAL  
 Complications Of Maternal Exposure To Therapeutic Drugs  
 Drug Exposure During Pregnancy  
 Pregnancy

Date:06/03/03ISR Number: 4122046-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0410340A  
 Age:74 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	100MG Per day 2 WK	Enterocolitis		Bupropion	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged		Haemorrhagic		Flagyl	SS	Glaxosmithkline	
Disability		Rash		Cipro	SS		
Other							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/03/03ISR Number: 4122048-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0410361A  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG per day	374 DAY	Abortion Spontaneous	Bupropion	PS	Glaxosmithkline	ORAL
			Drug Exposure During Pregnancy				

Date:06/04/03ISR Number: 4122389-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0301009A  
 Age:24 YR Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice per day	34 DAY	Sudden Death	Zyban	PS	Glaxosmithkline	ORAL

Date:06/04/03ISR Number: 4124276-3Report Type:Expedited (15-DaCompany Report #B0300463A  
 Age:47 YR Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Cardio-Respiratory Arrest Completed Suicide Toxicologic Test Abnormal	Wellbutrin (Formulation Unknown) (Buproppion Hydrochloride) Dextropropoxyphene (Formulation Unknown) (Dextropropoxyphene)	PS		ORAL
			Literature Health Professional	Alprazolam (Formulation Unknown) (Alparzolam)	SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Imipramine (Formulation Unknown) (Imipramine) Secobarbital Sodium (Formulation Unknown) (Secobarbital Sodium)	SS  SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Wellbutrin (Formulation Unknown) (Bupropion			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL				Hydrochloride)	PS		ORAL
				Nifedipine (Formulation Unknown) (Nifedipine)	SS		ORAL
ORAL				Tolcapone (Formulation Unknown) (Tolcapone)	SS		

Date:06/04/03ISR Number: 4124281-7Report Type:Expedited (15-DaCompany Report #B0300447A  
Age:47 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride) Theophylline (Formulation Unknown) (Theophylline)	PS		ORAL
ORAL				Vancomycin (Formulation Unknown) (Vancomycin)	SS		

Date:06/04/03ISR Number: 4124284-2Report Type:Expedited (15-DaCompany Report #B0300507A  
Age:29 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Ethanol (Formulation Unknown) (Alcohol) Sibutramine Hydrochloride	SS		



(Formulation  
Unknown)  
(Sibutramine SS

Date:06/04/03ISR Number: 4124285-4Report Type:Expedited (15-DaCompany Report #B0300503A  
Age:36 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Olanzapine (Formulation Unknown) (Olanzapine)	SS		
				Fluoxetine (Formulation Unknonw) (Fluoxetine)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/04/03ISR Number: 4124288-XReport Type:Expedited (15-DaCompany Report #B0300502A

Age:57 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Olanzapine (Formulation Unknown) (Olanzapine)	SS		
				Gabapentin (Formulation Unknown) (Gabapentin)	SS		

Date:06/04/03ISR Number: 4124290-8Report Type:Expedited (15-DaCompany Report #B0300506A

Age:35 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		ORAL
ORAL				Fluvoxamine (Formulation Unknown) (Fluvoxamine)	SS		
				Olanzapine (Formulation Unknown) (Olanzapine)	SS		

Date:06/04/03ISR Number: 4124294-5Report Type:Expedited (15-DaCompany Report #B0300510A

Age:41 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death	Cardio-Respiratory Arrest Completed Suicide	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS	ORAL
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Date:06/04/03ISR Number: 4124295-7Report Type:Expedited (15-DaCompany Report #B0300511A  
Age:24 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Completed Suicide	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL

Date:06/04/03ISR Number: 4124296-9Report Type:Expedited (15-DaCompany Report #B0300509A  
Age:45 YR Gender: I/FU:I

Outcome	PT	Report Source
Death	Completed Suicide	Literature Health

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Professional

Dose	Duration	Product	Role	Manufacturer	Route
ORAL		Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL

Date:06/04/03ISR Number: 4124297-0Report Type:Expedited (15-DaCompany Report #B0300504A  
Age:40 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Olanzapine (Formulation Unknown) (Olanzapine)	SS		

Date:06/04/03ISR Number: 4124298-2Report Type:Expedited (15-DaCompany Report #B0300505A  
Age:41 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Wellbutrin (Bupropion Hydorchloride)	PS		ORAL
ORAL				Imipramine (Imipramine)	SS		
				Desipramine (Desipramine)	SS		

Date:06/04/03ISR Number: 4124414-2Report Type:Expedited (15-DaCompany Report #2003022495  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 25 MG		Bipolar I Disorder Condition Aggravated	Consumer	Antivert (Meclizine) (Meclizine)	PS		ORAL
(DAILY), ORAL		Dizziness					
		Drug Ineffective		Metaxalone (Metaxalone)	SS		
		Drug Interaction		Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		
		Fall		Olanzapine (Olanzapine)	C		
		Mania		Rabeprazole Sodium (Rabeprazole Sodium)	C		
		Nausea		Trazodone (Trazodone)	C		
				Thyroid (Thyroid)	C		
				All Other Non-Therapeutic Products	C		
				Montelukast Sodium (Montelukast Sodium)	C		

Freedom Of Information (FOI) Report

Clopidogrel Sulfate (Clopidogrel Sulfate)	C
Potassium Chloride (Potassium Chloride)	C
Aporex (Paracetamol, Dextropropoxyphene Hydrochloride)	C
Allegra-D - Slow Release (Pseudoephedrine Hydrochloride, Fexofenadine)	C
Lomotil (Atropine Sulfate Diphenoxylate Hydrochloride)	C
Doxazosin (Doxazosin)	C
Valdecoxib (Valdecoxib)	C
Estradiol (Estradiol)	C
Sertraline Hydrochloride (Sertraline Hydrochloride)	C
Ginkgo Biloba (Ginkgo Biloba)	C
Lamotrigine (Lamotrigine)	C
Deloratadine (Desloratadine)	C
Ubidecarenone (Ubidecarenone)	C
Furosemide (Furosemide)	C
Estradiol (Estradiol)	C
Budesonide (Budesonide)	C
Baclofen (Baclofen)	C
Tramdol Hydrochloride (Tramadol Hydrochloride)	C
Librax	

(Chlordiazepoxide  
Hydrochloride,  
Clidinium Bromide) C  
All Other  
Therapeutic Products C

Date:06/04/03ISR Number: 4124509-3Report Type:Expedited (15-DaCompany Report #B0300450A  
Age:22 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Completed Suicide	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL  
 Hydrochloride) PS ORAL  
 Fluoxetine  
 (Formulation  
 Unknown)  
 (Fluoxetine) SS

Date:06/04/03ISR Number: 4124511-1Report Type:Expedited (15-DaCompany Report #B0300451A  
 Age:71 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL

ORAL

Date:06/04/03ISR Number: 4124512-3Report Type:Expedited (15-DaCompany Report #B0300452A  
 Age:28 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL

ORAL

Date:06/04/03ISR Number: 4124514-7Report Type:Expedited (15-DaCompany Report #B0300508A  
 Age:42 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL

ORAL

Ethanol (Formulation  
 Unknown) (Alcohol) SS



Date:06/04/03ISR Number: 4124720-1Report Type:Expedited (15-DaCompany Report #B0300294A  
Age:32 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Wellbutrin (Bupropion Hydrochloride) Amitriptyline	PS		
				(Amitriptyline) Haloperidol (Formulation Unknown) (Haloperidol)	SS		

Date:06/05/03ISR Number: 4122959-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0407071A  
Age:65 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening INTRAVENOUS	150MG Twice	Bradycardia	Health Professional	Wellbutrin	PS	Glaxosmithkline	
Other per day		Hypotension		Blood Transfusion	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

UNKNOWN 6.25MG Twice Coreg C Glaxosmithkline  
 per day

Date:06/05/03ISR Number: 4124772-9Report Type:Expedited (15-DaCompany Report #2003023103  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cleft Lip	Health	Neurontin			
		Maternal Drugs Affecting	Professional	(Gabapentin)	PS		ORAL
UNKNOWN		Foetus					
(UNKNOWN),		Pregnancy					
ORAL							

UNKNOWN	UNKNOWN			Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		
(UNKNOWN),							

UNKNOWN	UNKNOWN			Rizatriptan Benzoate (Rizatriptan Benzoate)	SS		
(UNKNOWN),							
UNKNOWN							

Date:06/05/03ISR Number: 4125359-4Report Type:Expedited (15-DaCompany Report #B0300490A  
 Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Ethanol Increased	Literature	Wellbutrin			
		Bradycardia	Health	(Formulation			
		Brain Death	Professional	Unknown) (Bupropion			

9 GRAM(S)/	Cardiac Arrest	Hydrochloride)	PS	ORAL
SINGLE DOSE/	Coma			
ORAL	Completed Suicide			
	Gastritis Haemorrhagic	Ethanol (Formulation		
	Grand Mal Convulsion	Unknown)(Alcohol)	SS	
	Hypotension			
	Intentional Misuse			
	Intestinal Ischaemia			
	Pneumonia Aspiration			
	Vomiting			

Date:06/05/03ISR Number: 4125459-9Report Type:Expedited (15-DaCompany Report #03-414-968-01  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Grand Mal Convulsion	Health Professional	Fluvoxamine Maleate Tablets 50 Mg	PS		ORAL
Intervention to 1 TAB DAILY							
Prevent Permanent PO							
Impairment/Damage 1 TAB 3 TIMES				Wellbutrin 100 Mg	SS		ORAL
A DAY PO							
1 TAB 2 TIMES				Vivactil 5 Mg	SS		ORAL
A DAY PO							
				Wellbutrin	C		
				Vivactil	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/06/03ISR Number: 4123709-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0398921A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death				Wellbutrin	PS	Glaxosmithkline	ORAL
45000MG See Hospitalization - dosage text			Intentional Misuse				
Initial or Prolonged			Somnolence				
			Tachycardia	Heroin	SS		
				Cocaine	SS		
				Valium	SS		

Date:06/09/03ISR Number: 4124549-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0411195A  
 Age: Gender:I I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1000MG Per Initial or Prolonged day				Bupropion	PS	Glaxosmithkline	ORAL
			Agitation				
			Confusional State				
			Mania				
			Panic Attack				
			Prescribed Overdose				
			Tremor				

Date:06/09/03ISR Number: 4124550-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0411263A  
 Age:18 MON Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Single Initial or Prolonged dose				Wellbutrin	PS	Glaxosmithkline	ORAL
			Accidental Exposure				

Date:06/09/03ISR Number: 4124551-2Report Type:Expedited (15-DaCompany Report #SK-GLAXOSMITHKLINE-B0118186A  
 Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	9 DAY	Areflexia					
		Feeling Abnormal					
		Hemiparesis					
		Hypoaesthesia					
		Nervousness					

Date:06/09/03ISR Number: 4124556-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0301370A  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Condition Aggravated		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	27 DAY						
Initial or Prolonged		Psoriasis		Kardegic	C		ORAL
		Rash Pustular		Praxilene	C		ORAL
				Colchimax	C		ORAL

Date:06/09/03ISR Number: 4125431-9Report Type:Direct Company Report #CTU 195299  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Bupropion Sr 150mg	PS		ORAL
150MG QD	ORAL						
		Drug Exposure Via Breast					
		Milk					
		Postictal State					
		Restlessness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/09/03ISR Number: 4126148-7Report Type:Expedited (15-DaCompany Report #B0300536A  
Age:40 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage	Duration Hallucination, Auditory Psychiatric Evaluation Abnormal	Study Literature Health Professional	Wellbutrin(Formulati on Unknown)(Bupropion Hydrochloride) Valproate Sodium Risperidone Perphenazine	PS C C C		

Date:06/10/03ISR Number: 4127715-7Report Type:Expedited (15-DaCompany Report #S03-USA-02425-01  
Age:52 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 10 MG QAM PO	Duration Coma Grand Mal Convulsion	Health Professional	Lexapro (Escitalopram)	PS		ORAL
150 MG BID	Injury Libido Decreased	Company Representative	Wellbutrin - Slow Release (Bupropion Hydrochloride)	SS		
100 MG QD			Wellbutrin (Bupropion Hydrochloride)	SS		

Date:06/11/03ISR Number: 4126384-XReport Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0372234B  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly	Duration Drug Exposure During Pregnancy Finger Deformity		Bupropion	PS	Glaxosmithkline	

Date:06/11/03ISR Number: 4126392-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0411383A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG per day	Abortion Spontaneous		Bupropion	PS	Glaxosmithkline	ORAL
		Drug Exposure During Pregnancy					
		Maternal Drugs Affecting Foetus					

Date:06/11/03ISR Number: 4126396-6Report Type:Expedited (15-DaCompany Report #SG-GLAXOSMITHKLINE-B0294984A  
Age:41 YR Gender:Male I/FU:F

Outcome	PT
Death	Abdominal Distension
Hospitalization - Initial or Prolonged	Accident
	Activated Partial Thromboplastin Time Prolonged
	Alanine Aminotransferase Increased
	Apoptosis
	Ascites
	Aspartate Aminotransferase

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Increased Blood Alkaline Phosphatase Increased Blood Bilirubin Increased				
UNKNOWN	150MG per day 10 DAY	Blood Creatinine Increased	Bupropion Hydrochloride	PS	Glaxosmithkline	
UNKNOWN	15MG per day 5 YR	Cholestasis	Carbimazole	SS		
UNKNOWN	10MG per day 5 YR	Coagulopathy	Propranolol	C		
UNKNOWN		Condition Aggravated 2 DAY	Paracetamol	C	Glaxosmithkline	
		Dialysis	Alcohol	C		ORAL
		Disseminated Intravascular Coagulation Dyspepsia Encephalopathy Epigastric Discomfort Granuloma Haemoglobin Decreased Hepatic Failure Hepatic Trauma Hepatorenal Syndrome Hypertrophy Hypokalaemia Hyponatraemia Hypotension Insomnia Intra-Abdominal Haemorrhage Jaundice Lethargy Metabolic Acidosis Mouth Haemorrhage Nausea Necrosis Prothrombin Time Prolonged Pulmonary Haemorrhage Pulmonary Oedema Rash Rectal Haemorrhage Refusal Of Treatment By Patient Renal Impairment Sepsis				



Urine Output Decreased

Date:06/11/03ISR Number: 4126399-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0299299A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	4 DAY	Dilatation Ventricular Hypertension					

Date:06/11/03ISR Number: 4126464-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0403696A  
Age:62 YR Gender:Female I/FU:F

Outcome	PT
Other	Deafness Neurosensory Dysphonia Speech Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Consumer	Wellbutrin Sr	PS	Glaxosmithkline	
			Paxil	SS	Glaxosmithkline	ORAL
			Remeron	SS		
12.5MG Per			Lopressor	C		ORAL
day						
.5MG As			Lorazepam	C		ORAL
required						

Date:06/12/03ISR Number: 4127889-8Report Type:Expedited (15-DaCompany Report #FLUV00203001421  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Health	Luvox (Fluvoxamine			
		Grand Mal Convulsion	Professional	Maleate)	PS		ORAL
50 MG DAILY							
PO				Wellbutrin			
				(Amfebutamone			
				Hydrochloride)	SS		ORAL
100 MG TID PO							
				Vivactil			
				(Protriptyline			
				Hydrochloride)	SS		ORAL
5 MG BID PO							

Date:06/12/03ISR Number: 4127898-9Report Type:Expedited (15-DaCompany Report #B0300447A US  
Age:47 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardio-Respiratory Arrest	Literature	Bupropion			
		Drug Level Above		Hydrochloride	PS		

ORAL	Therapeutic	Theophylline	SS	ORAL
		Vancomycin	SS	

Date:06/12/03ISR Number: 4129143-7Report Type:Expedited (15-DaCompany Report #B0300536A  
 Age:40 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - Initial or Prolonged Required	Hallucination, Auditory	Study Literature Health Professional	Bupropion Hydrochloride Valproate Sodium Risperidone Perphenazine	PS C C C		
Intervention to Prevent Permanent Impairment/Damage						

Date:06/13/03ISR Number: 4127904-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0412053A  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death 100MG Twice Other per day	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
2 DAY			Xanax Klonopin Zoloft	C C C		ORAL ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/13/03ISR Number: 4127905-3Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0295743A

Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Consumer	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
		Drug Interaction					
per day	19	DAY					
		Epilepsy		Trimipramine	SS		ORAL
25MG per day							
		Headache		Methotrimeprazine	SS		ORAL
5MG Per day							
		Loss Of Consciousness		Moclobemide	SS		ORAL
150MG Twice							
		Muscle Spasms					
per day							
		Salivary Hypersecretion		Alprazolam	C		ORAL
2MG per day							
		Tongue Biting					

Date:06/13/03ISR Number: 4127915-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0301595A

Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	10	DAY					
		Complications Of Maternal		Noroxine	SS	Glaxosmithkline	ORAL
11	DAY						
		Exposure To Therapeutic					
		Drugs					
		Maternal Drugs Affecting					
		Foetus					
		Pregnancy					

Date:06/13/03ISR Number: 4127918-1Report Type:Expedited (15-DaCompany Report #NZ-GLAXOSMITHKLINE-B0301669A

Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day							

Complications Of Maternal  
Exposure To Therapeutic  
Drugs  
Maternal Drugs Affecting  
Foetus  
Pregnancy

Date:06/13/03ISR Number: 4127920-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0301731A

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Life-Threatening	Abnormal Behaviour	Health	Zyban	PS	Glaxosmithkline	ORAL
2TABS per day 4 WK						
Hospitalization -	Amnesia	Professional	Alcohol	SS		
Initial or Prolonged	Head Banging		Temazepam	SS		
	Overdose					
	Suicide Attempt					

Date:06/13/03ISR Number: 4127923-5Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0038838A

Age:36 YR Gender:Male I/FU:F

Outcome	PT
Disability	Aggression
Other	Anxiety
	Bipolar Disorder
	Crying
	Depression
	Diarrhoea
	Disturbance In Attention

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	6 WK	Emotional Disorder Fatigue Gastrointestinal Disorder	Report Source				
UNKNOWN		Hyperhidrosis	Other	Zyban	PS	Glaxosmithkline	ORAL
		Hypomania		Seroxat	SS	Glaxosmithkline	
	92 DAY	Illusion		Sortis	C		ORAL
INTRAMUSCULAR		Migraine		Diclo	C	Glaxosmithkline	
INTRAMUSCULAR		Nausea		Dexa-Phlogont	C		
		Nervousness		Perenterol	C		ORAL
		Pain In Extremity		Mutaflor	C		ORAL
		Palpitations					
		Panic Reaction					
		Restlessness					
		Retching					
		Sleep Disorder					
		Suicidal Ideation					
		Tachycardia					
		Tongue Disorder					
		Vertigo					
		Weight Decreased					

Date:06/13/03ISR Number: 4138044-XReport Type:Periodic Company Report #A0364417A  
 Age:23 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Hypersensitivity	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
Intervention to Prevent Permanent 150 MG/TWICE Impairment/Damage PER DAY/ ORAL				Ethanol	C		

Date:06/13/03ISR Number: 4138045-1Report Type:Periodic Company Report #A0173731A  
 Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage PER DAY/ ORAL		Hypersensitivity	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:06/13/03ISR Number: 4138046-3Report Type:Periodic Company Report #A0402681A  
 Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged /PER DAY/ ORAL		Myocardial Infarction	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:06/13/03ISR Number: 4138047-5Report Type:Periodic Company Report #A0375316A  
 Age:19 YR Gender:Male I/FU:I

Outcome  
 Hospitalization -  
 Initial or Prolonged  
 Required  
 Intervention to

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Prevent Permanent Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG/TWICE		Hypersensitivity	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
PER DAY/	ORAL						

Date:06/13/03ISR Number: 4138048-7Report Type:Periodic Company Report #A0371219A  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Pain Pruritus Rash Swelling	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE							
PER DAY/	ORAL						

Date:06/13/03ISR Number: 4138049-9Report Type:Periodic Company Report #A0370912A  
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Chest Discomfort Chest Pain	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE							
PER DAY/	ORAL						

Date:06/13/03ISR Number: 4138050-5Report Type:Periodic Company Report #A0370366A  
 Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Anaphylactic Reaction	Health	Zyban Tablet-Zyban			



Initial or Prolonged Professional (Bupropion Hydrochloride) PS ORAL  
150 MG/TWICE  
PER DAY/ ORAL

Date:06/13/03ISR Number: 4138051-7Report Type:Periodic Company Report #A0369803A  
Age:24 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Cardiac Enzymes Increased Chest Pain	Consumer	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

ORAL

Date:06/13/03ISR Number: 4138052-9Report Type:Periodic Company Report #A0367670A  
Age:33 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required Intervention to Prevent Permanent 150 MG/ ORAL WK Impairment/Damage	Serum Sickness	Health Professional Company Representative	Zyban Tablet-Zyban (Bupropion Hydrochloride) Ibuprofen	PS C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/13/03ISR Number: 4138053-0Report Type:Periodic  
Age:23 YR Gender:Male I/FU:F

Company Report #A0364417A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage PER DAY/ ORAL		Hypersensitivity	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
				Ethanol	C		

Date:06/13/03ISR Number: 4138054-2Report Type:Periodic  
Age:36 YR Gender:Male I/FU:F

Company Report #A0173731A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage PER DAY/ ORAL		Hypersensitivity	Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:06/16/03ISR Number: 4128942-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0412212A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100MG Per day 1 MON		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
50MG See dosage text		Drug Abuser		Ultram	SS		ORAL
		Overdose					

Date:06/16/03ISR Number: 4128948-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0300655A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 3.7G per day	1 DAY	Agitation	Consumer	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged 6.25MG per	1 DAY	Balance Disorder		Xanax	C		ORAL
		Chorea					
		Coordination Abnormal					
		Dysarthria					
		Hallucinations, Mixed					
		Overdose					
		Suicide Attempt					
		Tremor					

Date:06/16/03ISR Number: 4128951-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0301940A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day	6 DAY	Brief Psychotic Disorder		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		With Marked Stressors					

Date:06/16/03ISR Number: 4128952-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0301984A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Per day	6 WK	Cerebral Haemorrhage		Zyban	PS	Glaxosmithkline	ORAL
Life-Threatening		Coma					
		Ruptured Cerebral					
		Aneurysm					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/16/03ISR Number: 4128953-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0302002A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day	22 DAY	Initial or Prolonged	Health Professional	Zyban	PS	Glaxosmithkline	ORAL
		Dermatitis Exfoliative Epidermal Necrosis Inflammation Leukocytosis Pyrexia Rash Papular Rash Psoriaform Rash Pustular Toxic Skin Eruption					

Date:06/16/03ISR Number: 4130322-3Report Type:Expedited (15-DaCompany Report #200318842BWH  
Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Disability		Initial or Prolonged	Health Professional	Cipro (Ciprofloxacin Hydrochloride)	PS		
		Diarrhoea Enterocolitis Haemorrhagic Rash	Other	Wellbutrin Sr (Amfebutamone Hydrochloride)	SS		ORAL
100 MG, TOTAL							
DAILY, ORAL				Metronidazole	SS		

Date:06/17/03ISR Number: 4129951-2Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0296637A  
Age:19 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Unknown		Initial or Prolonged	Consumer	Zyban	PS	Glaxosmithkline	ORAL
		Aptyalism Electrocardiogram Qt Prolonged Intentional Misuse Poisoning Somnolence		Norethisterone + Ethinyloestradiol	C		ORAL

Speech Disorder  
Tachycardia  
Tremor

Date:06/17/03ISR Number: 4130575-1Report Type:Direct  
Age:33 YR Gender:Female I/FU:I

Company Report #CTU 196033

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Stevens-Johnson Syndrome		Zyban (150 Mg)	PS		
SEE IMAGE	3 MON					
Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage						

Date:06/17/03ISR Number: 4131180-3Report Type:Expedited (15-DaCompany Report #03P-163-0221247-00  
Age:41 YR Gender: I/FU:I

Outcome	PT	Report Source
Death	Completed Suicide Multiple Drug Overdose	Literature Health

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Professional

Dose	Duration	Product	Role	Manufacturer	Route
		Imipramine (Janimine) (Imipramine Hcl) (Imipramine Hcl)	PS		
		Bupropion	SS		
		Desipramine	SS		

Date:06/17/03ISR Number: 4131528-XReport Type:Expedited (15-DaCompany Report #2003009863

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly 120 MG (BID) Other		Complications Of Maternal	Health	Geodon (Ziprasidone)	PS		
		Exposure To Therapeutic Drugs Congenital Central Nervous System Anomaly	Professional Company Representative	Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		
UNKNOWN	UNKNOWN	Congenital Oesophageal Anomaly Maternal Drugs Affecting Foetus Pregnancy					

Date:06/18/03ISR Number: 4131092-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0399473A

Age:18 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Depression	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
	MON	Intentional Misuse	Professional				
		Thermal Burn					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphasia	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Dysphemia	Professional	Zoloft	SS		
		Energy Increased					
		Feeling Jittery					
		Speech Disorder					

Date:06/18/03ISR Number: 4131208-0Report Type:Direct Company Report #CTU 196062  
Age:32 YR Gender:Female I/FU:I

Outcome	PT
Other	Activities Of Daily Living Impaired Amnesia Attention Deficit/Hyperactivity Disorder Brain Damage Drug Ineffective Dyslexia Fear

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Judgement Impaired Memory Impairment Speech Disorder	Report Source	Product	Role	Manufacturer	Route
150 MG 2 X				Wellbutrin Sr	PS		OTHER
DAY OTHER							

Date:06/19/03ISR Number: 4131494-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0402911A  
Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 200MG Unknown		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged Other		Encephalopathy Intentional Misuse Suicide Attempt Tachycardia		Antiepileptic Agent	C		

Date:06/20/03ISR Number: 4132301-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0371291A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600MG per day 2 YR		Confusional State	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
		Convulsion		Lamictal	SS	Glaxosmithkline	ORAL
		Crying		Wellbutrin	SS	Glaxosmithkline	ORAL
		Dysgeusia		Pain Medication	SS		ORAL
		Emotional Disorder		Klonopin	C		ORAL
		Gastric Disorder		Lorazepam	C		
		Head Injury		Ortho Tri-Cyclen	C		ORAL
		Insomnia					
		Mental Impairment					
		Nausea					
		Prescribed Overdose					
		Rash					
		Rash Psoriaform					
		Road Traffic Accident					
		Temporal Lobe Epilepsy					
		Vaginal Infection					



Vaginal Mycosis  
Vomiting  
Weight Decreased

Date:06/20/03ISR Number: 4132302-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0393612A  
Age:4 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 112MG Twice Initial or Prolonged per day 1 MON 1 DAY	Convulsion	Health Professional	Wellbutrin	PS	Glaxosmithkline	ORAL
			Benadryl	SS	Glaxosmithkline	
			Salbutamol	SS	Glaxosmithkline	

Date:06/20/03ISR Number: 4132304-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0402428A  
Age:52 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice Initial or Prolonged per day	Stevens-Johnson Syndrome	Health Professional	Wellbutrin	PS	Glaxosmithkline	ORAL
			Paxil	SS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/20/03ISR Number: 4132414-1Report Type:Expedited (15-DaCompany Report #US-ROCHE-339787  
Age:17 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 15 TABLETS.	Acidosis		Klonopin	PS	Roche	ORAL
Initial or Prolonged 15-20 TABLETS.	Arrhythmia		Wellbutrin Sr	SS		ORAL
	Drug Screen Positive					
120 TABLETS.	Electrocardiogram Qrs		Geodon	SS		ORAL
4 TABLETS.	Complex Prolonged		Ativan	SS		ORAL
	Electrocardiogram Qt Prolonged Hyperventilation Lethargy Multiple Drug Overdose Somnolence					

Date:06/23/03ISR Number: 4132926-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0402594A  
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death	Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL
	Drug Level Increased		Paxil	SS	Glaxosmithkline	ORAL
	Intentional Misuse		Benadryl	C	Glaxosmithkline	
			Doxylamine	C		
			Celexa	C		

Date:06/23/03ISR Number: 4132927-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0410340A  
Age:74 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 100MG Per day 27 DAY	Abdominal Pain		Bupropion	PS	Glaxosmithkline	ORAL
Hospitalization - 4 DAY	Diarrhoea		Flagyl	SS	Glaxosmithkline	ORAL

Initial or Prolonged Enterocolitis Cipro SS ORAL  
 3 DAY  
 Other Haemorrhagic  
 Giardiasis  
 Rash  
 Rectal Haemorrhage

Date:06/23/03ISR Number: 4132935-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0302412A  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Parapsoriasis		Bupropion	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	32 DAY						
20MG per day				Cipramil	C		ORAL
RESPIRATORY				Terbutaline	C		
(INHALATION)	500MCG per						
day							

Date:06/23/03ISR Number: 4134315-1Report Type:Direct Company Report #CTU 196325  
 Age:36 YR Gender:Female I/FU:I

Outcome PT  
 Hospitalization - Aggression  
 Initial or Prolonged Anger  
 Other Anxiety  
 Depression  
 Dizziness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Feeling Abnormal Mental Disorder Personality Change	Report Source	Product	Role	Manufacturer	Route
1 TABLET		Restlessness		Wellbutrin Sr 150 Mg Tablets Glaxo Wellcome	PS	Glaxowellcome	

TWICE DAILY

Date:06/24/03ISR Number: 4133949-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0302485A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Bradycardia	Health	Zyban	PS	Glaxosmithkline	ORAL
12G Single							
Hospitalization -		Coma	Professional				
dose	1 DAY						
Initial or Prolonged		Convulsion		Zyprexa	SS		ORAL
80TAB Single							
		Intentional Misuse					
dose							
		Pyrexia		Imovane	SS		ORAL
		Suicide Attempt					
		Tachycardia					

Date:06/24/03ISR Number: 4134113-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0300451A  
Age:71 YR Gender:I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Zyban	PS	Glaxosmithkline	ORAL

Date:06/24/03ISR Number: 4134114-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0300452A  
Age:28 YR Gender:I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Zyban	PS	Glaxosmithkline	ORAL

Date:06/24/03ISR Number: 4134115-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0300509A  
Age:45 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Bupropion	PS	Glaxosmithkline	ORAL

Date:06/24/03ISR Number: 4134116-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0300510A  
Age:41 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Completed Suicide		Bupropion	PS	Glaxosmithkline	ORAL

Date:06/24/03ISR Number: 4134117-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0300511A  
Age:24 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Completed Suicide		Bupropion	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/24/03ISR Number: 4134118-8Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0302490A  
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Leukopenia Neutropenia		Zyban	PS	Glaxosmithkline	

Date:06/25/03ISR Number: 4135074-9Report Type:Direct Company Report #CTU 196594  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Other Required Intervention to Prevent Permanent Impairment/Damage		Aggression Alcoholic Dependence Drug Abuser Intermittent Explosive Disorder Legal Problem Physical Assault Suicidal Ideation Verbal Abuse		Paxil 5mg Wellbutrin 150mg	PS SS		

Date:06/25/03ISR Number: 4135182-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0302713A  
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day	31 DAY	Amnesia Confusional State Head Banging Overdose Suicidal Ideation		Zyban No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:06/27/03ISR Number: 4136509-8Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0107035A  
Age:18 YR Gender:Male I/FU:F

Outcome PT

Death

Aggression  
Agitation  
Alcohol Poisoning  
Alcoholism  
Anorexia  
Anxiety  
Asthenia  
Chest Pain  
Completed Suicide  
Depersonalisation  
Depression  
Derealisation  
Dissociation  
Disturbance In Attention  
Dizziness  
Drug Abuser  
Dry Mouth  
Fear  
Feeling Abnormal  
Feeling Of Despair  
Head Injury  
Headache  
Hypochondriasis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
3 WK		Irritability Loss Of Consciousness Mania	Health	Zyban	PS	Glaxosmithkline	ORAL
37.5MG per day		Mental Disorder	Professional	Chloroquine	C		
		Mood Swings					
		Obsessive-Compulsive					
		Personality Disorder					
		Palpitations					
		Personality Change					
		Psychiatric Symptom					
		Restlessness					
		Social Avoidant Behaviour					
		Treatment Noncompliance					
		Vision Blurred					

Date:06/27/03ISR Number: 4136526-8Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0302778A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Acute Respiratory		Zyban	PS	Glaxosmithkline	
Initial or Prolonged per day	3 WK	Distress Syndrome					
		Convulsion		Methotrexate	C		
		Dermatomyositis		Corticosteroids	C		
		Renal Failure		Cyclophosphamide	C		

Date:06/27/03ISR Number: 4139048-3Report Type:Expedited (15-DaCompany Report #200301196  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Creatinine Abnormal	Consumer	Extra Strength			
500 MG BID PO		Blood Urea Abnormal	Other	Tylenol Product	PS		ORAL
100 MG TID PO		Hypokalaemia		Wellbutrin	SS		ORAL
		Renal Injury		Tylenol Arthritis			



Pain Er Caplet 650  
Mg/Caplet SS ORAL

1300 MG BID

PO

Date:06/30/03ISR Number: 4147697-1Report Type:Periodic Company Report #A211395  
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage (DAILY), ORAL	125 MG	Serotonin Syndrome	Health Professional	Zoloft (Sertraline) (Sertraline Hydrochloride)	PS		ORAL
150 MG (DAILY), ORAL				Wellbutrin (Bupropion) (Bupropion Hydrochloride)	SS		ORAL

Date:06/30/03ISR Number: 4148626-7Report Type:Periodic Company Report #2002066465  
Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	200 MG (BID),	Grand Mal Convulsion	Health Professional	Sertraline (Sertraline)	PS		ORAL

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Freedom Of Information (FOI) Report

ORAL						
300 MG (BID),				Bupropion Hydrochloride (Bupropion Hydrochloride)	SS	ORAL
ORAL						
5 MG (BID),				Olanzapine (Olanzapine)	SS	ORAL
ORAL						
(BID), ORAL				Allegra-D (Pseudoephedrine Hydrochloride, Fexofenadine)	SS	ORAL
200 MG (BID				Minocycline (Minocycline)	SS	

Date:07/01/03ISR Number: 4138565-XReport Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0414180A  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Abdominal Pain Upper		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day		Irritable Bowel Syndrome					
		Nausea Vomiting					

Date:07/01/03ISR Number: 4138566-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0296280A  
Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Unknown	23 DAY	Blood Creatine	Health	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged .5TAB Per day		Phosphokinase Increased	Professional	Soprol	C		ORAL

Other	Capillary Disorder	Microval	C	ORAL
	Dermatomyositis			
	Erythema			
	Myalgia			
	Skin Ulcer			

Date:07/01/03ISR Number: 4138873-2Report Type:Direct Company Report #CTU 196958  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Convulsion		Bupropion 100mg Tab	PS		ORAL
300MG PO QAM							
Intervention to		Fall		Oxycodone	C		
Prevent Permanent		Head Injury		Mvi	C		
Impairment/Damage		Loss Of Consciousness		Trazodone	C		
		Memory Impairment		Lorazepam	C		
		Tremor		Daypro	C		
				Nicotine Patch	C		
				Hctz	C		
				Felodipine	C		

Date:07/01/03ISR Number: 4140406-1Report Type:Expedited (15-DaCompany Report #2003165781US  
 Age:74 YR Gender:Female I/FU:I

Outcome  
 Life-Threatening  
 Hospitalization -  
 Initial or Prolonged

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Disability Required	Intervention to	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Condition Aggravated	Health Professional	Flagyl (Metronidazole) Tablet	PS		
Prevent Permanent Impairment/Damage		Diarrhoea		Wellbutrin - Slow Release (Amfebutamone Hydrochloride)	SS		ORAL
100 MG, QD,		Enterocolitis					
ORAL		Haemorrhagic Rash		Ciprofloxacin Hydrochloride (Ciprofloxacin Hydrochloride)	SS		

Date:07/02/03ISR Number: 4139571-1Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0414404A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abortion Spontaneous		Bupropion	PS	Glaxosmithkline	
Other		Complications Of Maternal Exposure To Therapeutic Drugs					
		Maternal Drugs Affecting Foetus					

Date:07/02/03ISR Number: 4139572-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0414610A  
 Age:75 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fall		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - 150MG Twice		Mental Status Changes					
Initial or Prolonged per day	2 WK	Psychotic Disorder		Mellaril	C		

Date:07/02/03ISR Number: 4139573-5Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0414773A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Bupropion	PS	Glaxosmithkline	
300MG per day		Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus		Dexedrine	C	Glaxosmithkline	

Date:07/03/03ISR Number: 4141767-XReport Type:Direct Company Report #CTU 197234  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG BID PO		Hypertension		Wellbutrin 100mg Bid	PS		ORAL

Date:07/07/03ISR Number: 4142114-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0405398A  
Age:43 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Lethargy Medication Error

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Prescribed Overdose Sensation Of Heaviness Tremor	Report Source	Product	Role	Manufacturer	Route
450MG Three times per day		Visual Disturbance	Health Professional	Wellbutrin	PS	Glaxosmithkline	ORAL
				Trazodone	C		
				Zyprexa	C		
				Zolpidem	C		
				Colace	C		
				Iron	C		
				Vicodin	C		
				Motrin	C	Glaxosmithkline	

Date:07/07/03ISR Number: 4142115-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0410077A  
Age:22 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice per day	8 MON	Death	Health Professional	Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN					Cocaine	SS		

Date:07/07/03ISR Number: 4142140-0Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0302485A  
Age:38 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged dose	12G Single dose	1 DAY	Bradycardia	Health Professional	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged dose	80TAB Single dose		Coma		Zyprexa	SS		ORAL
			Convulsion					
			Drug Ineffective Intentional Misuse		Imovane	SS		ORAL

Pyrexia  
Suicide Attempt  
Supraventricular  
Tachycardia

Date:07/07/03ISR Number: 4142145-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0303164A  
Age:56 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Anxiety		Zyban	PS	Glaxosmithkline	
UNKNOWN	150MG per day 29 DAY					
	Depression		Modecate	C		
INTRAMUSCULAR	25MG per day					
	Drug Ineffective		Kliofem	C		
UNKNOWN						
	Drug Withdrawal Syndrome		Diazepam	C		ORAL
10MG Three						
times per day						
			Procyclidine	C	Glaxosmithkline	ORAL
5MG Three						
times per day						

Date:07/07/03ISR Number: 4142149-7Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0041169A  
Age:62 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Asthenia
Initial or Prolonged	Diabetes Mellitus
	Fatigue
	Hepatitis B
	Jaundice

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Freedom Of Information (FOI) Report

Ocular Icterus

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1TAB Per day			Zyban	PS	Glaxosmithkline	ORAL

Date:07/07/03ISR Number: 4144601-7Report Type:Expedited (15-DaCompany Report #HQWYE509309JUN03  
 Age:17 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 4 TABLETS OF 0.5 MG (2MG) OVERDOSE	1 DAY	Arrhythmia Cardiotoxicity Depressed Level Of Consciousness Drug Screen Positive	Literature	Ativan (Lorazepam, Tablet)	PS		ORAL
AMOUNT, ORAL 15 TABLETS OF 0.5 MG (7.5 MG) OVERDOSE	1 DAY	Electrocardiogram Qrs Complex Prolonged Electrocardiogram Qt Corrected Interval Prolonged		Clonazepam (Clonazepam)	SS		ORAL
AMOUNT, ORAL 120 TABLETS OF 20 MG (2400 MG) OVERDOSE	1 DAY	Intentional Misuse Lethargy Somnolence		Geodon (Ziprasidone)	SS		ORAL
AMOUNT, ORAL 15-20 TABLETS	1 DAY			Wellbutrin-Slow Release (Amfebutamone Hydrochloride)	SS		ORAL



OF 150 MG,

OVERDOSE

AMOUNT, ORAL 1 DAY

Date:07/08/03ISR Number: 4142995-XReport Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0302778A

Age:32 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice	Acute Respiratory	Health	Zyban	PS	Glaxosmithkline	
Initial or Prolonged per day 3 WK	Distress Syndrome	Professional				
	Convulsion		Methotrexate	C		
	Dermatomyositis		Corticosteroids	C		
	Renal Failure		Cyclophosphamide	C		
			Pain Killers	C		

Date:07/08/03ISR Number: 4143272-3Report Type:Direct

Company Report #CTU 197451

Age:36 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 150MG 2 TIMES	Suicidal Ideation		Wellbutrin Sr 150mg	PS		ORAL

ORAL

Date:07/08/03ISR Number: 4155402-8Report Type:Direct

Company Report #CTU 198780

Age:46 YR Gender:Male I/FU:I

Outcome	PT
Other	Carpal Tunnel Syndrome Muscle Spasms Pharmaceutical Product

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Complaint  
Pharyngolaryngeal Pain

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
100 MG 3 TIMES/ ORAL			Bupropion 100mg Teva	PS	Teva	ORAL

Date:07/09/03ISR Number: 4143780-5Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0415583A  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dementia Hepatomegaly		Zyban	PS	Glaxosmithkline	ORAL

Date:07/10/03ISR Number: 4144474-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0297338A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150MG See Other dosage text	36 DAY	Ageusia Aggression Agitation Akathisia Asthenia Confusional State Depression Excitability Hyposmia Insomnia Mood Altered Muscle Twitching Somnolence Suicidal Ideation	Health Professional	Zyban Oligosol(S)	PS C	Glaxosmithkline	ORAL ORAL

Date:07/10/03ISR Number: 4146700-2Report Type:Expedited (15-DaCompany Report #2003-04-4611  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Lotrisone			
		Difficulty In Walking		(Clotrimazole/Betame			
		Drug Toxicity		thasone			
		Feeling Abnormal		Dipropionate) Cream	PS		
TRANSDERMAL	TOP-DERM						
		Hyperhidrosis		Zoloft	SS		
		Hypoaesthesia		Wellbutrin			
		Loss Of Consciousness		(Bupropion)	SS		
		Mania		Valium	C		
		Middle Insomnia					
		Nervous System Disorder					
		Pain					
		Rash					
		Tinnitus					
		Weight Increased					

Date:07/14/03ISR Number: 4146088-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0415814A  
Age:58 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Agitation
Initial or Prolonged	Anxiety

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Exposure To Therapeutic  
Drugs  
Maternal Drugs Affecting  
Foetus  
Premature Menopause

Date:07/15/03ISR Number: 4147153-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0380991A  
Age:30 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Anxiety
Disability	Asthenia
	Cold Sweat
	Crying
	Depression
	Diarrhoea
	Disability
	Disturbance In Attention
	Dizziness
	Drug Ineffective
	Energy Increased
	Faeces Discoloured

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG	Per day	Feeling Abnormal Headache Hyperhidrosis Insomnia		Zyban	PS	Glaxosmithkline	ORAL
		Mood Swings		Xanax	C		
		Muscle Spasms		Pepcid	C		
		Panic Attack		Nexium	C		
		Paranoia		Ativan	C		
		Pyrexia		Zyprexa	C		
		Tremor		Serzone	C		
		Vomiting		Gabitril	C		
		Weight Decreased					

Date:07/15/03ISR Number: 4147154-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0395212A  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	50MG Twice per day	Amnesia Convulsion Disability Fall Grand Mal Convulsion	Other	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:07/15/03ISR Number: 4147155-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0404161A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Carcinoid Syndrome Pheochromocytoma	Health Professional	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:07/15/03ISR Number: 4147161-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0290894A  
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Life-Threatening	Asthenia	Health	Zyban	PS	Glaxosmithkline	ORAL
81 DAY						
Hospitalization -	Blood Glucose	Professional	Aspirin	C	Glaxosmithkline	
UNKNOWN	1TAB per day					
Initial or Prolonged	Central Nervous System		Antibiotics	C		
UNKNOWN						
Other	Mass		Alcohol	C		
	Convulsion					
	Dizziness					
	Gliosarcoma					
	Glycosuria					
	Haematuria					
	Headache					
	Nausea					
	Somnolence					
	Ventricular Tachycardia					
	Vomiting					

Date:07/15/03ISR Number: 4147165-7Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0303269A  
Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Difficulty In Walking		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Gout					
per day							
				Allopurinol + Benzbromarone	C		ORAL
30MG per day	YR						

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

5MG per day	YR		Felodipine	C		ORAL
2.5MG per day	YR		Indapamide	C		ORAL
1MG per day	YR		Prazosin	C		ORAL
UNKNOWN			Simvastatin	C		
20MG per day	YR		Lipitor	C		ORAL

Date:07/17/03ISR Number: 4150924-8Report Type:Expedited (15-DaCompany Report #2003-04-4611  
 Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Lotrisone			
		Hyperhidrosis		(Clotrimazole/Betame			
		Loss Of Consciousness		thasone			
		Mania		Dipropionate) Cream	PS		
TOPICAL	TOP-DERM						
		Nervous System Disorder		Zoloft	SS		
		Weight Increased		Wellbutrin			
				(Bupropion)	SS		
				Valium	C		

Date:07/17/03ISR Number: 4194548-5Report Type:Periodic Company Report #HQ4699818OCT2002  
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis	Consumer	Advil (Ibuprofen,			
		Urticaria		Capsule, Soft			
				Gelatin)	PS		ORAL
ORAL							
				Wellbutrin - Slow			
				Release			
				(Amfebutamone			
				Hydrochloride, )	SS		ORAL
150 MG 2X PER							
1 DAY, ORAL							



Date:07/18/03ISR Number: 4149414-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0416863A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Antinuclear Antibody	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Positive					
per day	1	YR	Arthralgia				
			Systemic Lupus				
			Erythematosis				

Date:07/18/03ISR Number: 4149415-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0416890A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Abortion Spontaneous	Bupropion	PS	Glaxosmithkline	ORAL
150MG Per day		Complications Of Maternal					
		Exposure To Therapeutic					
		Drugs					
		Maternal Drugs Affecting					
		Foetus					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/18/03ISR Number: 4149419-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0301731A

Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Abnormal Behaviour	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Amnesia	Professional				
Hospitalization -		Confusional State		Alcohol	SS		
per day	31 DAY	Overdose		Temazepam	SS		
Initial or Prolonged		Suicide Attempt		No Concurrent Medication	C		
Other							

Date:07/18/03ISR Number: 4149420-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0302713A

Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Amnesia	Consumer	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Confusional State					
per day	31 DAY	Head Banging		No Concurrent Medication	C		
		Overdose					
		Suicidal Ideation					

Date:07/18/03ISR Number: 4149424-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0304114A

Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Follicle		Zyban	PS	Glaxosmithkline	ORAL
21 DAY		Stimulating Hormone Decreased		Microgynon	C		ORAL
		Fatigue		Asthma Medication	C		
		Menstruation Irregular					
		Thyroxine Decreased					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
50MG Twice		Convulsion					
per day		Disability		Vicodin	C		
		Fall		Allegra	C		
		Grand Mal Convulsion		Septra Ds	C	Glaxosmithkline	
		Postictal State		Sudafed	C	Glaxosmithkline	
				Nasacort	C		
				Cortisone	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Bupropion	PS	Glaxosmithkline	ORAL
Other				Alcohol	SS		
				Ms Contin	SS	Glaxosmithkline	
				Diphenhydramine	SS		
				Amitriptyline	SS		
				Pamelor	SS		
				Oxazepam	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/21/03ISR Number: 4150384-7Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0417270A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Failure		Bupropion	PS	Glaxosmithkline	ORAL
		Completed Suicide		Ethanol	SS		
		Fall		Zyprexa	SS		
		Intentional Misuse		Nortriptyline	SS		
				Xanax	SS		
				Acetaminophen	SS	Glaxosmithkline	
				Nexium	C		
				Nicorette	C	Glaxosmithkline	

Date:07/21/03ISR Number: 4150385-9Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0417271A

Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Bupropion	PS	Glaxosmithkline	
Other		Intentional Misuse		Benadryl	SS	Glaxosmithkline	
		Oedema		Ritalin	SS		
		Pulmonary Congestion		Codeine	SS		
				Acetaminophen	SS	Glaxosmithkline	

Date:07/21/03ISR Number: 4150386-0Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0417272A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Bupropion	PS	Glaxosmithkline	
Other		Intentional Misuse		Carbon Monoxide	SS		
		Poisoning		Desipramine	SS		

Date:07/21/03ISR Number: 4150403-8Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0304144A

Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Twice	Urinary Retention		Zyban	PS	Glaxosmithkline	

per day

Tamsulosin

C

Date:07/21/03ISR Number: 4152987-2Report Type:Expedited (15-DaCompany Report #200301196  
Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Creatinine Abnormal	Consumer	Extra Strength			
		Blood Urea Abnormal	Other	Tylenol Product	PS		ORAL
500 MG BID PO		Dysphagia		Wellbutrin	SS		ORAL
100 MG TID PO		Gastric Haemorrhage		Tylenol Arthritis			
		Hypokalaemia		Pain Er Caplet			
		Renal Injury		650mg/Caplet	SS		ORAL
1300 MG BID							
PO							

Date:07/22/03ISR Number: 4151338-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0404175A  
Age:51 YR Gender:Female I/FU:F

Outcome	PT
Other	Abdominal Distension
	Abdominal Pain Upper

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Dose	Duration	Bladder Cancer Flatulence Tinnitus	Report Source	Product	Role	Manufacturer	Route
150MG Twice				Lotronex	PS	Glaxosmithkline	ORAL
per day				Wellbutrin	SS	Glaxosmithkline	ORAL
				Tenormin	C		
				Klonopin	C		
				Claritin	C		
				Serzone	C		
				Estratest	C		

Date:07/22/03ISR Number: 4151341-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0409520A  
Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Jaundice		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day		Metastases To Liver Pancreatic Carcinoma		Celexa	C		

Date:07/22/03ISR Number: 4151364-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0304588A  
Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Headache		Zyban	PS	Glaxosmithkline	ORAL
1050MG per							
Initial or Prolonged		Intentional Misuse					
day	1 DAY	Suicide Attempt Tremor Vertigo					

Date:07/22/03ISR Number: 4151366-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0304615A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Discomfort		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	2 DAY	Dyspnoea		No Concurrent Medication	C		

Date:07/22/03ISR Number: 4154581-6Report Type:Expedited (15-DaCompany Report #S03-USA-02189-01  
 Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Dry Mouth Hepatitis A	Consumer	Celexa (Citalopram Hydrobromide)	PS		ORAL
20 MG QD PO		Pancreatic Carcinoma		Wellbutrin (Buprpion Hydrochloride)	SS		
200 MG QD							

Date:07/22/03ISR Number: 4154704-9Report Type:Expedited (15-DaCompany Report #2003029776  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Helicobacter Infection Mental Status Changes	Health Professional	Zoloft (Sertraline) Venlafaxine Hydrochloride	PS		
Other				(Venlafaxine Hydrochloride)	SS		

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Bupropion  
 Hydrochloride  
 (Bupropion Hydrochloride) SS  
 Clarithromycin  
 (Clarithromycin) SS  
 Atorvastatin  
 (Atorvastatin) C  
 Amoxicillin  
 (Amoxicillin) C

Date:07/23/03ISR Number: 4152091-3Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0417273A  
 Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia		Bupropion	PS	Glaxosmithkline	
		Coronary Artery Stenosis		Lorazepam	SS		
		Pulmonary Oedema		Cocaine	SS		

Date:07/23/03ISR Number: 4152093-7Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0417274A  
 Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Bupropion	PS	Glaxosmithkline	
		Polytraumatism		Flurazepam	SS		
		Road Traffic Accident		Dimenhydrinate	SS		

Date:07/23/03ISR Number: 4152094-9Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0417275A  
 Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Bupropion	PS	Glaxosmithkline	
		Drug Toxicity		Cocaine	SS		
				Hydrodiuril	C		
				Adalat	C	Glaxosmithkline	
				Catapres	C		
				Losec	C		



Date:07/23/03ISR Number: 4152095-0Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0417276A  
Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Bupropion	PS	Glaxosmithkline	
		Injury Asphyxiation		Alcohol	SS		

Date:07/23/03ISR Number: 4152096-2Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0417277A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Bupropion	PS	Glaxosmithkline	
		Injury Asphyxiation		Alcohol	SS		

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Freedom Of Information (FOI) Report

Date:07/23/03ISR Number: 4152097-4Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0417278A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest		Bupropion	PS	Glaxosmithkline	
		Drug Level Above Therapeutic					

Date:07/23/03ISR Number: 4152151-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0417468A

Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 40MG Per day		Abnormal Behaviour		Paxil	PS	Glaxosmithkline	ORAL
Initial or Prolonged 200MG Per day		Fatigue		Wellbutrin	SS	Glaxosmithkline	ORAL
Other 1200MG Per day		Mania		Lithium	SS	Glaxosmithkline	ORAL
				Zyprexa	C		

Date:07/23/03ISR Number: 4154876-6Report Type:Expedited (15-DaCompany Report #USA-2003-0008657

Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anorexia	Study Health Professional Other	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		ORAL
30 MG, TID, ORAL		Dehydration					
5 MG, PRN, ORAL		Dysarthria		Oxycodone Hydrochloride (Oxycodone Hydrochloride) Other	SS		ORAL
		Electrocardiogram Qrs Complex Prolonged					
		Electrocardiogram T Wave Peaked					

10 MG, NOCTE	Electroencephalogram Abnormal	Ambien (Zolpidem Tartrate)	SS	ORAL
PRN, ORAL	Emotional Distress			
	Insomnia	Wellbutrin (Amfebutamone Hydrochloride)	SS	ORAL
100 MG, BID, ORAL				
361 DAY		Investigational Oncology Drug ( )	SS	
		Neurontin (Gabapentin)	C	
		Tenormin (Atenolol)	C	
		Paxil (Paroxetine Hydrochloride)	C	
		Pamelor (Nortriptyline Hydrochloride)	C	

Date:07/23/03ISR Number: 4158491-XReport Type:Expedited (15-DaCompany Report #2003-07-1190  
Age:42 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Ischaemic Stroke	Foreign Health Professional Other	Viraferonpeg (Peg Interferon Alfa-2b Recombinant) Injectable Solution	PS		
SUBCUTANEOUS	SUBCUTANEOUS		Rebetol (Ribavirin)			

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1000 MG QD	Capsules	SS	ORAL
ORAL			
	Zyban (Bupropion Hcl)	SS	ORAL
ORAL			

Date:07/24/03ISR Number: 4153165-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0386261A  
 Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ageusia		Wellbutrin	PS	Glaxosmithkline	ORAL
25 DAY							
		Hyperaesthesia		Prozac	C		
20MG Per day		Weight Decreased					

Date:07/24/03ISR Number: 4153168-9Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0415390A  
 Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abasia		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	DAY						
		Arthralgia					
		Bronchospasm					
		Dysphagia					
		Dyspnoea					
		Hypersensitivity					
		Musculoskeletal Stiffness					
		Myalgia					
		Pain					
		Pharyngitis					
		Pharyngolaryngeal Pain					
		Pruritus					
		Rash Generalised					
		Rheumatoid Arthritis					
		Swelling					

Date:07/24/03ISR Number: 4153173-2Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0417279A  
Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Heart Injury Lung Injury		Bupropion	PS	Glaxosmithkline	

Date:07/24/03ISR Number: 4153187-2Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0041169A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1TAB Per day Initial or Prolonged		Asthenia  Diabetes Mellitus Fatigue Hepatic Cirrhosis Jaundice Ocular Icterus		Zyban	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/24/03ISR Number: 4153189-6Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0041333A  
 Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1TABS See Initial or Prolonged dosage text 21 DAY Disability 15MG Per day		Depression		Zyban	PS	Glaxosmithkline	ORAL
		Narcolepsy					
		Road Traffic Accident		Agopton	C		ORAL
		Sleep Attacks Suicidal Ideation					

Date:07/25/03ISR Number: 4154017-5Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0305031A  
 Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Per day 4 DAY		Diarrhoea Haemorrhagic		Bupropion	PS	Glaxosmithkline	
100MCG per day		Dyspepsia		Serevent	C	Glaxosmithkline	
		Nausea					
1200MCG per day		Palpitations		Becotide	C	Glaxosmithkline	
40MG per day				Omeprazole	C		

Date:07/28/03ISR Number: 4155157-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0417842A  
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100MG Twice per day 1 YR		Melanocytic Naevus		Wellbutrin	PS	Glaxosmithkline	ORAL
				Concurrent Medications	C		

Date:07/28/03ISR Number: 4155363-1Report Type:Direct  
Age:59 YR Gender:Female I/FU:I

Company Report #CTU 198741

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Memory Impairment		Wellbutrin Sr 150 Mg	PS		ORAL
150 MG 1/DAY							
ORAL							

Date:07/28/03ISR Number: 4155366-7Report Type:Direct  
Age:59 YR Gender:Female I/FU:I

Company Report #CTU 198743

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia		Wellbutrin 150 Mg	PS		ORAL
150 DAY ORAL							

Date:07/29/03ISR Number: 4156167-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0418387A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous Drug Exposure During Pregnancy		Bupropion	PS	Glaxosmithkline	ORAL

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Date:07/29/03ISR Number: 4156171-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0304591A  
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG per day		Convulsion		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Humerus Fracture		Atepadene	C		ORAL
				Duspatalin	C		ORAL
				Pentasa	C	Glaxosmithkline	ORAL

Date:07/29/03ISR Number: 4156180-9Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0305011A  
 Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day	30 DAY	Grand Mal Convulsion		Zyban	PS	Glaxosmithkline	
		Joint Dislocation					

Date:07/29/03ISR Number: 4157475-5Report Type:Direct Company Report #CTU 198898  
 Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ONE PO BID		Epistaxis		Wellbutrin 100mg Sr	PS		ORAL
		Hypertensive Crisis					

Date:07/29/03ISR Number: 4157491-3Report Type:Direct Company Report #CTU 198887  
 Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG QD PO		Dizziness		Zyban 150mg	PS		ORAL
4MG , INCR TO 8 PIECES PER				Nicotine 4g Gum	SS		ORAL



DAY PO

Date:07/29/03ISR Number: 4157539-6Report Type:Direct  
Age:28 YR Gender:Male I/FU:I

Company Report #CTU 198884

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Zyban 150mg	PS		ORAL
150MG, QD, PO				Nicotine 4mg Gum	SS		ORAL
4MG,							

INCREASED TO

8 PIECES/DAY

PO

Date:07/30/03ISR Number: 4157444-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0405379A  
Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Intentional Self-Injury					
		Obsessive Thoughts					
		Suicidal Ideation					

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Date:07/30/03ISR Number: 4157449-4Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0418052A  
 Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	13 DAY	Decreased		Zyban	PS	Glaxosmithkline	ORAL
		Chest Pain Dissociation Drug Toxicity Ear Congestion Erythema Hypersensitivity Hypotension Oedema Peripheral Oropharyngeal Swelling Pharyngolaryngeal Pain Pruritus					

Date:07/30/03ISR Number: 4157450-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0418605A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cerebrovascular Accident		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:07/30/03ISR Number: 4157805-4Report Type:Direct Company Report #CTU 198945  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Alopecia		Welbutrin Sr	PS		ORAL

Date:07/30/03ISR Number: 4161035-XReport Type:Expedited (15-DaCompany Report #S03-USA-03156-01  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Aggression	Celexa (Citalopram	
	Akathisia	Hydrobromide)	PS
	Depression	Wellbutrin	
	Euphoric Mood	(Bupropion	
	Psychiatric Symptom	Hydrochloride)	SS
	Sedation	Paxil (Paroxetine	
	Sexual Abuse	Hydrochloride)	SS
	Suicidal Ideation		
	Tension		

Date:07/30/03ISR Number: 4161181-0Report Type:Expedited (15-DaCompany Report #A03200300069  
Age:57 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Anorexia
Initial or Prolonged	Blood Creatinine
	Blood Potassium
	Cerebral Ischaemia
	Dehydration
	Dysarthria
	Electrocardiogram Qrs
	Complex Prolonged
	Electrocardiogram T Wave
	Peaked

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Dose	Duration	Electroencephalogram Abnormal Emotional Distress	Report Source	Product	Role	Manufacturer	Route
10 MG HS-ORAL		Family Stress Insomnia	Health Professional	Ambien-Zolpidem Tartrate-Tablet-10 Mg	PS		ORAL
30 MG				Investigational Drug Oxycontin-Oxycodone Hydrochloride	SS		ORAL
TID-ORAL							
5 MG PRN-ORAL				Oxycodone	SS		ORAL
100 MG				Wellbutrin-Bupropion Hydrochloride	SS		ORAL
BID-ORAL				Gabapentin	C		
				Atenolol	C		
				Paroxetine Hydrochloride	C		
				Nortriptyline Hydrochloride	C		

Date:07/31/03ISR Number: 4158460-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0417863A  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG Twice Initial or Prolonged per day	9 DAY	Cerebral Ischaemia Dehydration		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN		Dysarthria Electrocardiogram Qrs		Blinded Trial Medication	SS	Glaxosmithkline	
30MG Three times per day		Complex Prolonged Electrocardiogram T Wave		Oxycontin	SS		ORAL

5MG As	Peaked	Oxycodone	SS	ORAL
required	Electroencephalogram			
10MG At night	Abnormal	Ambien	SS	ORAL
600MG Twice	Emotional Distress	Neurontin	C	ORAL
per day	Family Stress			
50MG Per day	Ill-Defined Disorder	Tenormin	C	ORAL
20MG As	Insomnia	Paxil	C	Glaxosmithkline ORAL
required				
50MG At night		Pamelor	C	ORAL

Date:08/01/03ISR Number: 4160237-6Report Type:Direct Company Report #CTU 199059  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Toxicity		Wellbutrin Sr 100 Mg	PS		ORAL
ONE QAM ORAL				Risperdal 2 Mg			
				Janssen	SS	Janssen	ORAL
ONE BID ORAL				Benadryl	C		

Date:08/04/03ISR Number: 4160143-7Report Type:Periodic Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-12075131  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Drug Interaction International Normalised Ratio Decreased		Warfarin Sodium	PS	Bristol-Myers Squibb Company	
"As required"				Trazodone Hcl Tabs	SS		ORAL
				Wellbutrin	SS		ORAL

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Enoxaparin C  
 Olanzapine C  
 Clonazepam C  
 Docusate Sodium C  
 Sertraline Hcl C  
 Acetaminophen Elixir C  
 Ibuprofen C  
 Diphenhydramine C  
 Metoclopramide Hcl C

Date:08/04/03ISR Number: 4160709-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0395212A  
 Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 50MG Twice per day		Amnesia Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
		Disability Fall Grand Mal Convulsion Postictal State		Vicodin Allegra Septra Ds Sudafed Nasacort Cortisone Celexa	C C C C C C C	Glaxosmithkline Glaxosmithkline	

Date:08/04/03ISR Number: 4160712-4Report Type:Expedited (15-DaCompany Report #SG-GLAXOSMITHKLINE-B0294984A  
 Age:41 YR Gender:Male I/FU:F

Outcome	PT
Death	Abdominal Distension
Hospitalization - Initial or Prolonged	Activated Partial Thromboplastin Time Prolonged Alanine Aminotransferase Increased Ascites Aspartate Aminotransferase Increased Blood Alkaline Phosphatase Increased

Blood Bilirubin Increased  
Blood Creatinine  
Increased  
Cholestasis  
Coagulopathy  
Disseminated  
Intravascular Coagulation  
Dyspepsia  
Encephalopathy  
Epigastric Discomfort  
Granuloma  
Haemoglobin Decreased  
Hepatic Failure  
Hepatic Trauma  
Hepatorenal Syndrome  
Hypertrophy  
Hypotension  
Insomnia  
Intra-Abdominal

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Haemorrhage Jaundice Lethargy Metabolic Acidosis					
UNKNOWN	150MG per day	10 DAY Mouth Haemorrhage Nausea		Bupropion Hydrochloride	PS	Glaxosmithkline	
UNKNOWN	15MG per day	5 YR Necrosis		Carbimazole	SS		
UNKNOWN	10MG per day	5 YR Prothrombin Time		Propranolol	C		
UNKNOWN		Prolonged 2 DAY		Paracetamol	C	Glaxosmithkline	
		Pulmonary Haemorrhage Pulmonary Oedema Rash Rectal Haemorrhage Renal Impairment Sepsis Urine Output Decreased		Alcohol	C		ORAL

Date:08/04/03ISR Number: 4160718-5Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0295110A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150MG See Other dosage text	18 DAY	Abdominal Pain		Zyban	PS	Glaxosmithkline	ORAL
3TAB per day	4 DAY	Arthralgia Diarrhoea		Ercefuryl	C	Glaxosmithkline	ORAL
4 DAY		Erythema		Debridat	C		ORAL
3 DAY		Gastroenteritis		Ultra Levure	C		
		Pruritus Urticaria Vomiting					

Date:08/04/03ISR Number: 4160720-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0305473A  
Age:40 YR Gender:Male I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Sleep Apnoea Syndrome		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	85 DAY			Warfarin	C	Glaxosmithkline	ORAL

Date:08/04/03ISR Number: 4161258-XReport Type:Direct Company Report #USP 50185  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Wellbutrin	PS	Glaxo Wellcome	
CAPSULE				Zyban	SS	Glaxo Wellcome	

Date:08/04/03ISR Number: 4161477-2Report Type:Direct Company Report #USP 50459  
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Diovan	PS	Ciba Geigy	
CAPSULE				Zyban	SS	Glaxo Wellcome	
TABLET							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/05/03ISR Number: 4161651-5Report Type:Direct  
Age:25 YR Gender:Male I/FU:I

Company Report #CTU 199303

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dystonia		Wellbutrin Sr 150mg Glaxo	PS	Glaxo	ORAL
150MG DAILY							
ORAL							

Date:08/05/03ISR Number: 4165548-6Report Type:Expedited (15-DaCompany Report #HQWYE200125JUL03  
Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Coordination Abnormal Drug Interaction	Study	Effexor (Venlafaxine Hydrochloride, Tablet)	PS		ORAL
Other		Dysarthria					
ORAL							
		Feeling Drunk Mental Status Changes		Clarithromycin (Clarithromycin)	SS		ORAL
ORAL							
		Nuclear Magnetic Resonance Imaging Brain Abnormal		Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL
ORAL							
				Zoloft (Sertraline Hydrochloride)	SS		ORAL
ORAL							
				Atorvastatin (Atorvastatin)	C		
				Amoxicillin (Amoxicillin)	C		
				Atenolol (Atenolol)	C		
				Clonazepam (Clonazepam)	C		
				Aspirin (Acetylsalicylic Acid)	C		
				Omeprazole (Omeprazole)	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Coordination Abnormal Drug Interaction Dysarthria	Health Professional Other	Clarithromycin (Clarithromycin) (Clarithromycin)	PS		ORAL
PER ORAL	Lung Infiltration Mental Status Changes		Effexor (Venlafaxine Hydrochloride)	SS		ORAL
PER ORAL	Nuclear Magnetic Resonance Imaging		Zoloft (Sertraline Hydrochloride)	SS		ORAL
PER ORAL	Abnormal		Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
SEE IMAGE			Atorvastatin (Atorvastatin)	SS		
			Atenolol	C		
			Clonazepam	C		
			Acetylsalicylic Acid	C		
			Omeprazole	C		
			Fosinopril Sodium	C		
			Ranitidine Hydrochloride	C		
			Tetracycline	C		
			Bismuth			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Subsalicylate C  
Metronidazole C

Date:08/06/03ISR Number: 4162492-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0379918A  
Age:17 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Completed Suicide Grand Mal Convulsion Intentional Misuse Pupil Fixed Tachycardia		Bupropion Valproic Acid	PS SS	Glaxosmithkline	ORAL

Date:08/06/03ISR Number: 4162493-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0379979A  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Apnoea Cardio-Respiratory Arrest Completed Suicide Convulsion Cyanosis Intentional Misuse Pupil Fixed Ventricular Fibrillation		Bupropion	PS	Glaxosmithkline	ORAL

Date:08/06/03ISR Number: 4162494-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0379980A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Convulsion Intentional Misuse Liver Function Test		Bupropion Acetaminophen + Hydrocodone Carisoprodol	PS SS SS	Glaxosmithkline	ORAL ORAL
UNKNOWN		Abnormal Loss Of Consciousness Oxygen Saturation		Opiates	C		

Decreased  
Respiratory Rate  
Decreased  
Tachycardia

Date:08/06/03ISR Number: 4162495-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383124A  
Age:15 YR Gender:Male I/FU:F

Outcome	PT
Death	Aggression
Hospitalization -	Agitation
Initial or Prolonged	Blood Creatine Increased
	Brain Oedema
	Cardiac Arrest
	Coma
	Completed Suicide
	Grand Mal Convulsion
	Hallucination
	Hypotension
	Intentional Misuse

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
2 WK		Labile Blood Pressure Paralysis Pupillary Reflex Impaired Respiratory Arrest Status Epilepticus Tachycardia	Bupropion	PS	Glaxosmithkline	ORAL

Date:08/06/03ISR Number: 4162496-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383131A  
Age:42 YR Gender:Male I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Death		PT Cerebral Infarction Completed Suicide	Bupropion Methylphenidate	PS SS	Glaxosmithkline	ORAL
UNKNOWN		Cyanosis Drug Level Increased Intentional Misuse Loss Of Consciousness Nervous System Disorder Renal Failure Respiratory Failure Shock				

Date:08/06/03ISR Number: 4162500-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0411832A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Other 100MG Three times per day 3 MON		Drug Interaction Grand Mal Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
50MG Per day 26 DAY			Luvox	SS		ORAL
5MG Twice per day			Vivactil	SS		ORAL

Date:08/06/03ISR Number: 4162501-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0414552A  
Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	300MG per day	5 YR	Weight Increased	Wellbutrin	PS	Glaxosmithkline	ORAL
				Levothyroxine	C	Glaxosmithkline	
				Estradiol	C		
				Mvi	C		

Date:08/06/03ISR Number: 4162506-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0419143A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day		Blindness	Wellbutrin	PS	Glaxosmithkline	ORAL
			Vitamin A Deficiency	Effexor	C		
				Neurontin	C		

Date:08/06/03ISR Number: 4162508-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0419179A  
Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Agitation Anxiety Delusion

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Hallucination Mental Disorder				
Dose	Duration		Report Source	Product	Role	Manufacturer
				Wellbutrin	PS	Glaxosmithkline
100MG Per day	17 DAY					ORAL

Date:08/06/03ISR Number: 4162509-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0419316A  
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Caesarean Section		Bupropion	PS	Glaxosmithkline	
400MG per day						
	Cardiac Disorder					
	Cardiac Murmur					
	Cardiac Septal Defect					
	Drug Exposure During Pregnancy					
	Gastrointestinal Disorder					
	Insomnia					
	Malabsorption					
	Pregnancy					

Date:08/06/03ISR Number: 4162566-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0305578A  
Age:33 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Deep Vein Thrombosis		Zyban	PS	Glaxosmithkline	ORAL
150MG per day	18 DAY					

Date:08/06/03ISR Number: 4166233-7Report Type:Expedited (15-DaCompany Report #DSA\_23038\_2003  
Age:17 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - 2 MG ONE PO	Cardiotoxicity	Literature	Ativan	PS		ORAL
Initial or Prolonged 2400 MG ONCE	Drug Screen Positive	Health	Ziprasidone	SS		ORAL



PO	Electrocardiogram Qrs	Professional			
7.5 MG ONCE	Complex Prolonged		Klonopin	SS	ORAL
PO	Electrocardiogram Qt				
3000 MG ONCE	Corrected Interval Prolonged		Wellbutrin - Slow Release	SS	
PO	Intentional Misuse				
	Lethargy				
	Po2 Decreased				
	Pulmonary Function Test Abnormal				
	Somnolence				

Date:08/06/03ISR Number: 4166271-4Report Type:Expedited (15-DaCompany Report #USA-2003-0006512  
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Consumer	Oxycontin Tablets			
		Coma	Health	(Oxycodone			
		Drug Toxicity	Professional	Hydrochloride ) Cr			
		Hepatomegaly	Other	Tablet	PS		
		Portal Triaditis		Celexa (Citalopram			
		Postoperative Infection		Hydrobromide)	SS		
		Pulmonary Congestion		Carisoprodol			
		Pulmonary Oedema		(Carisoprodol)	SS		
				Meprobamate			

Freedom Of Information (FOI) Report

(Meprobamate) SS  
 Bupriopion  
 (Amfebutamone) SS

Date:08/07/03ISR Number: 4163287-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0419536A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300MG Per day	2 WK		Alanine Aminotransferase	Wellbutrin	PS	Glaxosmithkline	ORAL
			Increased				
			Aspartate				
			Aminotransferase				
			Increased				
			Chromaturia				
			Viral Infection				

Date:08/07/03ISR Number: 4163340-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0302485A  
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening			Atrioventricular Block	Zyban	PS	Glaxosmithkline	ORAL
12G Single							
Hospitalization -			Blood Creatine				
dose	1 DAY						
Initial or Prolonged			Phosphokinase Increased	Imovane	SS		ORAL
			Bradycardia	Zyprexa	C		ORAL
			Bundle Branch Block Right	Loxapac	C		
			Cardiogenic Shock				
			Coma				
			Diarrhoea				
			Electrocardiogram Qrs				
			Complex Prolonged				
			Hepatitis				
			Hyperthermia Malignant				
			Hypotension				
			Intentional Misuse				
			Lactic Acidosis				
			Lung Disorder				
			Mydriasis				
			Nodal Rhythm				
			Respiratory Failure				

Septic Shock  
Sinusitis  
Status Epilepticus  
Suicide Attempt  
Supraventricular  
Tachycardia  
Tachycardia

Date:08/07/03ISR Number: 4165239-1Report Type:Expedited (15-DaCompany Report #2003106276  
Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hypokalaemia Leukocytosis Myocardial Ischaemia Prinzmetal Angina	Foreign Literature Health Professional	Paracetamol/Pseudoep hedrine (Paracetamol, Pseudoephedrine)	PS		ORAL
9 TABLETS WITHIN 72 HOURS, ORAL		Respiratory Alkalosis		Bupropion (Bupropion)	SS		ORAL
ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL  
 Erythromycin  
 (Erythromycin) SS ORAL

Date:08/08/03ISR Number: 4164023-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0410328A  
 Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day	Abortion Induced		Bupropion	PS	Glaxosmithkline	ORAL
		Complications Of Maternal Exposure To Therapeutic Drugs Drug Exposure During Pregnancy Haemorrhage Maternal Drugs Affecting Foetus					

Date:08/09/03ISR Number: 4167362-4Report Type:Expedited (15-DaCompany Report #HQWYE525831JUL03  
 Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Gun Shot Wound Murder	Consumer Company Representative	Effexor (Venlafaxine Hydrochloride, Tablet)	PS		ORAL
				Ambien (Zolpidem Tartrate)	SS		
				Clonazepam (Clonazepam)	SS		
				Methylphenidate (Methylphenidate)	SS		
				Wellbutrin (Amfebutamone Hydrochloride)	SS		

Date:08/11/03ISR Number: 4203734-7Report Type:Periodic  
 Age:17 YR Gender:Male I/FU:I

Company Report #A0399789A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to Prevent Permanent ORAL Impairment/Damage		Hypersensitivity	Health Professional	Paxil (Formulation Unknown) (Paroxetine Hydrochloride)	PS		ORAL
				Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	SS		ORAL
ORAL							

Date:08/12/03ISR Number: 4165966-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0420287A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Supraventricular Extrasystoles		Wellbutrin	PS	Glaxosmithkline	ORAL
				Vioxx	C		
				Lopressor	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/12/03ISR Number: 4165980-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0306189A  
Age:64 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150MG Per day 62 DAY Hospitalization - Initial or Prolonged		Hereditary Angioedema	Zyban	PS	Glaxosmithkline	ORAL

Date:08/12/03ISR Number: 4166486-5Report Type:Direct Company Report #CTU 199790  
Age:25 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - 100 MG BID Initial or Prolonged ORAL		Abnormal Behaviour Anger	Wellbutrin 100 Mg Glaxowellcome	PS	Glaxowellcome	ORAL
Required Intervention to 150 MG SID Prevent Permanent AND ORAL Impairment/Damage		Feeling Of Despair Irritability Mania Mood Swings Self Mutilation Suicidal Ideation Suicide Attempt Thinking Abnormal	Wellbutrin 150 Mg Glaxowellcome	SS	Glaxowellcome	ORAL

Date:08/14/03ISR Number: 4167942-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0301940A  
Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Per day 6 DAY Initial or Prolonged		Brief Psychotic Disorder With Marked Stressors	Zyban	PS	Glaxosmithkline	ORAL

Date:08/14/03ISR Number: 4167951-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0306220A  
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Unknown	2 MON	Drug Withdrawal Syndrome	Zyban	PS	Glaxosmithkline	ORAL
Other			Rheumatoid Arthritis				

Date:08/14/03ISR Number: 4167953-0Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0039998A  
 Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Abnormal Dreams Angiopathy Anxiety Sleep Disorder Tinnitus	Zyban	PS	Glaxosmithkline	ORAL

Date:08/15/03ISR Number: 4168645-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0420979A  
 Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1TAB Twice Initial or Prolonged per day	2 MON		Cellulitis Oedema Pruritus Rash	Wellbutrin Lithium Diovan Armour Thyroid Glucophage	PS SS C C C	Glaxosmithkline Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/15/03ISR Number: 4168650-8Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0305867A  
 Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice				Zyban	PS	Glaxosmithkline	
Initial or Prolonged per day	21 DAY		Blood Creatine Phosphokinase Increased				
			Blood Creatine Phosphokinase Mb Increased Chest Pain				

Date:08/15/03ISR Number: 4168658-2Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0041169A  
 Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1TAB Per day	21 DAY			Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged UNKNOWN	75MG Twice		Diabetes Mellitus	Diclofenac	C		
per day	1 WK		Fatigue Hepatic Cirrhosis Jaundice Ocular Icterus				

Date:08/15/03ISR Number: 4171256-8Report Type:Direct Company Report #USP 56019  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
TABLET			Medication Error	Zovirax	PS	Glaxo Wellcome	
TABLET				Wellbutrin Sr	SS	Galxosmithkline	

Date:08/18/03ISR Number: 4171929-7Report Type:Expedited (15-DaCompany Report #2003-BP-05715RO  
 Age:64 YR Gender:Unknown I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Digoxin Elixir Usp, 0.05 Mg/Ml (Digoxin)	PS		ORAL
PO				Bupropion (Amfebutamone)	SS		ORAL
PO				Doxazosin (Doxazosin)	C		ORAL

Date:08/18/03ISR Number: 4172179-0Report Type:Expedited (15-DaCompany Report #HQ2220008MAY2002  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Human Chorionic Gonadotropin Positive Caesarean Section Complications Of Maternal Exposure To Therapeutic	Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		ORAL
SEE IMAGE		Drugs Drug Ineffective Pre-Eclampsia Pregnancy		Wellbutrin - Slow Release (Amfebutamone Hydrochloride,)	SS		ORAL
150 MG 2X PER		Pregnancy Test Urine					
1 DAY, ORAL		Positive Premature Baby					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/18/03ISR Number: 4172311-9Report Type:Expedited (15-DaCompany Report #HQWYE065613AUG03

Age:1 DY Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Apgar Score Low Caesarean Section Feeding Problem In Newborn Maternal Drugs Affecting	Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule Extended Release)	PS		
TRANSPLACENTAL	150 MG	3X PER Foetus					
1 DAY,		Neonatal Apnoeic Attack					
TRANSPLACENTA		Premature Baby					
L				Wellbutrin - Slow Release (Amfebutamone Hydrochloride)	SS		
TRANSPLACENTAL	150 MG	2X PER					
1 DAY,							
TRANSPLACENTA							
L							

Date:08/19/03ISR Number: 4170021-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0422196A

Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Back Pain Eye Irritation Eye Pain Headache Thinking Abnormal Visual Acuity Reduced		Clonazepam	C		

Date:08/19/03ISR Number: 4170022-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0422216A  
Age:83 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	100MG Per day 8 MON	Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged							

Date:08/19/03ISR Number: 4170028-8Report Type:Expedited (15-DaCompany Report #SK-GLAXOSMITHKLINE-B0118186A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day 9 DAY	Aggression		Zyban	PS	Glaxosmithkline	ORAL
Areflexia Feeling Abnormal Hemiparesis Hypoaesthesia Nervousness							

Date:08/21/03ISR Number: 4171553-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0422294A  
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	100MG Twice	Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged	per day 34 DAY	Faecaloma		Effexor Xr	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Birth Control Pills C

Date:08/21/03ISR Number: 4171559-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0422754A  
Age:10 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Twice Initial or Prolonged per day	6 YR	Abnormal Behaviour		Wellbutrin	PS	Glaxosmithkline	ORAL
50MG As required	10 YR	Aggression Insomnia Negativism Self Mutilation Suicide Attempt		Trazodone	SS		

Date:08/21/03ISR Number: 4171574-3Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0041696A  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Per day	61 DAY	Vasculitis		Zyban	PS	Glaxosmithkline	ORAL

Date:08/22/03ISR Number: 4172145-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0395212A  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 50MG Twice per day		Amnesia Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
		Disability Fall Grand Mal Convulsion Treatment Noncompliance		Vicodin Allegra Septra Ds Sudafed Nasacort Cortisone	C C C C C C	Glaxosmithkline	

Celexa C  
Oral Contraceptive C

Date:08/22/03ISR Number: 4172156-XReport Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0422610A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Intentional Self-Injury	Wellbutrin	PS	Glaxosmithkline	ORAL
30TAB Per day							
Initial or Prolonged			Laceration	Venlafaxine	SS		ORAL
10TAB Per day							
			Overdose	Estazolam	SS		ORAL
10TAB Per day			Suicide Attempt				

Date:08/22/03ISR Number: 4172162-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0422842A  
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Prescribed Overdose	Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Twice							
per day	3	WK	Urticaria				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/22/03ISR Number: 4172173-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0307137A  
 Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Disinhibition		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	30 DAY	Mania					
		Mood Swings		No Concurrent Medication	C		

Date:08/22/03ISR Number: 4173059-7Report Type:Direct Company Report #CTU 200472  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bruxism		Zoloft 50-100mg			
100MG Q.D.		Tooth Injury		Pfizer	PS	Pfizer	ORAL
ORAL							
		Trismus					
				Wellbutrin 150mg			
150MG B.I.D.				Glaxowellcome	SS	Glaxo Wellcome	ORAL
ORAL							

Date:08/25/03ISR Number: 4172883-4Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0294671A  
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Aphasia		Zyban	PS	Glaxosmithkline	ORAL
43 DAY							
Other		Cerebral Arteritis					
		Electroencephalogram					
		Abnormal					
		Grand Mal Convulsion					
		Headache					
		Mental Status Changes					

Date:08/25/03ISR Number: 4172884-6Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0295775A  
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Aphasia		Zyban	PS	Glaxosmithkline	ORAL
43 DAY							
Other		Cerebral Arteritis Cerebral Haemorrhage Computerised Tomogram Abnormal Electroencephalogram Abnormal Grand Mal Convulsion Headache Nausea Personality Change Vomiting					

Date:08/25/03ISR Number: 4172887-1Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-B0305654A  
Age:24 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Angioneurotic Oedema
Initial or Prolonged	Cough
Other	Drug Rash With Eosinophilia And Systemic Symptoms

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	21 DAY	Dysphonia Dyspnoea Electrocardiogram St Segment Elevation Eosinophils Urine Hepatitis Acute Hypersensitivity Hypoxia Liver Function Test Abnormal Muscle Necrosis Myalgia Myocarditis Myositis Obstructive Airways Disorder Pruritus Pyrexia Renal Disorder	Bupropion	PS	Glaxosmithkline	ORAL

Date:08/25/03ISR Number: 4173751-4Report Type:Direct  
Age:36 YR Gender:Female I/FU:I

Company Report #CTU 200588

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 20 DAILY ORAL		Asthenia Depression		Lexapro 20 Mg Threw Away	PS		ORAL
300 DAILY ORAL		Dizziness Dyspepsia Fatigue		Wellbutrin 100 Glaxosmithkline	SS	Glaxosmithkline	ORAL
		Hypoaesthesia Insomnia Malaise Paraesthesia Somnolence Weight Increased White Blood Cell Count Increased		Nasonex Motrin	C C		



Date:08/25/03ISR Number: 4174323-8Report Type:Direct  
Age:41 YR Gender:Female I/FU:I

Company Report #CTU 200622

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Wellbutrin 150 Mg	PS		ORAL
150 MG PO QHS		Nausea					

Date:08/25/03ISR Number: 4174358-5Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 200649

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash Maculo-Papular		Wellbutrin Sr 200	PS		ORAL
PO QD	7 DAY						
				Pravachol	C		
				Tenoratic	C		
				Synthroid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/25/03ISR Number: 4177879-4Report Type:Expedited (15-DaCompany Report #2003023103  
Age:1 DY Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Caesarean Section Cleft Lip	Health Professional	Neurontin (Gabapentin)	PS		ORAL
600 MG (300, BID), ORAL		Maternal Drugs Affecting Foetus		Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		ORAL
300 MG (150, BID), ORAL				Rizatriptan Benzoate (Rizatriptan Benzoate)	SS		

Date:08/25/03ISR Number: 4177883-6Report Type:Expedited (15-DaCompany Report #2003034365  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Caesarean Section Complications Of Maternal Exposure To Therapeutic Drugs	Health Professional	Neurontin (Gabapentin)	PS		ORAL
600 MG (300, BID), ORAL		Pregnancy Test Positive Uterine Dilation And Curettage		Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		ORAL
300 MG (150, BID), ORAL				Rizatriptan Benzoate (Rizatriptan Benzoate)	SS		

Date:08/26/03ISR Number: 4173455-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0295862A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged dosage text	34	DAY	Oedema Peripheral Psoriasis	Zyban	PS	Glaxosmithkline	ORAL

Elisor	C
Questran	C
Puva	C
Daivonex	C
Diprosone	C

Date:08/26/03ISR Number: 4173460-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0307344A  
Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Dysphemia Suicidal Ideation	Zyban Alcohol	PS C	Glaxosmithkline	ORAL

Date:08/26/03ISR Number: 4178818-2Report Type:Expedited (15-DaCompany Report #A0422244A  
Age: Gender:Unknown I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged Other	Aggression Agitation Confusional State Convulsion

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Complaint	Report Source	Product	Role	Manufacturer	Route
		Gastroenteritis Medication Error Nausea					
		Pharmaceutical Product Complaint Prescribed Overdose Vomiting	Literature Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		ORAL
ORAL				Bupropion Hydrochloride (Formulation Unknown) (Generic) (Bupropion Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	SS SS		

Date:08/27/03ISR Number: 4174059-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0307572A  
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day	3 WK	Dysarthria Dysgraphia Speech Disorder		Zyban Chondrosulf	PS C	Glaxosmithkline	ORAL

Date:08/28/03ISR Number: 4176001-8Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0304144A  
Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day		Urinary Retention		Zyban Tamsulosin	PS C	Glaxosmithkline	

Outcome	PT
Disability	Aggression
Other	Anxiety
	Bipolar Disorder
	Crying
	Depression
	Diarrhoea
	Disinhibition
	Disturbance In Attention
	Emotional Disorder
	Fatigue
	Gastroenteritis
	Gastrointestinal Disorder
	Gastrooesophageal Reflux
	Disease
	Hyperhidrosis
	Hypomania
	Illusion
	Irritable Bowel Syndrome
	Migraine
	Nausea
	Nervousness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	6 WK	Pain In Extremity Palpitations Panic Attack					
UNKNOWN		Retching		Zyban	PS	Glaxosmithkline	ORAL
		Sleep Disorder		Seroxat	SS	Glaxosmithkline	
		92 DAY					
		Suicidal Ideation		Sortis	C		ORAL
INTRAMUSCULAR		Tachycardia		Diclo	C	Glaxosmithkline	
INTRAMUSCULAR		Tongue Disorder		Dexa-Phlogont	C		
		Vertigo		Perenterol	C		ORAL
		Weight Decreased		Mutaflor	C		ORAL

Date:08/29/03ISR Number: 4175894-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0419536A  
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day 2 WK	Alanine Aminotransferase		Wellbutrin	PS	Glaxosmithkline	ORAL
		Increased					
		Aspartate					
		Aminotransferase					
		Increased					
		Chromaturia					
		Fatigue					
		Myalgia					
		Nausea					
		Viral Infection					

Date:08/29/03ISR Number: 4175896-1Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0420867A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	250MG Per day 1 YR	Liver Function Test		Wellbutrin	PS	Glaxosmithkline	ORAL
		Abnormal		Zocor	C		
				Altace	C		
				Pantaloc	C		

Date:08/29/03ISR Number: 4175900-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0423838A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Creatine		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Phosphokinase Increased					
per day		Drug Interaction		Cozar	SS		
UNKNOWN		Rhabdomyolysis					

Date:08/29/03ISR Number: 4176661-1Report Type:Direct Company Report #CTU 200951

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Burning Sensation		Wellbutrin 150 Mg			
Intervention to		Difficulty In Walking		Glaxosmithkline	PS	Glaxosmithkline	ORAL
SEE IMAGE							
Prevent Permanent		Pain					
Impairment/Damage		Pruritus					
		Pyrexia					
		Swelling Face					
		Urticaria					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/02/03ISR Number: 4177018-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0298243A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agoraphobia		Zyban	PS	Glaxosmithkline	ORAL
WK		Anorexia					
		Anxiety					
		Claustrophobia					
		Convulsion					
		Depression					
		Headache					
		Insomnia					
		Joint Swelling					
		Lethargy					
		Loss Of Libido					
		Mental Disorder					
		Oedema Peripheral					
		Pain In Extremity					
		Thinking Abnormal					
		Tremor					

Date:09/02/03ISR Number: 4177033-6Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0307344A

Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysphemia		Zyban	PS	Glaxosmithkline	ORAL
		Suicidal Ideation		Alcohol	C		

Date:09/02/03ISR Number: 4177034-8Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0307360A

Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Brain Stem Infarction		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	8 DAY	Wallenberg Syndrome					



Date:09/02/03ISR Number: 4177038-5Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0041169A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1TAB Per day	21 DAY	Asthenia		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged UNKNOWN	75MG Twice	Diabetes Mellitus		Diclofenac	C		
per day	1 WK	Fatigue					
		Hepatic Cirrhosis					
		Hepatic Neoplasm					
		Hepatitis B					
		Jaundice					
		Ocular Icterus					

Date:09/03/03ISR Number: 4178076-9Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0307425A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Twice		Bradycardia		Zyban	PS	Glaxosmithkline	
Hospitalization - per day		Headache					
Initial or Prolonged		Musculoskeletal Stiffness		Valsartan	C		
		Somnolence					
		Subarachnoid Haemorrhage					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/04/03ISR Number: 4179006-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0110609A  
 Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Application Site		Wellbutrin	PS	Glaxosmithkline	
Initial or Prolonged		Dermatitis		Nicoderm	SS	Glaxosmithkline	
		Body Temperature Increased					
		Erythema Multiforme					
		Hyperlipidaemia					
		Hypersensitivity					
		Leukocytosis					
		Pain					
		Prostatic Specific Antigen Increased					
		Pruritus					
		Purpura					
		Sinusitis					
		Urticaria					

Date:09/04/03ISR Number: 4179007-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0299049A  
 Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Application Site		Nicoderm Cq 21mg	PS	Glaxosmithkline	
TRANSDERMAL	21MG	Per day 7 DAY					
Initial or Prolonged		Dermatitis		Wellbutrin	SS	Glaxosmithkline	
		Body Temperature Increased					
		Erythema Multiforme					
		Hyperlipidaemia					
		Hypersensitivity					
		Leukocytosis					
		Pain					
		Prostatic Specific Antigen Increased					
		Pruritus					
		Purpura					
		Sinusitis					
		Urticaria					

Age: YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:09/04/03ISR Number: 4179862-1Report Type:Direct Company Report #CTU 201300

Age:17 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - PO 2 WK Initial or Prolonged		Dehydration Vomiting Weight Decreased		Atomoxetine Bupropion	PS SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/08/03ISR Number: 4180886-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0301940A  
 Age:54 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Per day 4 DAY		Brief Psychotic Disorder	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged 16MG Twice		With Marked Stressors	Medrol	C		ORAL
per day						
1UNIT Per day			Temazepam	C		ORAL
6UNIT per day			Dialgirex	C		ORAL

Date:09/08/03ISR Number: 4180911-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0308339A  
 Age:47 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability 150MG Per day 53 DAY		Blood Creatine	Zyban	PS	Glaxosmithkline	ORAL
738 DAY		Phosphokinase Increased	Co-Amoxiclav	C	Glaxosmithkline	ORAL
21MG per day		Muscle Rigidity	Niquitin	C	Glaxosmithkline	
		Myalgia				

Date:09/09/03ISR Number: 4181221-2Report Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-12335733  
 Age:66 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged INTRAVENOUS		Cerebral Haemorrhage Fall	Taxol	PS	Bristol-Myers Squibb Company	
Other INTRAVENOUS		Jugular Vein Thrombosis Laceration	Carboplatin	SS	Bristol-Myers Squibb Company	
AUC=6		Syncope	Coumadin	SS	Bristol-Myers Squibb Company	

1 MON Wellbutrin SS  
 Lipitor C  
 Atrovent C

Date:09/09/03ISR Number: 4181279-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0425104A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly	100MG Per day			Bupropion	PS	Glaxosmithkline	
		Valve Disorder					
		Drug Exposure During					
		Pregnancy					
		Pulmonary Valve					
		Incompetence					

Date:09/09/03ISR Number: 4188408-3Report Type:Expedited (15-DaCompany Report #S03-NOR-03725-01  
 Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Convulsion	Foreign	Cipramil (Citalopram			
Initial or Prolonged		Drug Interaction	Other	Hydrobromide)	PS		ORAL
40 MG QD PO							
		Loss Of Consciousness		Zyban (Bupropion			
		Tongue Biting		Hydrochloride)	SS		ORAL
150 MG BID PO							
				Zyban (Bupropion			
				Hydrochloride)	SS		ORAL
150 MG QD PO							
				Zopiklon (Zopiclone)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/10/03ISR Number: 4183039-3Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0422610A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	30TAB Per day	Overdose		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged	10TAB Per day	Self Mutilation		Venlafaxine	SS		ORAL
10TAB Per day		Suicide Attempt		Estazolam	SS		ORAL

Date:09/10/03ISR Number: 4183043-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0424726A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:09/10/03ISR Number: 4183045-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0424898A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:09/10/03ISR Number: 4183047-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0425148A

Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:09/10/03ISR Number: 4185410-2Report Type:Periodic Company Report #SOS-2003-039

Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Interaction	Health	Vivactil Tablets,			

5MG BID; ORAL	Grand Mal Convulsion	Professional	5mg	PS	ORAL
		Company	Luvox Tablets, 50mg	SS	ORAL
50MG QD; ORAL		Representative	Wellbutrin Tablets, 100mg	SS	ORAL
100 MG TID;					
ORAL					

Date:09/11/03ISR Number: 4183389-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0425336A  
 Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Agitation Suicide Attempt		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:09/11/03ISR Number: 4187003-XReport Type:Direct Company Report #CTU 201618  
 Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG PO QD		Drug Hypersensitivity Oedema Peripheral  Serum Sickness Urticaria		Wellbutrin 200 Mg Qd	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/12/03ISR Number: 4184643-9Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0425393A  
 Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice	Death	Pharmaceutical Product	Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - per day	Initial or Prolonged	Complaint		Paxil	C	Glaxosmithkline	
UNKNOWN	Other			Insulin	C		
INTRAVENOUS							

Date:09/12/03ISR Number: 4184651-8Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0308326A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG per day 1 DAY	Cardiac Disorder		Zyban	PS	Glaxosmithkline	

Date:09/16/03ISR Number: 4186902-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0425504A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:09/16/03ISR Number: 4186911-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0425700A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:09/16/03ISR Number: 4186913-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0425713A  
 Age: YR Gender: I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:09/16/03ISR Number: 4190329-7Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20030902464  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Toxicity	Health Professional	Risperdal (Risperidone)	PS		ORAL
Death	2 MG, 2 IN 1 DAY, ORAL						
				Wellbutrin Sr (Bupropion Hydrochloride)	SS		ORAL
	100 MG, 1 IN 1 DAY, ORAL						

Date:09/17/03ISR Number: 4188732-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0309048A  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Circulatory Collapse		Bupropion	PS	Glaxosmithkline	ORAL
Life-Threatening	150MG Twice Other per day	Convulsion					
	31 DAY	Migraine		Unspecified Inhaler	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/17/03ISR Number: 4188740-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0425927A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Visual Acuity Reduced		Wellbutrin Celexa	PS C	Glaxosmithkline	ORAL

Date:09/17/03ISR Number: 4188741-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0426117A

Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error Pharmaceutical Product Complaint		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:09/17/03ISR Number: 4188742-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0426129A

Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Per day		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:09/17/03ISR Number: 4188743-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0426142A

Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:09/19/03ISR Number: 4189275-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0309060A

Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Zyban	PS	Glaxosmithkline	ORAL
17 DAY							

Feeling Drunk  
Psoriasis  
Somnolence

No Concurrent  
Medications C

Date:09/22/03ISR Number: 4190395-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0426581A  
Age:11 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
0 DAY		Drug Ineffective		Paxil	PS	Glaxosmithkline	ORAL
150MG Per day	2 YR	Mania		Wellbutrin	SS	Glaxosmithkline	ORAL
		Suicidal Ideation		Zyrtec	C	Glaxosmithkline	
				Risperdal	C		
				Trazodone	C		
				Lamictal	C	Glaxosmithkline	ORAL

Date:09/24/03ISR Number: 4192999-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0425709A  
Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
4 YR		Drug Exposure During		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Pregnancy		Zyprexa	C		
		Eclampsia		Lithium	C	Glaxosmithkline	
		Pregnancy		Depakote	C		
		Tremor					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/24/03ISR Number: 4193001-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0426614A

Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:09/24/03ISR Number: 4193014-0Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0309003A

Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Aptyalism Electrocardiogram Qt Prolonged Intentional Misuse Poisoning Somnolence Speech Disorder Tachycardia Tremor		Zyban Synfase	PS C	Glaxosmithkline	ORAL ORAL

Date:09/24/03ISR Number: 4193019-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0309572A

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other 150MG per day	28 DAY	Depression		Zyban	PS	Glaxosmithkline	ORAL
1TAB Per day		Gingival Pain		Kliofem	C		ORAL
10MG Per day		Gingival Swelling		Propanolol	C		ORAL
		Insomnia Oedema Mouth Oral Pain					

Date:09/24/03ISR Number: 4198239-6Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20030904723

Age:43 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Multiple Drug Overdose	Literature Health Professional	Risperdal (Risperidone) Unspecified	PS		ORAL
ORAL			Distributor	Paroxetine (Paroxetine) Bupropion (Bupropion)	SS SS		

Date:09/24/03ISR Number: 4199156-8Report Type:Expedited (15-DaCompany Report #USA-2003-0009569  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiomegaly	Health Professional	Oxycontin Cr	PS		
MG		Haemorrhage Pneumonia	Professional Other	Bupropion (Amfebutamone)	SS		
MG		Toxicologic Test Abnormal		Olanzapine (Olanzapine)	SS		
MG				Venlafaxine (Venlafaxine)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/25/03ISR Number: 4194031-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0426180A  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Twice	Anxiety		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization - per day	15 DAY	Clonus					
Initial or Prolonged Disability		Dizziness					
Other		Drug Hypersensitivity					
		Encephalopathy					
		Hyperhidrosis					
		Muscle Spasms					
		Opisthotonus					
		Rash					

Date:09/25/03ISR Number: 4194032-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0426921A  
 Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:09/25/03ISR Number: 4194035-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0427051A  
 Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	
		Pharmaceutical Product Complaint					

Date:09/25/03ISR Number: 4194047-0Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0309137A  
 Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See		Arthralgia		Zyban	PS	Glaxosmithkline	ORAL

Initial or Prolonged Feeling Cold  
dosage text 20 DAY  
General Physical Health  
Deterioration  
Myalgia  
Pyrexia  
Rash Pruritic

Date:09/25/03ISR Number: 4196195-8Report Type:Direct Company Report #CTU 202578  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG TWICE		Circulatory Collapse Convulsion		Welbutrin 150 Mg Glaxosmithkline	PS	Glaxosmithklin	
A DAY							

Date:09/25/03ISR Number: 4196619-6Report Type:Direct Company Report #CTU 202567  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - BUPROPION Initial or Prolonged ORAL		Angioneurotic Oedema Urticaria Vasculitis		Bupropion	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/26/03ISR Number: 4195320-2Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0425393A

Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice	Death	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - per day	Initial or Prolonged	Dizziness	Professional				
20MG Per day	Other	Pharmaceutical Product		Paxil	C	Glaxosmithkline	ORAL
INTRA VENOUS	Complaint			Insulin	C		
				Lorazepam	C		
				Micardis	C	Glaxosmithkline	
				Cephalexin	C	Glaxosmithkline	

Date:09/26/03ISR Number: 4195323-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0427337A

Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Three	Renal Failure		Wellbutrin	PS	Glaxosmithkline	ORAL
times per day							

Date:09/26/03ISR Number: 4195326-3Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0299573A

Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN	Initial or Prolonged	Hypokalaemia		Zyban	PS	Glaxosmithkline	
300MG per day	Tremor	Muscular Weakness		Aprovel	C		

Date:09/26/03ISR Number: 4195334-2Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0309807A

Age:52 YR Gender:Male I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG per day	Bronchopneumonia		Zyban	PS	Glaxosmithkline	ORAL
		Death					

Date:09/26/03ISR Number: 4199884-4Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20030904632  
 Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Acidosis Base Excess Blood Bicarbonate Decreased Completed Suicide	Literature Health Professional Distributor	Tylox (Oxycodone/Acetaminophen/Dextromethorphan/Pseudoephedrine) Capsules	PS		ORAL
ORAL		Drug Level Increased Drug Screen Positive Hypotension Overdose Po2 Increased Ventricular Tachycardia		Acetaminophen/Codeine (Acetaminophen/Codeine) Tablets Acetaminophen/Aspirin/Caffeine (All Other Therapeutic Products) Bupropion (Bupropion) Aspirin (Acetylsalicylic Acid) Metformin (Metformin)	SS SS SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Citalopram (Citalopram)	SS
Diethylpropion (Amfepramone Hydrochloride)	SS
Oil Base Paint	SS
Oil Base Paint (All Other Non-Therapeutic Products)	SS

Date:09/29/03ISR Number: 4196777-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0426919A  
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Myocardial Infarction					
per day	1	MON		Xanax	C		
				Alcohol	C		

Date:09/29/03ISR Number: 4196782-7Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0427237A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Cholesterol Increased Convulsion Hypertension		Zyban	PS	Glaxosmithkline	ORAL

Date:09/29/03ISR Number: 4196784-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0427367A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:09/29/03ISR Number: 4196786-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0427510A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:09/29/03ISR Number: 4196789-XReport Type:Expedited (15-DaCompany Report #IE-GLAXOSMITHKLINE-B0293852A  
Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Behaviour		Zyban	PS	Glaxosmithkline	ORAL
Other		Agitation					
150MG		Confusional State		Alcohol	SS		
Variable dose	8 DAY	Unknown					
UNKNOWN	2UNIT	Drug Interaction					
		Memory Impairment					
		Paranoia					
		Verbal Abuse					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/29/03ISR Number: 4197754-9Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #CTU 202773

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Bupropion Hydrochloride	PS	Geneva Pharmceutical	ORAL
75 MG TID							
ORAL				Bupropion Hydrochloride	SS	Geneva Pharmceutical	

Date:09/29/03ISR Number: 4202063-5Report Type:Expedited (15-DaCompany Report #200304730  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Drowning	Literature Other	Unspecified Acetaminophen Product	PS		
75 G		Drug Level Above					
		Therapeutic Intentional Misuse		Bupropion Alprazolam	SS SS		

Date:09/30/03ISR Number: 4197998-6Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0427236A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Electrocardiogram Qt		Bupropion	PS	Glaxosmithkline	
Other UNKNOWN	150MG	Unknown Prolonged		Celexa	SS		ORAL
300MG Unknown		Grand Mal Convulsion		Methotrimeprazine	SS		
UNKNOWN		Intentional Misuse					

Date:09/30/03ISR Number: 4197999-8Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0427238A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Electrocardiogram Qt		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Unknown						
		Prolonged		Celexa	SS		ORAL
300MG	Unknown						
		Grand Mal Convulsion		Methotrimeprazine	SS		
UNKNOWN							

Date:09/30/03ISR Number: 4198012-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0307622A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Psoriasis		Zyban	PS	Glaxosmithkline	ORAL
		Rash		Flucloxacillin	C	Glaxosmithkline	
		Rash Erythematous					

Date:09/30/03ISR Number: 4200612-4Report Type:Direct Company Report #CTU 202818  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin Sr 200mg	PS		ORAL
200 MG	PO BID			Ortho Evra	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/30/03ISR Number: 4200613-6Report Type:Direct  
Age:55 YR Gender:Female I/FU:I

Company Report #CTU 202879

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain		Paxil 20 Mg One			
20 GM ONE		Diarrhoea		Daily	PS		
DAILY		Dizziness					
100 MG ONE		Pharmaceutical Product		Wellbutrin 100 Mgs			
DAILY AND		Complaint		Daily 1/2 Then Bid	SS		
THEN BID		Tinnitus					

Date:09/30/03ISR Number: 4202892-8Report Type:Expedited (15-DaCompany Report #SUS1-2003-00289  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Idiopathic	Health	Adderall			
Initial or Prolonged		Thrombocytopenic Purpura	Professional	Xr(Amphetamine			
60 MG,			Other	Aspartate,	PS		
1X/DAY:QD				Amphetamine Sulfate,			
150,				Dextroamphetamine			
1X/DAY:QD				Effexor-Xr(Venlafaxi	SS		
150MG 2X/DAY				ne Hydrochloride)			
				Wellbutrin - Slow			
				Release(Bupropion	SS		
				Hydrochloride)			

Date:09/30/03ISR Number: 4202923-5Report Type:Expedited (15-DaCompany Report #200304863  
Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Acidosis	Literature	Unspecified			
Hospitalization - Initial or Prolonged		Completed Suicide	Other	Apap/Dm/Pse Product			
PO		Hypotension		Unknown	PS		ORAL
Required Intervention to Prevent Permanent Impairment/Damage		Multiple Drug Overdose		Acetaminophen/Aspirin/Caffeine	SS		ORAL
PO		Ventricular Tachycardia		Bupropion (Long-Acting)	SS		ORAL
PO				Metformin	SS		
PO				Citalopram	SS		ORAL
PO				Acetaminophen/Oxycodone	SS		ORAL
PO				Acetaminophen/Codeine	SS		ORAL
PO				Diethylpropion	SS		ORAL
PO				Oil-Based Paint	SS		
PO				Aspirin	SS		ORAL

Date:09/30/03ISR Number: 4203077-1Report Type:Expedited (15-DaCompany Report #NSADSS2003011298  
Age:42 YR Gender:Female I/FU:F

Outcome	PT
Death	Anaemia
Hospitalization - Initial or Prolonged	Anger
	Chronic Obstructive Pulmonary Disease
	Coma

Freedom Of Information (FOI) Report

Dose	Duration	Confusional State Depression Disturbance In Attention	Report Source	Product	Role	Manufacturer	Route
SEE IMAGE		Drug Ineffective	Health	Risperdal			
		Female Orgasmic Disorder	Professional	(Risperidone)			
		Ganglion		Tablets	PS		ORAL
		Hallucination, Auditory		Effexor (Venlafaxine			
		Headache		Hydrochloride)	SS		
		Insomnia		Wellbutrin			
		Mania		(Bupropion			
		Memory Impairment		Hydrochloride)	SS		
		Pneumonia		Paxil (Paroxetine			
		Pulmonary Hypertension		Hydrochloride)	SS		
		Renal Failure		Remeron			
		Restlessness		(Mirtazapine)	SS		
		Sleep Apnoea Syndrome		Zoloft (Sertraline			
		Suicidal Ideation		Hydrochloride)	C		
		Tachycardia		Xanax (Alprazolam)	C		
		Thinking Abnormal		Klonopin			
		Tumour Excision		(Clonazepam)	C		
		Weight Decreased		Lithium (Lithium)	C		
	Weight Increased		Lasix (Furosemide)	C			
			Ambien (Zolpidem				
			Tartrate)	C			
			Zyprexa (Olanzapine)	C			
			Neurontin				
			(Gabapentin)	C			
			Synthroid				
			(Levothyroxine				
			Sodium)	C			
			Valium (Diazepam)	C			
			Geodon (Ziprasidone				
			Hydrochloride)	C			
			Provigil (Modafinil)	C			
			Medicine For Htn				
			(Antihypertensives)	C			
			Medicinese For				
			Diabetes (Drug Used				
			In Diabetes)	C			
			Zenical (Orlistat)	C			



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tic		Wellbutrin	PS	Glaxosmithkline	ORAL
.25MG At		Tourette'S Disorder		Risperdal	C		
night							

Date:10/02/03ISR Number: 4199974-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0426614A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	

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Freedom Of Information (FOI) Report

Date:10/02/03ISR Number: 4199976-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0427051A

Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
		Medication Error Pharmaceutical Product Complaint					

Date:10/02/03ISR Number: 4200864-0Report Type:Direct Company Report #CTU 203000

Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr 100 Mg Tablet	PS		
Required Intervention to Prevent Permanent Impairment/Damage		Rash Urticaria		Hydrochlorothiazide 25mg Tab	SS		
				Bupropion (Wellbutrin Sr)	C		
				Hydrochlorothiazide	C		
				Simvastatin	C		

Date:10/03/03ISR Number: 4200860-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0427852A

Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
		Medication Error Pharmaceutical Product Complaint					

Date:10/03/03ISR Number: 4200861-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0427881A

Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
		Medication Error Pharmaceutical Product Complaint					

Date:10/03/03ISR Number: 4200862-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0427883A

Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
		Medication Error Pharmaceutical Product Complaint					

Date:10/03/03ISR Number: 4201737-XReport Type:Direct Company Report #CTU 203103

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr 150 Mg	PS	Glaxo Smith Kline	ORAL
Other		Medication Error		Glaxo Smith Kline			
150 MG ONCE							
DAILY ORAL							
150 MG ONCE				Wellbutrin Xl 150 Mg	SS		ORAL
DAILY ORAL							

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Freedom Of Information (FOI) Report

Date:10/03/03ISR Number: 4206023-XReport Type:Expedited (15-DaCompany Report #2003039894

Age:45 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature	Doxepin (Caps)			
Other		Completed Suicide	Health	(Doxepin)	PS		ORAL
ORAL			Professional	Bupropion (Bupropion)	SS		ORAL
ORAL				Clorazepate Dipotassium (Clorazepate Dipotassium)	SS		ORAL
ORAL				All Other Therapeutic Products	SS		ORAL

Date:10/06/03ISR Number: 4201882-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0427597A

Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Complications Of Maternal		Bupropion	PS	Glaxosmithkline	
100MG Twice		Exposure To Therapeutic					
per day		Drugs					
		Intentional Misuse					
		Stillbirth					

Date:10/06/03ISR Number: 4201883-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0427597B

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Exposure During		Bupropion	PS	Glaxosmithkline	
100MG Twice		Pregnancy					
Congenital Anomaly		Fat Atrophy					
per day							

Flushing  
Head Deformity  
Muscle Atrophy  
Skin Maceration  
Umbilical Cord  
Abnormality

Date:10/06/03ISR Number: 4201885-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0427884A

Age: YR Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error		Wellbutrin	PS	Glaxosmithkline	
		Pharmaceutical Product Complaint		Wellbutrin	SS	Glaxosmithkline	

Date:10/06/03ISR Number: 4201887-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0428071A

Age:43 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	5 DAY	Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
		Pharmaceutical Product Complaint		Wellbutrin	SS	Glaxosmithkline	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/06/03ISR Number: 4201888-XReport Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0428095A  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour		Wellbutrin	PS	Glaxosmithkline	ORAL
68 DAY		Insomnia		Effexor Xr	C		ORAL
225MG per day		Mania					

Date:10/06/03ISR Number: 4201890-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0428121A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Twice		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
per day							

Date:10/06/03ISR Number: 4201892-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0428137A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abasia		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN		Rash					
		Stevens-Johnson Syndrome					

Date:10/06/03ISR Number: 4205663-1Report Type:Expedited (15-DaCompany Report #2003039883  
 Age:17 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardiac Arrest	Literature	Sertraline			
Other		Completed Suicide	Health	(Sertraline)	PS		ORAL
ORAL		Drug Level Increased	Professional	Risperidone			

ORAL	Respiratory Arrest	(Risperidone)	SS	ORAL
		Bupropion (Bupropion)	SS	ORAL

Date:10/06/03ISR Number: 4205741-7Report Type:Expedited (15-DaCompany Report #2003039667  
 Age:76 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health	Amlodipine (Amlodipine)	PS		ORAL
ORAL			Professional	Atenolol (Atenolol)	SS		ORAL
ORAL				Bupropion (Bupropion)	SS		ORAL

Date:10/07/03ISR Number: 4202507-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0428237A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Wellbutrin	SS	Glaxosmithkline	
150MG Per day	5 DAY						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/07/03ISR Number: 4202508-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0428238A

Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Medication Error Pharmaceutical Product Complaint	Wellbutrin	PS	Glaxosmithkline	

Date:10/07/03ISR Number: 4202509-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0428252A

Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 200MG Twice Initial or Prolonged per day 10 YR Disability			Hip Fracture Pain	Wellbutrin	PS	Glaxosmithkline	ORAL
				Effexor	C		
				Unknown Supplement	C		
				Guaifenesin	C		
				Oral Contraceptives	C		
				Allegra	C		
				Corticosteroids (Unspecified)	C		
				Zyrtec	C	Glaxosmithkline	

Date:10/07/03ISR Number: 4202512-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0428688A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Medication Error	Wellbutrin	PS	Glaxosmithkline	

Date:10/07/03ISR Number: 4202513-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0428692A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Medication Error	Wellbutrin	PS	Glaxosmithkline	



Date:10/07/03ISR Number: 4202515-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0428716A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:10/07/03ISR Number: 4206707-3Report Type:Expedited (15-DaCompany Report #2003178859US  
Age:28 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Calan (Verapamil) Tablet Venlafaxine (Venlafaxine) Bupropion (Amfebutamone)	PS SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/07/03ISR Number: 4206710-3Report Type:Expedited (15-DaCompany Report #2003178477US  
Age:47 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature	Xanax (Alprazolam)			
ORAL		Completed Suicide	Health	Tablet	PS		ORAL
		Intentional Misuse	Professional	Metadon Pharmacia (Methadone Hydrochloride) Tablet	SS		
				Bupropion (Amfebutamone)	SS		

Date:10/08/03ISR Number: 4206888-1Report Type:Direct Company Report #USP 080184  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error		Wellbutrin	PS	B-W	
TABLET				Wellbutrin	SS	B-W	

Date:10/08/03ISR Number: 4207435-0Report Type:Expedited (15-DaCompany Report #2003034365  
Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Caesarean Section	Consumer	Neurontin			
600 MG (300,		Complications Of Maternal	Health	(Gabapentin)	PS		ORAL
BID), ORAL		Exposure To Therapeutic	Professional				
		Drugs		Bupropion			
300 MG (150,		Maternal Drugs Affecting		Hydrochloride			
BID), ORAL		Foetus		(Bupropion			
		Pregnancy Test Positive		Hydrochloride)	SS		ORAL
				Rizatriptan Benzoate (Rizatriptan			

Date:10/09/03ISR Number: 4205033-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0428689A  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	200MG per day	Rash Generalised		Wellbutrin	PS	Glaxosmithkline	
150MG Single		Systemic Lupus		Wellbutrin	SS	Glaxosmithkline	ORAL
dose		Erythematosis					
		Urticaria		Paxil Cr	C	Glaxosmithkline	
				Valium	C		

Date:10/09/03ISR Number: 4209192-0Report Type:Expedited (15-DaCompany Report #KII-2003-0002499  
 Age:43 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Acidosis
Initial or Prolonged	Alanine Aminotransferase
Other	Aspartate
	Aminotransferase
	Blood Alkaline
	Phosphatase
	Blood Potassium
	Blood Pressure Systolic

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Increased Depressed Level Of Consciousness Epistaxis	Health Professional	Oxycodone Hydrochloride (Similar To Nda 20-553) (Oxycodone Hydrochloride)	PS		ORAL
ORAL		Haematocrit Haemoglobin Heart Rate Increased Intentional Misuse Multiple Drug Overdose					
		Pharyngeal Haemorrhage Prothrombin Time		Coumadin(Warfarin Sodium)	SS		ORAL
ORAL		Respiratory Rate Decreased		Wellbutrin(Amfebutam one Hydrochloride)	SS		ORAL
ORAL		Suicide Attempt		Seroquel(Quetiapine)	SS		ORAL
ORAL		White Blood Cell Count		Acetaminophen(Parace tamol)	SS		

Date:10/10/03ISR Number: 4205867-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0422842A  
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Dexedrine	C	Glaxosmithkline	
3 WK							

Date:10/10/03ISR Number: 4205870-8Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0429044A  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Neutropenia		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
150MG Twice							
per day	MON			Adriamycin	SS		
UNKNOWN				Taxol	SS		
UNKNOWN							

Date:10/10/03ISR Number: 4205874-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0429227A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Psychotic Disorder	Wellbutrin	PS	Glaxosmithkline	ORAL
2	MON		Suicide Attempt	Alcohol Celexa	SS C		

Date:10/10/03ISR Number: 4205878-2Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0307425A  
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening			Headache	Zyban	PS	Glaxosmithkline	
150MG Twice							
Hospitalization -			Musculoskeletal Stiffness				
per day							
Initial or Prolonged			Somnolence Subarachnoid Haemorrhage	Valsartan	C		

Date:10/10/03ISR Number: 4205897-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0311254A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -			Road Traffic Accident	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/10/03ISR Number: 4209002-1Report Type:Direct  
Age:34 YR Gender:Male I/FU:I

Company Report #CTU 203607

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100MG 4.5 ORAL		Confusional State Depression Loss Of Employment		Bupropion 100 Mg Mylan	PS	Mylan	ORAL
100MG 4.5 ORAL		Pharmaceutical Product Complaint Sleep Disorder Somnolence		Bupropion 100 Mg Tevau	SS	Tevau	ORAL

Date:10/13/03ISR Number: 4207221-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0412091A  
Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	10 MON	Dizziness Drug Exposure During Pregnancy Fatigue Pregnancy Premature Baby Stillbirth		Wellbutrin Prenatal Vitamins	PS C	Glaxosmithkline	ORAL

Date:10/13/03ISR Number: 4207222-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0412091B  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Twice Congenital Anomaly per day		Abdominal Wall Anomaly Abdominal Wall Disorder Atrial Septal Defect Chorioamnionitis		Wellbutrin	PS	Glaxosmithkline	

Cleft Palate  
 Congenital Anomaly  
 Congenital Lymphoedema  
 Dizziness  
 Drug Exposure During  
 Pregnancy  
 Fatigue  
 Funnel Chest Acquired  
 Kyphosis  
 Lung Disorder  
 Lymphadenopathy  
 Placental Necrosis  
 Pregnancy  
 Premature Baby  
 Stillbirth

Date:10/13/03ISR Number: 4207224-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0427597A  
 Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complications Of Maternal		Bupropion	PS	Glaxosmithkline	
100MG Twice		Exposure To Therapeutic					
per day		Drugs					
		Intentional Misuse					
		Maternal Drugs Affecting					
		Foetus					
		Stillbirth					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/13/03ISR Number: 4207225-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0427597B

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	100MG Twice		Complications Of Maternal	Bupropion	PS	Glaxosmithkline	
	Congenital Anomaly per day		Exposure To Therapeutic				
			Drugs				
			Drug Exposure During				
			Pregnancy				
			Flushing				
			Muscle Atrophy				
			Overdose				
			Skin Maceration				
			Stillbirth				
			Umbilical Cord				
			Abnormality				

Date:10/13/03ISR Number: 4207229-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0429063A

Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day		Abortion Spontaneous	Bupropion	PS	Glaxosmithkline	
			Drug Exposure During				
			Pregnancy				
			Maternal Drugs Affecting				
			Foetus				
			Pregnancy				

Date:10/13/03ISR Number: 4207247-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0310773A

Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day 8 DAY		Atrial Fibrillation	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged			Congestive Cardiomyopathy				
			Dyspnoea				



Date:10/13/03ISR Number: 4207449-0Report Type:Expedited (15-DaCompany Report #PHEH2003US07744  
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose		Zelnorm	PS	Novartis Sector: Pharma	
6 mg, BID				Wellbutrin	SS		

Date:10/14/03ISR Number: 4207748-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0371291A  
Age:36 YR Gender:Female I/FU:F

Outcome	PT
Other	Confusional State Convulsion Crying Dysgeusia Emotional Disorder Gastric Disorder Grand Mal Convulsion Head Injury Insomnia Nausea

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300MG Twice per day	2 YR	Prescribed Overdose Psoriasis Rash Road Traffic Accident	Wellbutrin	PS	Glaxosmithkline	ORAL
		Thinking Abnormal				
		Vaginal Infection	Lamictal	SS	Glaxosmithkline	ORAL
		Vaginal Mycosis	Wellbutrin	SS	Glaxosmithkline	ORAL
		Vomiting	Pain Medication	C		ORAL
		Weight Decreased	Klonopin	C		ORAL
			Lorazepam	C		
			Ortho Tri-Cyclen	C		ORAL
TRANSDERMAL			Nicotine Patch	C	Glaxosmithkline	
			Nicotine Gum	C	Glaxosmithkline	ORAL

Date:10/14/03ISR Number: 4207780-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0426614A  
Age: Gender: I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
PT		Medication Error	Wellbutrin	PS	Glaxosmithkline	

Date:10/14/03ISR Number: 4209852-1Report Type:Expedited (15-DaCompany Report #2003AP03513  
Age:76 YR Gender: I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Atenolol	PS		
		Multiple Drug Overdose	Bupropion	SS		
			Amlodipine	SS		

Date:10/15/03ISR Number: 4208697-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0429976A  
Age: YR Gender: I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
PT		Medication Error	Wellbutrin	PS	Glaxosmithkline	

Date:10/15/03ISR Number: 4208698-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0429978A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	11 DAY		Hyperhidrosis	Wellbutrin	PS	Glaxosmithkline	ORAL
			Insomnia	Alprazolam	C		
			Medication Error				
			Pollakiuria				

Date:10/15/03ISR Number: 4208699-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0429983A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Medication Error	Wellbutrin	PS	Glaxosmithkline	ORAL
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/03ISR Number: 4208703-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0430268A

Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:10/15/03ISR Number: 4208723-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0311479A

Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL
Other		Drug Interaction		Microgynon	SS		ORAL
UNKNOWN	1TAB Per day	Unintended Pregnancy		No Concurrent Medication	C		

Date:10/15/03ISR Number: 4211352-XReport Type:Expedited (15-DaCompany Report #200318842BWH

Age:74 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Enterocolitis	Health	Cipro			
Life-Threatening Hospitalization - 100 MG, TOTAL		Haemorrhagic	Professional	(Ciprofloxacin)	PS		ORAL
Initial or Prolonged DAILY, ORAL 5 DAY Disability		Giardiasis	Other				
		Rash		Wellbutrin Sr (Bupropion Hydrochloride)	SS		
				Metronidazole	SS		
3 DAY							

Date:10/16/03ISR Number: 4209763-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0125519A

Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL

Disability	Bronchial Oedema	Acular	C		
INTRAOCULAR					
	Cardiac Failure	Vancenase Aq	C	Glaxosmithkline	NASAL
	Congestive	Diet Pep	C		
	Coagulopathy	Ginko Biloba	C		
	Coronary Artery				
	Atherosclerosis				
	Gait Disturbance				
	Hepatic Steatosis				
	Malaise				
	Myocardial Ischaemia				
	Nephrosclerosis				
	Polytraumatism				
	Pulmonary Congestion				
	Thrombotic				
	Thrombocytopenic Purpura				

Date:10/16/03ISR Number: 4209769-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0423710A  
Age:52 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL
		Depression		Luvox	C		
		Drug Toxicity		Klonopin	C		
1MG As		Overdose					
directed		Restlessness					
		Status Epilepticus					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/03ISR Number: 4209770-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0425135A

Age: YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:10/16/03ISR Number: 4209777-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0303990A

Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG See Initial or Prolonged dosage text 1 MON		Muscle Haemorrhage Muscle Rupture Phlebitis		Zyban	PS	Glaxosmithkline	ORAL

Date:10/16/03ISR Number: 4210542-XReport Type:Direct Company Report #CTU 203927

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other Required 150MG 2X Intervention to DAILY ORAL Prevent Permanent Impairment/Damage		Dizziness Nausea Pharmaceutical Product Complaint		Wellbutrin 150 Mg Glaxo	PS	Glaxo	ORAL

Date:10/17/03ISR Number: 4210624-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0429382A

Age:72 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization - Initial or Prolonged Other		Grand Mal Convulsion Headache Loss Of Consciousness		Wellbutrin Wellbutrin	PS SS	Glaxosmithkline Glaxosmithkline	ORAL

Date:10/17/03ISR Number: 4210625-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0429515A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Medication Error	Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day							

Date:10/17/03ISR Number: 4210626-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0429678A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Medication Error	Wellbutrin	PS	Glaxosmithkline	

Date:10/17/03ISR Number: 4210627-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0429687A  
Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Medication Error	Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/17/03ISR Number: 4214646-7Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20030904723  
 Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bradycardia Cardiac Arrest Decerebration	Literature Health Professional	Risperdal (Risperidone) Unspecified			ORAL
ORAL		Eye Movement Disorder Hyperpyrexia Mental Status Changes Multiple Drug Overdose Muscle Rigidity Pyrexia	Distributor	Paroxetine (Paroxetine) Bupropion (Bupropion) Clonazepam (Clonazepam)	PS SS SS SS		

Date:10/20/03ISR Number: 4211989-8Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0398320A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Electrocardiogram Qt Prolonged Grand Mal Convulsion Intentional Misuse		Bupropion Citalopram Methotrimeprazine	PS SS SS	Glaxosmithkline	

Date:10/20/03ISR Number: 4211993-XReport Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0427236A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Electrocardiogram Qt Prolonged		Bupropion Celexa	PS SS	Glaxosmithkline	ORAL
UNKNOWN	150MG	Unknown					
300MG	Unknown						
UNKNOWN		Grand Mal Convulsion Intentional Misuse		Methotrimeprazine	SS		

Date:10/20/03ISR Number: 4211994-1Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0427238A  
 Age: Gender: I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Unknown	Electrocardiogram Qt		Wellbutrin	PS	Glaxosmithkline	ORAL
	300MG Unknown	Prolonged		Celexa	SS		ORAL
UNKNOWN		Grand Mal Convulsion		Methotrimeprazine	SS		

Date:10/20/03ISR Number: 4212009-1Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0308919A  
Age:45 YR Gender:Male I/FU:F

Outcome	PT
Life-Threatening	Ageusia
Hospitalization -	Amnesia
Initial or Prolonged	Chest Discomfort
Other	Diarrhoea
	Dizziness
	Dyspnoea
	Feeling Abnormal
	Feeling Hot And Cold
	Hallucination
	Headache
	Immobile
	Lethargy
	Muscle Spasms
	Nausea

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Paraesthesia Respiratory Failure Thirst	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
8	DAY			Zyban	PS	Glaxosmithkline	ORAL
10MG per day	YR			Norvasc	C		ORAL

Date:10/20/03ISR Number: 4212015-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0311639A  
Age:44 YR Gender:Male I/FU:I

		PT Back Pain	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Death				Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	3 DAY	Cardio-Respiratory Arrest Myocardial Infarction Nausea					

Date:10/20/03ISR Number: 4212018-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0311648A  
Age:46 YR Gender:Female I/FU:I

		PT Cardiac Arrest	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Death				Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	WK	Pulmonary Oedema		Daonil	C		
				Glucophage	C		
				Kardegic	C		
				Mediator	C		ORAL
				Tenormine	C		
				Coversyl	C		
				Vastarel	C		

Date:10/20/03ISR Number: 4212026-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0429530A  
Age: Gender: I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose							

Hospitalization - Drug Exposure During Wellbutrin PS Glaxosmithkline  
200MG Per day  
Initial or Prolonged Pregnancy  
Tremor Neonatal

Date:10/20/03ISR Number: 4212027-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0429574A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Bupropion	PS	Glaxosmithkline	
150MG Twice		Drug Exposure During					
per day		Pregnancy					

Date:10/20/03ISR Number: 4212028-5Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0306220A  
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Inflammation		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown 2 MON		Joint Swelling					
Other		Musculoskeletal Pain					
		Streptococcal Infection					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/20/03ISR Number: 4212029-7Report Type:Expedited (15-DaCompany Report #DK-GLAXOSMITHKLINE-B0309119A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyskinesia Dysstasia Hypokinesia Musculoskeletal Discomfort		Zyban	PS	Glaxosmithkline	ORAL

Date:10/21/03ISR Number: 4215154-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0311425A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 2 WK Initial or Prolonged		Aggression  Personality Change		Zyban  Hrt	PS  C	Glaxosmithkline	ORAL

Date:10/22/03ISR Number: 4213784-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0412091A  
Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other 150MG Twice  per day	10 MON	Complications Of Maternal Exposure To Therapeutic Drugs Dizziness Drug Exposure During Pregnancy Fatigue Intra-Uterine Death		Wellbutrin  Prenatal Vitamins	PS  C	Glaxosmithkline	ORAL

Date:10/22/03ISR Number: 4213785-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0412091B  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death	Abdominal Wall Anomaly	Wellbutrin	PS	Glaxosmithkline
150MG Twice				
Congenital Anomaly	Arteriopathic Disease			
per day				
	Atrial Septal Defect			
	Chorioamnionitis			
	Cleft Palate			
	Drug Exposure During			
	Pregnancy			
	Foetal Disorder			
	Intra-Uterine Death			
	Jaw Fracture			
	Kyphosis			
	Lymphangiectasia			
	Macrogathia			
	Pectus Excavatum			
	Traumatic Delivery			

Date:10/22/03ISR Number: 4213789-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0428071A  
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	5 DAY		Medication Error	Wellbutrin	PS	Glaxosmithkline	ORAL
			Pharmaceutical Product Complaint	Wellbutrin	SS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/22/03ISR Number: 4213790-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0430630A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
		Medication Error					

Date:10/22/03ISR Number: 4213826-4Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0312169A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
Other		Visual Acuity Reduced					

Date:10/24/03ISR Number: 4217108-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0430978A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
		Medication Error					

Date:10/24/03ISR Number: 4217109-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0430981A

Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Medication Error					

Date:10/24/03ISR Number: 4217113-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0431158A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -		Amnesia					
300MG Per day 1 WK							
Initial or Prolonged		Convulsion		None	C		
		Joint Dislocation					
		Tongue Biting					

Date:10/24/03ISR Number: 4217115-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0431173A  
Age:36 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1 WK Initial or Prolonged	Dizziness	Road Traffic Accident	Wellbutrin Prozac	PS C	Glaxosmithkline	

Date:10/24/03ISR Number: 4217127-XReport Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0312102A  
Age:26 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 22 DAY Initial or Prolonged	Arthralgia Body Temperature Increased Eyelid Oedema Face Oedema Myalgia Oedema Peripheral Urticaria		Zyban	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/24/03ISR Number: 4219794-3Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20031003781

Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Sleep Walking	Consumer	Concerta (Methylphenidate Hydrochloride) Sustained Release Tablets	PS		ORAL
54 MG, 1 IN 1 DAY, ORAL				Wellbutrin (Bupropion Hydrochloride)	SS		
150 MG							

Date:10/24/03ISR Number: 4219800-6Report Type:Expedited (15-DaCompany Report #HQWYE065613AUG03

Age:1 DY Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Apgar Score Low Caesarean Section Feeding Problem In Newborn Maternal Drugs Affecting	Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		ORAL
75 MG 3X PER 1 DAY, ORAL		Foetus		Wellbutrin - Slow Release (Amfebutamone Hydrochloride)	SS		
TRANSPLACENTAL	150 MG 2X PER	Neonatal Apnoeic Attack Premature Baby					

1 DAY,

TRANSPLACENTA

L



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Caesarean Section Complications Of Maternal Exposure To Therapeutic Drugs Drug Ineffective	Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		ORAL
75 MG 3X PER		Pre-Eclampsia					
11DAY, ORAL		Pregnancy Premature Baby		Wellbutrin- Slow Release (Amfebutamone	SS		ORAL
150 MG 2X PER							
1 DAY, ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - Initial or Prolonged Disability Required Intervention to Prevent Permanent Impairment/Damage ORAL		Condition Aggravated Enterocolitis Haemorrhagic Rash	Health Professional	Flagyl (Metronidazole) Tablet Wellbutrin - Slow Release (Amfebutamone Hydrochloride)	PS   SS		ORAL
100 MG, QD,							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Ciprofloxacin  
Hydrochloride  
(Ciprofloxacin  
Hydrochloride) SS

FOR 5 DAYS

Date:10/27/03ISR Number: 4218507-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0430634A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	
		Pharmaceutical Product Complaint					

Date:10/27/03ISR Number: 4218509-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0431387A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:10/27/03ISR Number: 4218511-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0431427A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Depression		Paxil	PS	Glaxosmithkline	ORAL
80MG Per day	8 YR	Drug Ineffective		Wellbutrin	SS	Glaxosmithkline	ORAL
		Drug Withdrawal Syndrome		Paxil Cr	SS	Glaxosmithkline	ORAL
75MG Per day		Headache		Ortho Novum 1/35	C		
		Nausea		Advair	C	Glaxosmithkline	
		Panic Attack		Prednisone	C		
		Suicidal Ideation		Zyrtec	C	Glaxosmithkline	
		Thinking Abnormal		Flonase	C	Glaxosmithkline	
		Weight Increased		Albuterol	C	Glaxosmithkline	
				Ritalin	C		
				Multivitamin	C		

Date:10/27/03ISR Number: 4220887-5Report Type:Expedited (15-DaCompany Report #HQWYE337420OCT03  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Drug Interaction Unintended Pregnancy	Health Professional Other	Microgynon (Levonorgestrel/Ethinyl Estradiol, Tablet)	PS		ORAL
1 TABLET							
DAILY							
150MG				Zyban (Amfebutamone Hydrochloride, )	SS		ORAL
TABLETS;							

Date:10/28/03ISR Number: 4219549-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0423710A  
Age:52 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death							
75MG Per day	2 DAY	Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 DAY	Intentional Misuse		Wellbutrin	SS	Glaxosmithkline	ORAL
1MG As		Restlessness Status Epilepticus		Luvox Klonopin	C C		

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Freedom Of Information (FOI) Report

directed

4MG Twice per

day

200MG At

night

Gabapril

C

Seroquel

C

Date:10/28/03ISR Number: 4222755-1Report Type:Expedited (15-DaCompany Report #2003112988

Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension	Consumer	Lipitor	PS		ORAL
Other		Balance Disorder		(Atorvastatin)			
20 MG		Flatulence					
(DAILY), ORAL		Gastrointestinal Motility Disorder		Amoxicillin	SS		
		Gastroesophageal Reflux Disease		(Amoxicillin)			
		Haematemesis		Bupropion			
		Impaired Healing		Hydrochloride			
		Liver Function Test Abnormal		(Bupropion Hydrochloride)	SS		
		Memory Impairment		Levothyroxine Sodium			
		Muscular Weakness		(Levothyroxine Sodium)	C		
		Myalgia		Insulin (Insulin)	C		
		Pain		Dexamfetamine			
		Periodontal Disease		(Dexamfetamine)	C		

Date:10/28/03ISR Number: 4222758-7Report Type:Expedited (15-DaCompany Report #USA-2003-0006512

Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Consumer	Oxycontin			
		Coma	Health	Tablets(Oxycodone			
		Drug Toxicity	Professional	Hydrochloride) Cr			

	Hepatomegaly	Other	Tablet	PS	ORAL
40 MG, BID,					
ORAL	Myalgia				
	Portal Triaditis		Celexa (Citalopram Hydrobromide)	SS	
4 DAY	Post Procedural Pain				
	Postoperative Infection		Carisoprodol (Carisoprodol)	SS	
4 DAY	Pulmonary Congestion				
	Pulmonary Oedema		Meprobamate (Meprobamate)	SS	
4 DAY					
			Bupropion (Amfebutamone)	SS	
4 DAY					
			Biaxin (Clarithromycin)	C	

Date: 10/29/03  
 ISR Number: 4220678-5  
 Report Type: Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0430634A  
 Age:            Gender:            I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error		Wellbutrin	PS	Glaxosmithkline	
		Pharmaceutical Product Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/29/03ISR Number: 4220685-2Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0307425A  
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Twice	Aneurysm		Zyban	PS	Glaxosmithkline	
Hospitalization - per day		Blood Pressure Increased					
Initial or Prolonged		Bradycardia Headache Musculoskeletal Stiffness Somnolence Subarachnoid Haemorrhage		Valsartan	C		

Date:10/29/03ISR Number: 4223117-3Report Type:Expedited (15-DaCompany Report #USA-2003-0007154  
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose Alcohol Use Atherosclerosis Brain Herniation Brain Oedema Cardiomegaly Congestive Cardiomyopathy Drug Abuser Drug Screen Positive Drug Toxicity Ecchymosis Scab	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553) (Oxycodone Hydrochloride) Bupropion (Amfebutamone) Ethanol (Ethanol)	PS SS SS		

Date:10/29/03ISR Number: 4223361-5Report Type:Expedited (15-DaCompany Report #USA-2003-0007532  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiomegaly Completed Suicide Intentional Misuse Multiple Drug Overdose Pulmonary Congestion	Health Professional Other	Oxycodone Hydrochloride Hydrocodone Bitartrate (Similar To Ind	PS		

59,175) (Naltrexone,  
Paracetamol, SS  
Acetaminophen (Parace  
tamol) SS  
Mirtazapine (Mirtazap  
ine) SS  
Alprazolam (Alprazola  
m) SS  
Olanzapine (Olanzapin  
e) SS  
Sertraline (Sertralin  
e) SS  
Trazodone (Trazodone) SS  
Bupropion (Amfebutamo  
ne) SS

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/30/03ISR Number: 4222178-5Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0398320A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Aspiration		Wellbutrin	PS	Glaxosmithkline	
Initial or Prolonged		Depressed Level Of Consciousness		Citalopram	SS		
Other		Dystonia		Methotrimeprazine	SS		
		Electrocardiogram Qt Prolonged					
		Fall					
		Grand Mal Convulsion					
		Intentional Misuse					

Date:10/30/03ISR Number: 4222187-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0432011A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	1 DAY	Confusional State		Wellbutrin	PS	Glaxosmithkline	ORAL
		Lethargy					
		Medication Error					

Date:10/30/03ISR Number: 4222193-1Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0312414A  
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Bradycardia		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Cardiac Enzymes Increased		Ascorbic Acid	C	Glaxosmithkline	
		Epilepsy		Aspirin Junior	C	Glaxosmithkline	
		Malaise		Multivitamin With Zinc	C		
		Syncope		Selenium	C		
		Ventricular Fibrillation		Glucosamine Sulphate	C		
		White Blood Cell Count Increased					

Date:10/30/03ISR Number: 4223978-8Report Type:Expedited (15-DaCompany Report #S03-USA-04440-01  
Age:17 YR Gender:Male I/FU:I



Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
10 MG QD PO		Confusional State Convulsion	Health Professional	Lexapro (Escitalopram)	PS		ORAL
200 MG BID		Gastrooesophageal Reflux Disease Staring		Wellbutrin (Bupropion Hydrochloride)	SS		
100 MG BID		Throat Irritation		Wellbutrin (Bupropion Hydrochloride)	SS		
100 MG QD				Wellbutrin (Bupropion Hydrochloride)	SS		
				Neurontin (Gabapentin)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/31/03ISR Number: 4223303-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0357840A

Age:37 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 5 MON	Abdominal Discomfort		Paxil	PS	Glaxosmithkline	
Hospitalization - Initial or Prolonged	Abortion Spontaneous		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
Other .5MG As required	Aggression		Effexor	C		
	Agitation		Ativan	C		
	Agoraphobia					
25MG Per day	Alcoholism		Atenolol	C		
	Anger		Alcohol	C		
	Anxiety		Valium	C		ORAL
	Blood Creatinine Decreased					
	Blood Sodium Decreased					
	Chest Discomfort					
	Confusional State					
	Convulsion					
	Dependence					
	Dizziness					
	Drug Abuser					
	Drug Dependence					
	Drug Ineffective					
	Drug Withdrawal Syndrome					
	Dyspepsia					
	Dyspnoea					
	Feeling Hot					
	Flat Affect					
	Hypoaesthesia					
	Influenza					
	Insomnia					
	Intentional Self-Injury					
	Ketoacidosis					
	Loss Of Consciousness					
	Mental Disorder					
	Nausea					
	Nightmare					
	Panic Disorder					
	Paraesthesia					
	Pco2 Decreased					
	Platelet Count Increased					
	Pyrexia					

Suicidal Ideation  
Suicide Attempt  
Tremor  
Urine Analysis Abnormal

Date:10/31/03ISR Number: 4223315-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0424693A

Age:21 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	2 DAY	Atrioventricular Block Second Degree Bradycardia Fatigue Syncope Vasovagal		Wellbutrin	PS	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/31/03ISR Number: 4223317-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0426335A  
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Amnesia					
16 WK				Zyprexa	C		
10MG Per day		Drug Ineffective					
600MG Twice		Grand Mal Convulsion		Trileptal	C		
per day		Tongue Biting					

Date:10/31/03ISR Number: 4223318-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0428237A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Medication Error					
150MG Per day	5 DAY			Wellbutrin	SS	Glaxosmithkline	

Date:10/31/03ISR Number: 4223319-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0431158A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -		Amnesia					
300MG Per day	1 WK			None	C		
Initial or Prolonged		Convulsion					
		Joint Dislocation					
		Tongue Biting					

Date:10/31/03ISR Number: 4223326-3Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0307425A  
 Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Life-Threatening 150MG Twice	Aneurysm	Zyban	PS	Glaxosmithkline
Hospitalization - per day	Bradycardia			
Initial or Prolonged	Headache	Valsartan	C	
	Musculoskeletal Stiffness			
	Somnolence			
	Subarachnoid Haemorrhage			

Date:10/31/03ISR Number: 4224878-XReport Type:Expedited (15-DaCompany Report #KII-2003-0004112  
Age:43 YR Gender:Male I/FU:I

Outcome	PT
Death	Acid Base Balance
Life-Threatening	Abnormal
Hospitalization -	Blood Glucose Abnormal
Initial or Prolonged	Blood Lactic Acid
Other	Increased
	Blood Pressure Decreased
	Blood Pressure Increased
	Body Temperature
	Increased
	Bradycardia
	Cardiac Arrest
	Coma
	Depression
	Electrocardiogram Qt
	Corrected Interval
	Prolonged
	Electroencephalogram
	Abnormal

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Heart Rate Increased Miosis Multiple Drug Overdose	Report Source	Product	Role	Manufacturer	Route
		Muscle Twitching	Health	Oxycontin Cr	PS		
		Mydriasis	Professional	Methadone			
		Posturing	Other	(Methadone)	SS		
		Pupil Fixed		Trazodone(Trazodone)	SS		
400 MG		Respiratory Rate Decreased		Gabapentin(Gabapenti n)	SS		
200 MG		Respiratory Rate Increased		Quetiapine(Quetiapin e)	SS		
20 MG				Citalopram(Citalopra m)	SS		
1 MG				Clonazepam (Clonazepam)	SS		
4 MG, ORAL				Zanaflex(Tizanidine Hydrochloride)	SS		ORAL
150 MG				Bupropion(Amfebutamo ne)	SS		
10 MG				Cyclobenzaprine(Cycl obenzaprine)	SS		
50 MG				Amitriptyline(Amitri ptyline)	SS		
				Estalopram()	SS		

Date:10/31/03ISR Number: 4225127-9Report Type:Direct  
Age: Gender: I/FU:I

Company Report #USP 56117

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin Sr	PS	Glaxo Smithkline	
TABLET,EXTEND							
ED RELEASE							
TABLET,				Wellbutrin Xl	SS	Glaxosmithkline	

EXTENDED

RELEASE

Date:11/03/03ISR Number: 4224539-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0431972A  
Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	5 YR	Arthritis					
		Psoriasis		Synthroid	C	Glaxosmithkline	
				Atenolol	C		
				Cozar	C		
				Thiazide	C		
				Fosamax	C		

Date:11/03/03ISR Number: 4224540-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0432114A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Clumsiness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day		Dysgraphia					
		Muscle Rigidity		Risperdal	C		
		Neuroleptic Malignant Syndrome					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/03ISR Number: 4224542-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0432276A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:11/03/03ISR Number: 4224546-4Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0309306A

Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Discomfort		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	5 DAY						
Initial or Prolonged		Respiratory Distress		Duphaston	C		ORAL
1UNIT See							
		Urticaria Generalised					
dosage text							

Date:11/03/03ISR Number: 4224548-8Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0309578A

Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Apathy		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
Initial or Prolonged		Asthenia					
per day	9 DAY						
		Tongue Disorder		Alcohol	C		
		Tremor		Tobacco	C		
		Vertigo					

Date:11/03/03ISR Number: 4224556-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0312731A

Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Psoriasis		Zyban	PS	Glaxosmithkline	ORAL
150MG per day	7 DAY						
				Bendrofluazide	C	Glaxosmithkline	ORAL
2.5MG per day							



Date:11/03/03ISR Number: 4225263-7Report Type:Direct Company Report #CTU 205089  
Age:15 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Urticaria		Wellbutrin XL	PS		
1 PER DAY							
Intervention to Prevent Permanent Impairment/Damage							

Date:11/03/03ISR Number: 4226117-2Report Type:Expedited (15-DaCompany Report #2003-DE-05117GD  
Age:20 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Completed Suicide	Literature	Theophylline	PS		
Death		Multiple Drug Overdose		Bupropion (Amfebutamone)	SS		
				Heroin (Diamorphine0	SS		

Date:11/03/03ISR Number: 4226470-XReport Type:Expedited (15-DaCompany Report #K200301715  
Age:20 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Completed Suicide	Literature	Quibron-T/Sr			
Death		Intentional Misuse	Health Professional	(Theophylline) Tablet, 300mg	PS		ORAL
ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL  
 INHALATION  
 Bupropion (Amfebutamone) SS  
 Heroin (Diamorphine) SS

Date:11/03/03ISR Number: 4226849-6Report Type:Expedited (15-DaCompany Report #2003-DE-05279GD  
 Age:42 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature	Methadone (Methadone)	PS		ORAL
PO		Intentional Misuse		Diazepam Intensol (Diazepam)	SS		
				Bupropion (Amfebutamone)	SS		

Date:11/03/03ISR Number: 4226988-XReport Type:Expedited (15-DaCompany Report #2003-DE-05212GD  
 Age:47 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature	Alprazolam	PS		
		Completed Suicide		Methadone (Methadone)	SS		
		Multiple Drug Overdose		Bupropion (Amfebutamone)	SS		

Date:11/03/03ISR Number: 4227089-7Report Type:Expedited (15-DaCompany Report #2003-DE-05417GD  
 Age:38 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Literature	Alprazolam	PS		ORAL
PO		Completed Suicide		Paracetamol (Paracetamol)	SS		ORAL
		Respiratory Arrest		Bupropion			

PO

Date:11/04/03ISR Number: 4225469-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0303990A  
Age:45 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Muscle Haemorrhage		Zyban	PS	Glaxosmithkline	ORAL
150MG See						
Initial or Prolonged	Muscle Rupture					
dosage text	1 MON					

Date:11/04/03ISR Number: 4225471-5Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0312384A  
Age:51 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Life-Threatening	Hepatic Trauma		Zyban	PS	Glaxosmithkline	ORAL
5 WK						
Hospitalization -	Pelvic Fracture					
Initial or Prolonged	Suicide Attempt					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/04/03ISR Number: 4225476-4Report Type:Expedited (15-DaCompany Report #FI-GLAXOSMITHKLINE-B0313032A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema		Zyban	PS	Glaxosmithkline	ORAL
1TAB See		Oropharyngeal Swelling					
dosage text	15	DAY					

Date:11/04/03ISR Number: 4227174-XReport Type:Expedited (15-DaCompany Report #031023-PM0146-00  
Age:45 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardio-Respiratory Arrest	Literature	Tranxene			
		Completed Suicide	Health	(Clorazepate			
			Professional	Dipotassium)	PS		ORAL
PO				Doxepin	SS		ORAL
PO				Bupropion			
				(Long-Acting)	SS		ORAL
PO							

Date:11/05/03ISR Number: 4226759-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0432609A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:11/05/03ISR Number: 4226760-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0432618A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Lymphadenopathy		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Twice		Pitting Oedema					
per day	3	WK					

Date:11/05/03ISR Number: 4226772-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0312762A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Agitation		Zyban	PS	Glaxosmithkline	ORAL
150MG See							
Hospitalization -		Anxiety					
dosage text	3 WK						
Initial or Prolonged		Insomnia					
		Overdose					
		Suicidal Ideation					
		Suicide Attempt					

Date:11/05/03ISR Number: 4226779-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0313313A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Discomfort		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	11 DAY						
		Dizziness					
		Dyspnoea					
		Feeling Abnormal					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/06/03ISR Number: 4227868-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0354550A

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 20MG Per day	Abdominal Discomfort		Paxil	PS	Glaxosmithkline	ORAL
Initial or Prolonged 150MG Twice	Aggression		Wellbutrin	SS	Glaxosmithkline	ORAL
Other per day	Agitation					
	Anger		Ritalin	C		
	Anorexia		Amoxicillin	C	Glaxosmithkline	
	Anxiety					
	Asthenia					
	Confusional State					
	Coordination Abnormal					
	Depression					
	Disturbance In Attention					
	Dizziness					
	Drug Ineffective					
	Drug Withdrawal Syndrome					
	Dystonia					
	Fatigue					
	Gait Disturbance					
	Homicidal Ideation					
	Influenza					
	Intentional Self-Injury					
	Irritability					
	Lethargy					
	Malaise					
	Memory Impairment					
	Myalgia					
	Nausea					
	Nightmare					
	Obsessive Thoughts					
	Obsessive-Compulsive Disorder					
	Paraesthesia					
	Paranoia					
	Pyrexia					
	Suicidal Ideation					
	Suicide Attempt					
	Tremor					
	Vertigo					

Outcome	PT
Hospitalization -	Cataract
Initial or Prolonged	Confusional State
Other	Crying
	Dissociation
	Drooling
	Drug Ineffective
	Dysgeusia
	Dyskinesia
	Emotional Disorder
	Excoriation
	Feeling Jittery
	Gastric Disorder
	Grand Mal Convulsion
	Head Injury
	Insomnia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Twice per day	2 YR	Mania Musculoskeletal Stiffness Nail Disorder					
		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
		Pain					
600MG per day		Prescribed Overdose		Lamictal	SS	Glaxosmithkline	ORAL
		Psoriasis		Wellbutrin	SS	Glaxosmithkline	ORAL
		Road Traffic Accident		Pain Medication	SS		ORAL
		Scratch		Wellbutrin	SS	Glaxosmithkline	ORAL
		Thinking Abnormal		Ultracet	SS		
		Tooth Loss		Seroquel	SS		ORAL
		Vaginal Infection		Nicotine Gum	SS	Glaxosmithkline	ORAL
		Vaginal Mycosis		Zyprexa	SS		ORAL
		Vomiting		Klonopin	C		ORAL
		Weight Decreased		Lorazepam	C		
		Weight Increased		Ortho Tri-Cyclen	C		ORAL
TRANSDERMAL				Nicotine Patch	C	Glaxosmithkline	
				Effexor	C		
				Lescol	C		

Date:11/06/03ISR Number: 4227922-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0313314A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 6 DAY		Convulsion		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged Disability		Facial Palsy		Thyroxine	C	Glaxosmithkline	
		Monoplegia					

Date:11/06/03ISR Number: 4227923-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0313315A  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Twice		Arthralgia		Zyban	PS	Glaxosmithkline	ORAL



per day 18 DAY Eye Swelling  
UNKNOWN Rash Pruritic  
Swelling Face

Viagra C

Date:11/06/03ISR Number: 4227925-4Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0313519A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Overdose		Zyban	PS	Glaxosmithkline	

Date:11/07/03ISR Number: 4228850-5Report Type:Expedited (15-DaCompany Report #PHEH2003US07744  
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 6 mg, BID		Completed Suicide Drug Screen Positive		Zelnorm	PS	Novartis Sector: Pharma	
150 mg, BID	87840MIN	Overdose		Wellbutrin	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229253-XReport Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0095466A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Chest Pain					
UNKNOWN		Heart Rate Increased		Singulair	SS		

Date:11/07/03ISR Number: 4229254-1Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0124877A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
450MG per day	7 MON	Muscle Atrophy					
		Rhabdomyolysis					
		Weight Decreased					

Date:11/07/03ISR Number: 4229255-3Report Type:Periodic  
 Age:49 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0145456A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
250MG per day		Photosensitivity Reaction					
20MG Per day				Prozac	C		
80MG Twice				Oxycontin	C		
per day				Antibiotic	C		

Date:11/07/03ISR Number: 4229256-5Report Type:Periodic  
 Age:51 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0150345A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Twice		Cheilitis		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	5	DAY	Dizziness				
			Drug Ineffective	Synthroid	C	Glaxosmithkline	
			Herpes Simplex	Estrogen	C		
			Influenza Like Illness	Benzalin	C		
			Lip Pain	Estratest	C		
			Nausea				
			Somnolence				

Date:11/07/03ISR Number: 4229261-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0154448A  
 Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Depression					
300MG Twice		Drug Ineffective		Lithium	C	Glaxosmithkline	ORAL
per day		Pharmaceutical Product					
		Complaint		Klonopin	C		ORAL

Date:11/07/03ISR Number: 4229267-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0156099A  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1 WK	Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Fatigue		Synthroid	C	Glaxosmithkline	
		Hypoaesthesia					
		Myalgia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229268-1Report Type:Periodic  
Age:52 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0157048A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness		Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Twice							
per day	13	DAY					
		Rash					
		Tinnitus					

Date:11/07/03ISR Number: 4229269-3Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0360415A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erythema Multiforme		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	13	DAY					
		Stevens-Johnson Syndrome					

Date:11/07/03ISR Number: 4229270-XReport Type:Periodic  
Age:32 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0369580A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Feeling Abnormal	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1	DAY					
		Headache					
		Nausea		Wellbutrin	C	Glaxosmithkline	ORAL
100MG Twice							
per day	3	DAY					
		Pain In Jaw					
		Paraesthesia					
		Paranoia					
		Pharyngolaryngeal Pain					

Date:11/07/03ISR Number: 4229271-1Report Type:Periodic  
Age:62 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0371577A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400MG See			Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
dosage text		10 YR	Anxiety					
			Dizziness		Premarin	C		
			Drug Ineffective		Lopressor	C		
			Euphoric Mood		Trileptal	C		
			Rash		Buspar	C		
			Rosacea		Neurontin	C		
			Stress					
			Vision Blurred					

Date:11/07/03ISR Number: 4229272-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0372805A  
Age:33 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Anaphylactic Reaction	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice			Oedema Peripheral	Professional				
per day		3 WK	Pruritus					
			Rash Generalised					
			Swelling Face					

Date:11/07/03ISR Number: 4229279-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0374355A  
Age:56 YR Gender:Female I/FU:F

Outcome	PT
	Abdominal Discomfort
	Abdominal Pain Upper

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	143 DAY	Eructation Feeling Hot Food Intolerance Hepatocellular Damage Weight Decreased	Wellbutrin	PS	Glaxosmithkline	ORAL
			Glucophage	C		
			Amaryl	C		
			Avandia	C	Glaxosmithkline	
			Calcium	C		
			Vitamin	C		
			Prinivil	C		
			Levoxyl	C	Glaxosmithkline	
			Unknown Medication	C		

Date:11/07/03ISR Number: 4229280-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0375304A  
 Age:41 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose Other 400MG Per day	YR	Dry Mouth Hyperhidrosis Rash	Wellbutrin	PS	Glaxosmithkline	ORAL
			Atacand	C		
			Orth Novum 777	C		
			Amitriptyline	C		

Date:11/07/03ISR Number: 4229281-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0375429A  
 Age: Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose 150MG Unknown		Drug Ineffective Pharmaceutical Product Complaint	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229282-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0376012A  
 Age:46 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	100MG	Per day	13 DAY	Anxiety	Wellbutrin	PS	Glaxosmithkline	ORAL
	25MG	Unknown		Chest Pain	Imipramine	C		ORAL
				Dyspnoea	Lipitor	C		
				Swelling Face				
				Urticaria				

Date:11/07/03ISR Number: 4229284-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0377324A  
Age:24 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG	Three		Grand Mal Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
		times per day	6 MON		Xanax	C		ORAL
	.25MG	As						
		required			Oral Contraceptive	C		
					Zyrtec	C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229285-1Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0379148A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Twice		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	6	AMnesia					
		Tremor					

Date:11/07/03ISR Number: 4229286-3Report Type:Periodic  
Age:64 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0379578A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Twice		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Stool Analysis Abnormal					
				Multivitamin	C		
				Calcium	C		
				Fosamax	C		
70MG Weekly				Effexor	C		ORAL
75MG Twice							
per day							
				Activelle	C		ORAL
1MG Per day							

Date:11/07/03ISR Number: 4229287-5Report Type:Periodic  
Age:64 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0379734A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Ejaculation Delayed		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2	Insomnia					
		Irritability					
		Tremor					



Date:11/07/03ISR Number: 4229288-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0380279A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropion	PS	Glaxosmithkline	ORAL
Other		Convulsion		Depakote	SS		ORAL

Date:11/07/03ISR Number: 4229289-9Report Type:Periodic  
Age:34 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0380405A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
21 DAY							
Other							

Date:11/07/03ISR Number: 4229290-5Report Type:Periodic  
Age:51 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0380854A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Flatulence		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Gastrointestinal Pain					
		Pharmaceutical Product Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229291-7Report Type:Periodic  
Age:57 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0381642A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Alkaline		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
50MG Per day		Phosphatase Increased		Zoloft	SS		ORAL
200MG Per day		Fatigue Hepatitis Lethargy Liver Function Test Abnormal Weight Decreased					

Date:11/07/03ISR Number: 4229292-9Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382141A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Twice		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 WK			Entex Pse	C		
				Humibid La	C		
				Acyclovir	C	Glaxosmithkline	
				Ibuprofen	C	Glaxosmithkline	
				Trazodone	C		
				Claritin	C		

Date:11/07/03ISR Number: 4229293-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382142A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hypothyroidism		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229294-2Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382155A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Stomatitis		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2	MON					

Date:11/07/03ISR Number: 4229295-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0382156A  
 Age:26 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	4	DAY		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229296-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0382157A  
 Age:34 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Weight Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2	YR		Vioxx	C		
UNKNOWN	25MG	Unknown		Synthroid	C	Glaxosmithkline	
				Multivitamin	C		
				Claritin D	C		
				Calcium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229297-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382159A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229298-XReport Type:Periodic  
 Age:13 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382162A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	
Other				Diflucan	SS		ORAL
250MG Per day	1 YR						

Date:11/07/03ISR Number: 4229299-1Report Type:Periodic  
 Age:31 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0382178A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR						

Date:11/07/03ISR Number: 4229300-5Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0382181A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gait Disturbance		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	4 MON	Hypoaesthesia		Estrogen	C		
		Nausea					
		Paraesthesia					
		Paranoia					
		Tinnitus					

Date:11/07/03ISR Number: 4229301-7Report Type:Periodic  
 Age:17 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0382233A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day	1 DAY	Pharmaceutical Product		Wellbutrin	PS	Glaxosmithkline	ORAL
		Complaint Somnolence Urticaria					

Date:11/07/03ISR Number: 4229302-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0382262A  
 Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Bruxism		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229303-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0382282A  
 Age:39 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	6 MON			Zoloft	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229304-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382336A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Unknown		Feeling Abnormal					
		Nervousness					

Date:11/07/03ISR Number: 4229305-4Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382337A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	1 WK	Muscle Contractions					
		Involuntary		Synthroid	C	Glaxosmithkline	
				Allegra	C		

Date:11/07/03ISR Number: 4229306-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382399A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Pharmaceutical Product		Unknown Medication	C		
		Complaint					
		Stool Analysis Abnormal					

Date:11/07/03ISR Number: 4229307-8Report Type:Periodic  
Age:51 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0382431A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Pharmaceutical Product					
per day	5 MON	Complaint					
		Stool Analysis Abnormal		Imitrex	C	Glaxosmithkline	
				Ambien	C		

Date:11/07/03ISR Number: 4229309-1Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0382447A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Urinary Tract Infection		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229310-8Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382455A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Paraesthesia					
per day	17 DAY			Menest	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229311-XReport Type:Periodic  
Age:53 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0382461A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Pharmaceutical Product		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	2 WK	Complaint		Lipitor	C		
		Stool Analysis Abnormal		Ziac	C		
				Paxil	C	Glaxosmithkline	
				Xanax	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229312-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382463A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229313-3Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382467A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	WK	Middle Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Estrogen	C		

Date:11/07/03ISR Number: 4229314-5Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382468A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	3 WK	Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
				Klonopin	C		

Date:11/07/03ISR Number: 4229315-7Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0382496A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Abuser Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229316-9Report Type:Periodic  
Age:41 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0382605A



Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	2 YR	Pharmaceutical Product Complaint Stool Analysis Abnormal Vomiting		Wellbutrin	PS	Glaxosmithkline	ORAL
				Glucophage	C		
				Neurontin	C		
				Betapace	C	Glaxosmithkline	
				Lanoxin	C	Glaxosmithkline	
				Buspar	C		

Date:11/07/03ISR Number: 4229318-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0382608A  
Age:40 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	3 MON	Agitation Insomnia Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229319-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0382653A  
Age:18 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229320-0Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382681A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice				Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3	MON					
				Mobic	C		

Date:11/07/03ISR Number: 4229321-2Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382684A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice				Wellbutrin	PS	Glaxosmithkline	ORAL
per day	6	WK					

Date:11/07/03ISR Number: 4229322-4Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382694A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
1 YR				Wellbutrin	PS	Glaxosmithkline	ORAL
				Paxil	C	Glaxosmithkline	
				Ambien	C		
				Hrt	C		

Date:11/07/03ISR Number: 4229323-6Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382778A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day				Wellbutrin	PS	Glaxosmithkline	ORAL
				Hormone Replacement	C		

Date:11/07/03ISR Number: 4229324-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382789A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229325-XReport Type:Periodic  
Age:11 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382801A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Effect Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day							
10MG Per day				Adderall	C		
50MG At night				Trazodone	C		
10MG Twice				Adderall Xr	C		
per day							

Date:11/07/03ISR Number: 4229326-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382804A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Irritability		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229327-3Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382811A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Urticaria					
per day	16 DAY			Tylenol	SS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229328-5Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382835A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -		Intentional Misuse	Professional	Celexa	SS		
Initial or Prolonged				Alcohol	SS		
Other							

Date:11/07/03ISR Number: 4229329-7Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382843A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day							

Date:11/07/03ISR Number: 4229330-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382844A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229331-5Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382846A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Fall Salivary Hypersecretion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229332-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0382852A  
 Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	4 YR	Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
		Fatigue		Zoloft	C		
		Headache		Klonopin	C		
		Somnolence					

Date:11/07/03ISR Number: 4229333-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0382858A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG per day		Thrombocytopenia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Effexor	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229334-0Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382860A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day							

Date:11/07/03ISR Number: 4229335-2Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382896A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	6 WK						

Date:11/07/03ISR Number: 4229336-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382900A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Jittery		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229337-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382930A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	5 DAY						

Date:11/07/03ISR Number: 4229338-8Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0382969A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK			Vitamins	C		

Date:11/07/03ISR Number: 4229339-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0383080A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia Strabismus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229340-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0383081A  
 Age:19 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Urticaria					

Date:11/07/03ISR Number: 4229341-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0383086A  
 Age: Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229342-XReport Type:Periodic  
Age:69 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383169A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	MON			Lexapro	C		
5MG Per day				Vitamins	C		
				Lorazepam	C		

Date:11/07/03ISR Number: 4229343-1Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383255A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Female Sexual Dysfunction		Wellbutrin	PS	Glaxosmithkline	ORAL
1 MON				Celexa	SS		
UNKNOWN		Urticaria					

Date:11/07/03ISR Number: 4229344-3Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383267A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day				Diltiazem	C	Glaxosmithkline	
				Toprol Xl	C		
				Prevacid	C		
				Lipitor	C		
				Plavix	C		
				Cyclobenzaprine	C		
				Darvocet N	C		



Date:11/07/03ISR Number: 4229345-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0383269A  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyskinesia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229346-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0383403A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Body Dysmorphic Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Disturbance In Attention					

Date:11/07/03ISR Number: 4229347-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0383404A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
400MG per day 4 YR		Depression		Xanax	C		ORAL
.25MG		Diarrhoea					
Variable dose		Medication Residue					
		Pharmaceutical Product					
		Complaint					
		Stool Analysis Abnormal					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229349-2Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383427A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Twice	Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
	per day	Irritability					
	5 YR						

Date:11/07/03ISR Number: 4229350-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383431A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menorrhagia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229351-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383441A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229352-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383449A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
				Paxil	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229353-4Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383566A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Twice	Loss Of Libido		Wellbutrin	PS	Glaxosmithkline	ORAL

Sexual Dysfunction

per day 1 YR

Clonazepam C

UNKNOWN 2MG Twice per

day

Date:11/07/03ISR Number: 4229354-6Report Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0383567A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bipolar Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Dyspnoea					
per day		Urticaria					

Date:11/07/03ISR Number: 4229355-8Report Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0383598A

Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Carpal Tunnel Syndrome		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Plantar Fasciitis					
per day				Valtrex	C	Glaxosmithkline	
				Buspar	C		
				Ritalin	C		
				Trazodone	C		
				Lamictal	C	Glaxosmithkline	
				Vivactil	C		
				Hctz	C		

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Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229356-XReport Type:Periodic  
Age:47 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0383603A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Aphthous Stomatitis		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Three							
times per day	7	DAY					
				Remeron	C		
				Paxil Cr	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229357-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383634A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							
		Hypertension		Zocor	C		
				Prevacid	C		

Date:11/07/03ISR Number: 4229358-3Report Type:Periodic  
Age:41 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0383637A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day	DAY						
		Pharmaceutical Product		Lithium	C	Glaxosmithkline	
		Complaint		Risperdal	C		
		Stool Analysis Abnormal		Effexor Xr	C		
				Ativan	C		
				Buspar	C		
				Geodon	C		

Date:11/07/03ISR Number: 4229359-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383659A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Per day 3 DAY	Eye Movement Disorder	Wellbutrin	PS	Glaxosmithkline	ORAL
	Hypoaesthesia				
	Rash Vesicular				

Date:11/07/03ISR Number: 4229360-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0383667A  
 Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day 2 WK		Dyspepsia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Fatigue		Nitro Patch	C	Glaxosmithkline	
		Feeling Abnormal		Nitrostat	C	Glaxosmithkline	
		Headache		Atenolol	C		
		Laziness		Multi-Vitamins	C		
				Ranitidine	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229361-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0383709A  
 Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	DAY	Hyperhidrosis		Wellbutrin	PS	Glaxosmithkline	ORAL
		Memory Impairment					
		Nausea					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229362-5Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383720A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day	2 MON	Back Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
		Chest Discomfort		Klonopin	C		
		Dysphagia		Buspirone	C		
		Eructation					
		Musculoskeletal Stiffness					

Date:11/07/03ISR Number: 4229367-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383738A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Nausea					
		Restlessness		Paxil	SS	Glaxosmithkline	ORAL
		Tremor					

Date:11/07/03ISR Number: 4229368-6Report Type:Periodic  
Age:50 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0383766A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
250MG per day	6 YR	Chest Discomfort		Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Ineffective					
		Dysphagia					
		Dyspnoea					
		Feeling Abnormal					
		Insomnia					
		Respiratory Failure					
		Speech Disorder					

Date:11/07/03ISR Number: 4229369-8Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383767A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1	MON					

Date:11/07/03ISR Number: 4229370-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0383774A  
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
150MG Twice							
per day							

Date:11/07/03ISR Number: 4229371-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0383800A  
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1	Swelling					
		Urticaria					
	DAY						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229376-5Report Type:Periodic  
Age:69 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383802A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG per day	3 DAY	Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Per day		Nightmare		Moduretic	C		ORAL
40MG Per day		Tinnitus		Zocor	C		
				Vitamin	C		

Date:11/07/03ISR Number: 4229378-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383808A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Coordination Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
				Paxil	SS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229379-0Report Type:Periodic  
Age:20 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383809A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Balance Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	5 MON	Dizziness					
5 DAY		Hypoaesthesia		Zithromax	C		
		Nausea					
		Pollakiuria					
		Weight Decreased					

Date:11/07/03ISR Number: 4229386-8Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0383849A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Feeling Abnormal  
Feeling Jittery  
Headache

Wellbutrin

PS

Glaxosmithkline

ORAL

Date:11/07/03ISR Number: 4229390-XReport Type:Periodic  
Age:40 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0383881A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Pharmaceutical Product	Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice			Complaint				
per day			Stool Analysis Abnormal				

Date:11/07/03ISR Number: 4229391-1Report Type:Periodic  
Age:46 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0383886A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Alopecia	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day				Concerta	C		
36MG Per day				Multivitamin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229392-3Report Type:Periodic  
Age:47 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0383930A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Bupropion	PS	Glaxosmithkline	ORAL
1 YR							
		Drug Interaction		Paxil	SS	Glaxosmithkline	ORAL
1 YR							
		Road Traffic Accident					

Date:11/07/03ISR Number: 4229394-7Report Type:Periodic  
Age:47 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0383950A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229395-9Report Type:Periodic  
Age:18 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383963A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Per day	1 WK	Drug Screen False		Wellbutrin	PS	Glaxosmithkline	ORAL
		Positive					

Date:11/07/03ISR Number: 4229396-0Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0383966A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Asthenia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	6 MON	Balance Disorder					
		Dizziness		Morphine	C		
		Feeling Abnormal		Robaxin	C		
500MG Twice							
per day		Feeling Drunk					

	Flat Affect	Gabapentin	C
	Sedation	Tegretol	C
50MG At night	Transient Ischaemic	Hydroxyzine	C
	Attack	Amitriptyline	C
50MG At night			

Date:11/07/03ISR Number: 4229397-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0383969A  
 Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
400MG per day	6 WK						

Date:11/07/03ISR Number: 4229398-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0383981A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Cheilitis		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	WK						

Date:11/07/03ISR Number: 4229399-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0384016A  
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG In the		Insomnia					
morning	6 WK	Mood Swings		Prenatal Vitamins	C		
				Progestin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Protonix C  
Amoxicillin C Glaxosmithkline

Date:11/07/03ISR Number: 4229400-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0384025A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day							

Date:11/07/03ISR Number: 4229401-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0384026A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharmaceutical Product		Wellbutrin	PS	Glaxosmithkline	ORAL
2 MON		Complaint					
		Stool Analysis Abnormal					

Date:11/07/03ISR Number: 4229402-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0384031A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229403-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0384034A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Food Poisoning					
150MG Twice							
per day							

30MG Per day Vomiting Paxil C Glaxosmithkline ORAL

Date:11/07/03ISR Number: 4229404-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0384061A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229405-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0384067A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharyngitis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229406-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0384072A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL

6 WK

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229407-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0384115A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
				Topamax	SS		
UNKNOWN							

Date:11/07/03ISR Number: 4229408-4Report Type:Periodic  
Age:38 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0384116A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgraphia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Heart Rate Increased		Prozac	C		
		Illiteracy		Trazodone	C		
		Performance Status Decreased					
		Vision Blurred					

Date:11/07/03ISR Number: 4229409-6Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0384117A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Tamoxifen	C		

Date:11/07/03ISR Number: 4229410-2Report Type:Periodic  
Age:67 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0384130A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Exfoliative		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4 YR	Dry Skin		Estrace	C		
UNKNOWN	1MG Unknown	Skin Exfoliation					

Date:11/07/03ISR Number: 4229411-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0384213A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229412-6Report Type:Periodic  
Age:28 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0384590A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Grand Mal Convulsion		Ortho Cyclen	C		
150MG Per day	5 DAY	Irritability					

Date:11/07/03ISR Number: 4229413-8Report Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0384599A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mood Swings		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1 MON						

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Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229414-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0384600A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cystitis Urinary Tract Infection		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229415-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0384603A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea Pharmaceutical Product Complaint		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229416-3Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0384607A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Three times per day 7 WK		Headache					
81MG per day		Hordeolum		Aspirin	C	Glaxosmithkline	
				Vitamin E	C		
				Calcium	C		
				Vitamin C	C	Glaxosmithkline	
				Centrum	C		

Date:11/07/03ISR Number: 4229417-5Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0384909A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Migraine		Xanax	C		



Date:11/07/03ISR Number: 4229418-7Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0384935A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Crying		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3 WK	Dry Mouth					
		Headache					
		Hypersensitivity					
		Weight Decreased					

Date:11/07/03ISR Number: 4229419-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0384936A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Per day		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229420-5Report Type:Periodic  
Age:37 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0384940A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Per day	5 DAY	Local Swelling		Wellbutrin	PS	Glaxosmithkline	ORAL
		Rash		Avapro	C		

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Freedom Of Information (FOI) Report

Mobic

C

Date:11/07/03ISR Number: 4229421-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0384995A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Twice per day		Accidental Overdose Asthenia Confusional State Dizziness Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229422-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0385077A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	5 DAY	Anger Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
				Prozac	C		

Date:11/07/03ISR Number: 4229423-0Report Type:Periodic  
Age:56 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0385078A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	1 MON	Anger Hyperhidrosis		Wellbutrin	PS	Glaxosmithkline	ORAL
20MG Per day	MON			Paxil	C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229424-2Report Type:Periodic  
Age:30 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0385146A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Burning Sensation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Pruritus					
per day	15 DAY	Urticaria					

Date:11/07/03ISR Number: 4229425-4Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385149A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Headache					
per day		Tinnitus					

Date:11/07/03ISR Number: 4229426-6Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385151A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Unknown	1 YR	Insomnia					
		Lethargy					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229428-XReport Type:Periodic  
 Age:54 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385155A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice per day	5 DAY	Anxiety Dizziness Tic Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229429-1Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385157A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229430-8Report Type:Periodic  
 Age:89 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385158A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown	2 DAY	Malaise Vomiting		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229431-XReport Type:Periodic  
 Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385252A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229432-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0385255A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hordeolum		Wellbutrin	PS	Glaxosmithkline	ORAL
3	WK						

Date:11/07/03ISR Number: 4229433-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0385261A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Depression					
per day	3 DAY	Dizziness		Hormone Replacement			
		Hyperhidrosis		Therapy	C		
		Insomnia					

Date:11/07/03ISR Number: 4229434-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0385277A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pain In Extremity		Wellbutrin	PS	Glaxosmithkline	ORAL
400MG per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229435-7Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385283A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day				Benadryl Asthma Medication	C C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229436-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385292A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229437-0Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385408A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Somnolence		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day		Vomiting		Lithium Methadone	C C	Glaxosmithkline Glaxosmithkline	

Date:11/07/03ISR Number: 4229445-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385419A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229446-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0385464A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	6	MON					

Date:11/07/03ISR Number: 4229447-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0385525A

Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ageusia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Dysgeusia					
per day	2	MON					
UNKNOWN		Parosmia		Prozac	C		
40MG Per day				Roxicodone	C		
5MG Twice per							
day				Oxycontin	C		
20MG Twice							
per day				Flexeril	C		
UNKNOWN							
10MG Twice							
per day							

Date:11/07/03ISR Number: 4229448-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0385532A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229449-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385537A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
				Depakote	SS		ORAL

Date:11/07/03ISR Number: 4229450-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385539A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Per day		Drug Screen False		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN		Positive		Aspirin	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229451-5Report Type:Periodic  
Age:67 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0385542A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	13 DAY	Constipation					
40MG Per day		Hangover		Coumadin	C	Glaxosmithkline	ORAL
		Sebaceous Glands		Zocor	C		ORAL
		Overactivity					
		Suicidal Ideation					
		Throat Tightness					

Date:11/07/03ISR Number: 4229453-9Report Type:Periodic  
Age:80 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385554A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG At		Blood Glucose Increased		Wellbutrin	PS	Glaxosmithkline	ORAL



night  
DAY  
Dry Mouth  
Fatigue

Celexa C  
Altace C  
Amaryl C  
Aciphex C  
Lipitor C  
Lopressor C

Date:11/07/03ISR Number: 4229455-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0385582A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Interaction		Flomax	SS		
		Urinary Incontinence					

UNKNOWN

Date:11/07/03ISR Number: 4229456-4Report Type:Periodic  
Age:35 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0385587A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Amnesia		Ativan	C		ORAL
		Aura					
1MG Per day		Convulsion					
		Grand Mal Convulsion					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229457-6Report Type:Periodic  
Age:15 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0385704A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Drug Screen Positive		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229458-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385738A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Twice per day		Pharmaceutical Product Complaint Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229459-XReport Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385746A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day 2 YR		Retinal Vascular Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229460-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385762A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash Throat Tightness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229461-8Report Type:Periodic  
Age:48 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0385787A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG In the morning		Blood Glucose Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
20MG Per day				Prozac	C		
15MG At night				Remeron	C		
				Asa	C	Glaxosmithkline	
				Mvi	C		
				Vioxx	C		

Date:11/07/03ISR Number: 4229462-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0385819A  
Age:30 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Twice per day	6 DAY	Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
				Diclofenac	C		

Date:11/07/03ISR Number: 4229463-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0385825A  
Age:10 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	1 MON	Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Eye Swelling					
		Fatigue		Amoxicillin	C	Glaxosmithkline	
		Oedema Peripheral		Trimex	C	Glaxosmithkline	
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229473-4Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385893A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown	YR	Pharmaceutical Product		Wellbutrin	PS	Glaxosmithkline	ORAL
		Complaint Somnolence		Fluoxetine	C		

Date:11/07/03ISR Number: 4229474-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385895A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Electrocardiogram Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229475-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385896A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 WK		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229476-XReport Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385899A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 WK	Hypoaesthesia Nausea Palpitations Throat Tightness Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229477-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0385950A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:11/07/03ISR Number: 4229478-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0385977A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Delirium		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged							

Date:11/07/03ISR Number: 4229479-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0385984A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
YR							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229480-1Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0385987A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1	MON					

Date:11/07/03ISR Number: 4229481-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386031A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Loss Of Consciousness		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1	MON					
		Road Traffic Accident					
				Elavil	C	Glaxosmithkline	
				Norvasc	C		
				Blood Pressure Medication	C		

Date:11/07/03ISR Number: 4229482-5Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386069A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Interaction		Wellbutrin	PS	Glaxosmithkline	ORAL
		Hot Flush					

Date:11/07/03ISR Number: 4229483-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386071A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229484-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0386081A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharmaceutical Product Complaint Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229485-0Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386100A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day				Zoloft	C		
				Depakote	C		
				Valium	C		

Date:11/07/03ISR Number: 4229486-2Report Type:Periodic  
Age:38 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0386162A

Outcome	PT
	Contusion Formication Joint Swelling Muscle Tightness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Pruritus Urticaria					
Dose	Duration		Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	18 DAY			Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229491-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0386170A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice per day		Hypokalaemia					

Date:11/07/03ISR Number: 4229492-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0386175A  
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day							

Date:11/07/03ISR Number: 4229493-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0386188A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - 150MG Twice Initial or Prolonged per day Other				Ritalin Unknown Medication	C C		



Date:11/07/03ISR Number: 4229494-1Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386189A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Drug Ineffective	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3	MON	Weight Increased				

Date:11/07/03ISR Number: 4229495-3Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386191A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day			Abdominal Pain Upper	Wellbutrin	PS	Glaxosmithkline	ORAL
			Chest Pain	Levothroid	C	Glaxosmithkline	
			Dizziness	Alcohol	C		
			Headache				
			Panic Attack				

Date:11/07/03ISR Number: 4229501-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386193A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Dizziness	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1	WK					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229502-8Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386230A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3 DAY			Xanax	C		

Date:11/07/03ISR Number: 4229503-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386256A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Liver Function Test Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229504-1Report Type:Periodic  
Age:42 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0386296A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	WK	Flushing Headache Hypertension Insomnia					

Date:11/07/03ISR Number: 4229505-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386326A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1 YR	Swelling					

Vasodilatation

Prozac  
Avalide

C  
C

Date:11/07/03ISR Number: 4229506-5Report Type:Periodic  
Age:10 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0386338A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 21 DAY		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Arthralgia		Penicillin	SS	Glaxosmithkline	
Other		Eye Swelling		Benadryl	C	Glaxosmithkline	
		Lethargy		Zyrtec	C	Glaxosmithkline	
		Oedema Peripheral		Zithromax	C		
		Serum Sickness		Cleocin	C		
		Urticaria					

Date:11/07/03ISR Number: 4229512-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386396A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharmaceutical Product Complaint Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229513-2Report Type:Periodic  
Age:44 YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0386425A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Levbid	C		
UNKNOWN				Allergy Shots	C		
UNKNOWN				Levsin Sl	C		
UNKNOWN	.125MG Per						
day							

Date:11/07/03ISR Number: 4229514-4Report Type:Periodic  
Age:43 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0386462A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bradyphrenia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Memory Impairment		Prozac	C		
per day							
10MG Per day							

Date:11/07/03ISR Number: 4229515-6Report Type:Periodic  
Age:48 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0386520A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Rash Generalised					
per day	2 WK	Serum Sickness					
		Urticaria					

Date:11/07/03ISR Number: 4229526-0Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386521A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blister		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Depression					
per day	2 WK	Insomnia					
		Pruritus					
		Rash					
		Rash Erythematous					

Date:11/07/03ISR Number: 4229527-2Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386522A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
25MG Per day	6 WK	Insomnia					

Date:11/07/03ISR Number: 4229528-4Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386524A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2 MON			Pravachol	C		
				Blood Pressure			
				Medication	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229529-6Report Type:Periodic  
Age:37 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0386525A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphemia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG In the							
morning							
2 YR				Paxil	C	Glaxosmithkline	ORAL
				Unknown Medication	C		

Date:11/07/03ISR Number: 4229530-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386526A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:11/07/03ISR Number: 4229531-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386533A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229532-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386558A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypotension		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229533-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386560A

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229534-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0393385A  
Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Irritability		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Nervousness					

Date:11/07/03ISR Number: 4229535-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0393392A  
Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Rash					
per day	16 DAY						

Date:11/07/03ISR Number: 4229536-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0393397A  
Age:74 YR Gender:Female I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Tremor					
per day	MON						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229537-5Report Type:Periodic  
Age:57 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393403A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscular Weakness		Wellbutrin	PS	Glaxosmithkline	ORAL
YR				Celexa	C		
YR							

Date:11/07/03ISR Number: 4229538-7Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0393409A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229539-9Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0393511A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229541-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393546A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229542-9Report Type:Periodic  
Age:17 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393552A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypotrichosis		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							



per day 4 MON Sleep Disorder  
 Weight Decreased Adderall Xr C  
 Orthotricyclen C

Date:11/07/03ISR Number: 4229543-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0393777A  
 Age:12 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Twice		Abdominal Distension		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	451 DAY	Swelling Face					
UNKNOWN	1MG Per day	Weight Increased		Risperdal	C		
UNKNOWN	150MG Twice	663 DAY		Lithium	C	Glaxosmithkline	
per day	532 DAY						
UNKNOWN	150MG Per day	991 DAY		Zoloft	C		
UNKNOWN	.5MG Per day	3 YR		Clozapine	C		

Date:11/07/03ISR Number: 4229544-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0393797A  
 Age:9 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
85 DAY		Cyanosis					
Hospitalization -		Musculoskeletal Stiffness					
Initial or Prolonged		Opisthotonus					
Other							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229545-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393805A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229546-6Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393817A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK	Chest Wall Pain		Birth Control Pills	C		
				Chondroitin	C		

Date:11/07/03ISR Number: 4229547-8Report Type:Periodic  
 Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393842A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK						

Date:11/07/03ISR Number: 4229548-XReport Type:Periodic  
 Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393847A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	DAY	Bladder Pain		Celexa	C		
		Disturbance In Attention		Remeron	C		
		Feeling Abnormal		Vitamins	C		
		Insomnia		Ultram	C		
		Nightmare		Nexium	C		

Date:11/07/03ISR Number: 4229553-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0393859A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypotrichosis		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/07/03ISR Number: 4229554-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0393861A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1 WK						

Date:11/07/03ISR Number: 4229558-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0393863A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction Priapism		Wellbutrin Trazodone	PS C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229559-4Report Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393886A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2	MON					

Date:11/07/03ISR Number: 4229560-0Report Type:Periodic  
Age:51 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0393971A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased		Wellbutrin Metformin	PS C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229561-2Report Type:Periodic  
Age:33 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0393983A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bruxism		Wellbutrin	PS	Glaxosmithkline	ORAL
1 YR		Drug Ineffective Nervousness Toothache					

Date:11/07/03ISR Number: 4229562-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393989A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day	33	DAY					

Date:11/07/03ISR Number: 4229563-6Report Type:Periodic  
Age:69 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394031A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
225MG per day	6 WK	Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
.25MG Twice		Decreased Appetite		Klonopin	C		
per day		Dry Mouth					
		Headache		Tylenol	C	Glaxosmithkline	
		Nausea					
		Nervousness					
		Palpitations					
		Sleep Disorder					

Date:11/07/03ISR Number: 4229564-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0394056A  
Age:46 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3 YR			Clinoril	C		
				Atacand	C		
				Plaquenil	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229565-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394063A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice		Erythema		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	4 MON	Rash					
		Urticaria					

Date:11/07/03ISR Number: 4229566-1Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394067A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229567-3Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0394204A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Twice		Bronchospasm		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Depression					
		Hypersensitivity					

Date:11/07/03ISR Number: 4229568-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394264A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 WK	Rash					

Date:11/07/03ISR Number: 4229569-7Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394270A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Accidental Overdose	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
			Headache				
per day	2	MON		Advair	C	Glaxosmithkline	
RESPIRATORY							
(INHALATION)							
				Vicodin	C		
				Nebulizer	C		
				Nicotine Patch	C	Glaxosmithkline	
INTRADERMAL							

Date:11/07/03ISR Number: 4229570-3Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394274A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Drug Ineffective	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
			Therapeutic Response				
per day			Unexpected	Paxil	C	Glaxosmithkline	
				Lipitor	C		
				Vioxx	C		
				Hrt	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229571-5Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394314A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Flatulence		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	6	MON					
		Nausea					
		Vomiting		Celebrex	C		
				Prilosec	C		

Date:11/07/03ISR Number: 4229572-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394327A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Amenorrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1	WK					

Date:11/07/03ISR Number: 4229573-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394329A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2	MON		Wellbutrin	PS	Glaxosmithkline	ORAL
		Blood Potassium Increased					
				Serzone	C		
				Thyroid Medication	C		

Date:11/07/03ISR Number: 4229574-0Report Type:Periodic  
Age:28 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0394436A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2	WK					
		Rash Generalised					



Date:11/07/03ISR Number: 4229575-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0394439A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flushing Hyperhidrosis Premature Menopause		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229576-4Report Type:Periodic  
Age: Gender:I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394531A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229577-6Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394639A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Insomnia					
per day				Celexa	C		
40MG In the							
morning							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229578-8Report Type:Periodic  
Age:37 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0394644A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	9 MON	Anxiety Diarrhoea Flushing Night Sweats		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229579-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394708A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Three times per day		Erectile Dysfunction Hypersomnia Sexual Dysfunction		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229580-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394730A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229581-8Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394818A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Per day		Amenorrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
				Prempro	C		
				Synthroid	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229582-XReport Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394819A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amenorrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day	4	MON					

Date:11/07/03ISR Number: 4229583-1Report Type:Periodic  
Age:25 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394821A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Loss Of Consciousness		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day		Syncope		Valproic Acid	C		
		Tachycardia		Alka Seltzer	C		

Date:11/07/03ISR Number: 4229584-3Report Type:Periodic  
Age:50 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0394830A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	5	DAY		Aciphex	C		
		Insomnia		Flexeril	C		
		Pharyngolaryngeal Pain					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229585-5Report Type:Periodic  
Age:28 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0394866A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	46 DAY	Convulsion Grand Mal Convulsion Tonic Clonic Movements					

Date:11/07/03ISR Number: 4229586-7Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394872A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Euphoric Mood		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1 WK	Gastric Disorder Palpitations					

Date:11/07/03ISR Number: 4229587-9Report Type:Periodic  
Age:29 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0394992A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Fall					
per day	12 MON	Feeling Abnormal Loss Of Consciousness Road Traffic Accident Skin Laceration		Zoloft	C		

Date:11/07/03ISR Number: 4229588-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395022A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Convulsion Wellbutrin PS Glaxosmithkline  
UNKNOWN

Date:11/07/03ISR Number: 4229589-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395040A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Hair Metal Test					
		Parosmia					
		Skin Odour Abnormal					

Date:11/07/03ISR Number: 4229590-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395159A  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Myalgia					
per day	2 WK						

Date:11/07/03ISR Number: 4229591-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395171A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day		Migraine		Imitrex	SS	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229592-2Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395176A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Twice	Delusion		Wellbutrin	PS	Glaxosmithkline	ORAL
	per day	Paranoia					

Date:11/07/03ISR Number: 4229593-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395177A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyskinesia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Prozac	C		

Date:11/07/03ISR Number: 4229594-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395213A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	1 DAY	Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229595-8Report Type:Periodic  
Age:12 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395271A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	100MG Twice	Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
	per day	Anger					
	1 MON	Headache					

Date:11/07/03ISR Number: 4229596-XReport Type:Periodic  
Age:22 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395380A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1	MON					
		Obsessive-Compulsive Disorder		Prozac	C		
				Zyprexa	C		

Date:11/07/03ISR Number: 4229597-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395382A  
 Age:54 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Blood Pressure Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	21	DAY					
		Confusional State					
		Impatience		Atenolol	C		
		Irritability		Aleve	C		
		Memory Impairment					
		Personality Change					
		Stress					

Date:11/07/03ISR Number: 4229610-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395387A  
 Age:85 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Hallucination		Wellbutrin	PS	Glaxosmithkline	ORAL
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229611-3Report Type:Periodic  
Age:84 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395411A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	2 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
		Agitation					
		Insomnia					
		Nausea					

Date:11/07/03ISR Number: 4229612-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395419A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Rash					

Date:11/07/03ISR Number: 4229613-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395421A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion		Elavil	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229614-9Report Type:Periodic  
Age:40 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0395423A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	200MG Twice			Wellbutrin	PS	Glaxosmithkline	ORAL
	per day	2 YR		Glucophage	C		
		Anxiety					
		Drug Ineffective					
		Insomnia					
		Irritability					



Date:11/07/03ISR Number: 4229615-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395437A  
Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Clumsiness		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Disturbance In Attention					
per day	6	Feeling Abnormal					
		DAY					

Date:11/07/03ISR Number: 4229616-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395466A  
Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eye Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Vision Blurred					
per day	2						
		MON					

Date:11/07/03ISR Number: 4229617-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395482A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	7						
		DAY					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229618-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395487A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Myalgia Serum Sickness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229619-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395503A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229620-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395510A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
300MG per day		Feeling Abnormal Sexual Dysfunction		Comtrex (Daytime)	SS		

Date:11/07/03ISR Number: 4229626-5Report Type:Periodic  
Age:46 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0395658A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Urticaria		Premarin	C		
150MG Per day 1 DAY				Acyclovir	C	Glaxosmithkline	
				Celebrex	C		

Date:11/07/03ISR Number: 4229627-7Report Type:Periodic  
Age:54 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0395664A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
3 WK				Remeron	C		

Date:11/07/03ISR Number: 4229628-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395682A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Zyprexa	C		
				Unknown Medication	C		

Date:11/07/03ISR Number: 4229629-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395690A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229630-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395728A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229631-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0395729A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Twice per day	2 MON	Burning Sensation Pruritus Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229632-0Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395831A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Anxiety Heart Rate Increased		Wellbutrin Zebeta	PS C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229633-2Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395835A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
450MG per day	3 MON	Alopecia Hypotrichosis		Wellbutrin Oxycontin Nortriptyline	PS C C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229634-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395875A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229635-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395878A  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nightmare		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Prozac	C		

Date:11/07/03ISR Number: 4229636-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395880A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229637-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395884A  
Age:21 YR Gender:Female I/FU:I

Outcome	PT
Other	Eczema Erythema Pallor

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Stevens-Johnson Syndrome

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	34 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229647-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395888A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Unknown	5	MON	Energy Increased	Wellbutrin	PS	Glaxosmithkline	ORAL
100U Per day	1	YR	Hyperhidrosis	Zoloft	SS		ORAL
600U Twice per day		YR	Weight Increased	Trileptal	C		ORAL
5MG Per day		YR		Lopressor	C		ORAL
20U Per day		YR		Furosemide	C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229648-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395891A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Stool Analysis Abnormal	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229649-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0396026A  
Age:90 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Unknown			Diplopia	Wellbutrin	PS	Glaxosmithkline	ORAL

Nausea  
Tremor  
Vision Blurred

Concurrent  
Medications

C

Date:11/07/03ISR Number: 4229650-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396027A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Insomnia Irritability		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229651-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396034A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depersonalisation Dissociation Feeling Abnormal Feeling Cold Flight Of Ideas Headache Insomnia Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

150MG Twice  
per day

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229652-6Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396039A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	6 DAY	Rash	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229653-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396040A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Pruritus Rash	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229655-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396044A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Twice per day		Dry Mouth Dyspnoea Nasal Congestion Nasal Dryness	Wellbutrin Bextra Norco Trazodone Singulair Zyrtec Prevacid	PS C C C C C C	Glaxosmithkline Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229656-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396169A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Twice per day		Blood Glucose Increased Drug Interaction	Wellbutrin	PS	Glaxosmithkline	ORAL



UNKNOWN Tinnitus Avandamet SS Glaxosmithkline  
UNKNOWN Insulin SS

Date:11/07/03ISR Number: 4229657-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0396265A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Depression					
per day	1 YR	Drug Ineffective		Prozac	C		
20MG Unknown							

Date:11/07/03ISR Number: 4229658-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0396267A  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disturbance In Attention		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Dizziness					
per day	11 DAY	Insomnia					
		Nausea					
		Tinnitus					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229665-4Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396268A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Bleeding Time Prolonged	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	5	MON	Menopausal Symptoms				
			Menorrhagia	Klonopin	C		

Date:11/07/03ISR Number: 4229666-6Report Type:Periodic  
Age:77 YR Gender:I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0396289A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day			Atrial Fibrillation	Wellbutrin	PS	Glaxosmithkline	ORAL
				Ambien	C		
				Clonopin	C		

Date:11/07/03ISR Number: 4229667-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396290A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	14	DAY	Blood Pressure Increased	Wellbutrin	PS	Glaxosmithkline	ORAL
21MG Unknown			Choking	Nicotine	C	Glaxosmithkline	NASAL
			Cough				
			Drug Ineffective				
			Energy Increased				
			Headache				
			Libido Increased				
			Malaise				
			Mental Impairment				
			Therapeutic Response				
			Unexpected				
			Weight Increased				

Date:11/07/03ISR Number: 4229668-XReport Type:Periodic  
Age:29 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0396319A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blindness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Blindness Transient		Zoloft	C		ORAL
100MG Per day	MON	Muscle Contractions Involuntary Muscle Twitching Vision Blurred Vomiting					

Date:11/07/03ISR Number: 4229669-1Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396322A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 YR			Prozac	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229670-8Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0396463A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Irritability		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229671-XReport Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396473A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Intentional Misuse Psychotic Disorder Suicide Attempt		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229672-1Report Type:Periodic  
Age:49 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396618A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229673-3Report Type:Periodic  
Age:42 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0396652A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day	8 DAY	Emotional Distress Nausea		Thyroid Medication Birth Control	C C		

Date:11/07/03ISR Number: 4229674-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396653A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Electroencephalogram Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229675-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0396655A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG In the morning		Sleep Apnoea Syndrome		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229676-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0396659A  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Extrasystoles Musculoskeletal Stiffness		Wellbutrin	PS	Glaxosmithkline	ORAL
				Lithium	C	Glaxosmithkline	
				Nitroglycerin	C	Glaxosmithkline	
				Plavix	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229678-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0396718A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229679-4Report Type:Periodic  
Age:12 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0396731A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complex Partial Seizures		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Three		Convulsion					
times per day		Tongue Paralysis		Risperidal	C		ORAL

Date:11/07/03ISR Number: 4229680-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396739A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
Initial or Prolonged				Lexapro	C		
per day	MON						

Date:11/07/03ISR Number: 4229681-2Report Type:Periodic  
Age:41 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0396760A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Twice		Pharmaceutical Product	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 MON	Complaint	Professional				
40MG Per day		Retching		Prozac	C		ORAL

10MG At night				Ambien	C		ORAL
				Provigil	C		ORAL
				Gabatril	C		ORAL

Date:11/07/03ISR Number: 4229682-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0396771A  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Trigeminal Neuralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Medroxyprogesterone	C		

Date:11/07/03ISR Number: 4229683-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0396784A  
 Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day 1 DAY		Swelling Face		Vitamins	C		
				Ambien	C		

Date:11/07/03ISR Number: 4229684-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0397023A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229685-XReport Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0397029A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Heart Rate Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/07/03ISR Number: 4229686-1Report Type:Periodic  
 Age:40 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0397033A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated		Wellbutrin	PS	Glaxosmithkline	ORAL
Life-Threatening							
150MG Per day 10 DAY							
Other		Hypertension					

Date:11/07/03ISR Number: 4229687-3Report Type:Periodic  
 Age:53 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397040A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort		Wellbutrin	PS	Glaxosmithkline	ORAL
1 YR							
UNKNOWN	45MG Per day	Feeling Abnormal		Remeron	C		
		Insomnia					
		Palpitations					
		Restlessness					
		Tinnitus					

Date:11/07/03ISR Number: 4229688-5Report Type:Periodic  
 Age:76 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397041A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG In the							



morning 2 YR Drug Ineffective  
 Pharmaceutical Product  
 Complaint  
 Lorazepam C  
 Calcium C  
 Vitamin C C Glaxosmithkline  
 500MG Unknown  
 Norvasc C

Date:11/07/03ISR Number: 4229690-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0397116A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229691-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0397132A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Snoring		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Nicoderm	C	Glaxosmithkline	
per day							
INTRADERMAL							

Date:11/07/03ISR Number: 4229692-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0397133A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229693-9Report Type:Periodic  
Age:32 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0397143A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Tinnitus					

Date:11/07/03ISR Number: 4229694-0Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397148A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3	WK					

Date:11/07/03ISR Number: 4229695-2Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0397150A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229696-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397152A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accidental Overdose		Wellbutrin	PS	Glaxosmithkline	ORAL
		Disturbance In Attention		Prozac	C		
		Dysphasia		Xanax	C		
		Feeling Abnormal					
		Heart Rate Increased					
		Incoherent					
		Panic Reaction					

Date:11/07/03ISR Number: 4229697-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0397154A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229698-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0397161A  
Age:34 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Joint Swelling					
per day	18 DAY	Rash					

Date:11/07/03ISR Number: 4229699-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0397163A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bruxism		Wellbutrin	PS	Glaxosmithkline	ORAL
400MG Per day	3 MON	Muscle Tightness		Perphenazine	C	Glaxosmithkline	
				Unknown Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229700-3Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397168A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	7 WK	Chest Pain Flushing Heart Rate Increased Increased Appetite Insomnia Nausea		Wellbutrin  Oral Contraceptive	PS  C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229701-5Report Type:Periodic  
Age:32 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397176A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day 30MG At night 75MG Per day	4 MON	Hypersomnia Somnolence		Wellbutrin Remeron Effexor	PS C C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229702-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397200A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN UNKNOWN	225MG Unknown 175MG Unknown	Pruritus Urticaria		Wellbutrin Effexor Xr Unithroid	PS C C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229710-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397211A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day	28 DAY	Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
			Dry Mouth		Nexium	C		
			Rash Erythematous		Actigall	C		
			Sinus Congestion		Sam E	C		
			Urine Output Decreased		Tylenol Pm	C		
UNKNOWN					Trazodone	C		
UNKNOWN					Librium	C		

Date:11/07/03ISR Number: 4229711-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0397347A  
Age:65 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	3	DAY	Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
			Dyspnoea		Baby Aspirin	C	Glaxosmithkline	
	50MG Twice		Heart Rate Increased		Cataflam	C		
		per day						

Date:11/07/03ISR Number: 4229712-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0397352A  
Age:35 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	200MG	In the morning	Yawning		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN	50MG	Per day			Zoloft	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229713-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397375A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Drug Effect Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
per day							

Date:11/07/03ISR Number: 4229714-3Report Type:Periodic  
Age:36 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0397382A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100VA Twice		Neuralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	36 DAY	Paraesthesia		Lipitor	C		ORAL

Date:11/07/03ISR Number: 4229715-5Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0397383A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229716-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397515A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Per day	1 YR	Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229718-0Report Type:Periodic  
Age:29 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0397522A

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	8 DAY						

Date:11/07/03ISR Number: 4229719-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0397547A  
 Age: Gender:Female I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Acne		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229720-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0397552A  
 Age:57 YR Gender:Male I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Life-Threatening		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
350MG per day	6 YR						
Other		Grand Mal Convulsion		Nortriptyline	C		ORAL
50MG Per day							
				Aricept	C		ORAL
10MG Per day							
				Buspar	C		ORAL
30MG Unknown							

Date:11/07/03ISR Number: 4229721-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0397568A  
 Age:75 YR Gender:Male I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Swelling		Wellbutrin	PS	Glaxosmithkline	ORAL

100MG Twice

per day

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229722-2Report Type:Periodic  
Age:54 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397572A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Hyperhidrosis		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	15 DAY			Ultram	C		
				Albuterol	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229723-4Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397573A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	5 DAY	Insomnia		Ambien	C		
				Unknown Medication	C		

Date:11/07/03ISR Number: 4229724-6Report Type:Periodic  
Age:53 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0397578A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 DAY	Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
		Fall		Birth Control Pills	C		
				Vicodin	C		
				Valium	C		

Date:11/07/03ISR Number: 4229725-8Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0397628A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anger		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:11/07/03ISR Number: 4229726-XReport Type:Periodic  
 Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397633A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3	WK		Vitamins	C		
				Lipitor	C		
				Prevacid	C		

Date:11/07/03ISR Number: 4229727-1Report Type:Periodic  
 Age:25 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397634A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dreamy State		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day	10	DAY					
25MG At night		Impaired Driving Ability		Paxil	SS	Glaxosmithkline	
		Somnolence					
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229728-3Report Type:Periodic  
Age:18 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397638A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin Sr	PS	Glaxosmithkline	
Other							
UNKNOWN	150MG	Per day	MON				

Date:11/07/03ISR Number: 4229729-5Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0397799A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Twice						
per day							

Date:11/07/03ISR Number: 4229730-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397817A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Twice						
per day	6	DAY					
		Abnormal Dreams					
		Anxiety		Vitamins	C		
		Hypoaesthesia		Flax Seed Oil	C		
		Neck Pain					
		Nervousness					
		Palpitations					

Date:11/07/03ISR Number: 4229731-3Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397826A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG	Per day						

Date:11/07/03ISR Number: 4229732-5Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397837A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Dry Mouth		Astelin	C		NASAL
		Feeling Jittery		Vitamins	C		
		Heart Rate Increased					
		Hyperhidrosis					
		Menstrual Disorder					

Date:11/07/03ISR Number: 4229733-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397984A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eczema Weeping		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Pruritus					
per day	2 WK						

Date:11/07/03ISR Number: 4229735-0Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397992A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -							
150MG Twice							
Initial or Prolonged							
per day	16 DAY			Oscal	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

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Date:11/07/03ISR Number: 4229736-2Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0398041A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Intentional Misuse		Wellbutrin	PS	Glaxosmithkline	NASAL

Date:11/07/03ISR Number: 4229737-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398042A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	18 MON	Night Sweats		Prevacid	C		
		Weight Decreased		Lodine	C		
400MG Per day				Zanaflex	C		

Date:11/07/03ISR Number: 4229738-6Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398172A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Three							
Initial or Prolonged		Dry Mouth					
times per day	25 DAY	Grand Mal Convulsion					
		Head Injury					
		Incontinence					

Date:11/07/03ISR Number: 4229739-8Report Type:Periodic  
Age:19 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398183A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Convulsion Wellbutrin PS Glaxosmithkline  
300MG Per day Lexapro C

Date:11/07/03ISR Number: 4229740-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0398211A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Unknown						

Date:11/07/03ISR Number: 4229741-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0398316A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Unknown 3 WK	Dry Mouth Headache Insomnia Stress					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229742-8Report Type:Periodic  
Age:76 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398332A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
3	MON			Oxazepam	C		

Date:11/07/03ISR Number: 4229743-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398333A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
				Strattera	SS		
UNKNOWN							

Date:11/07/03ISR Number: 4229744-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398363A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Syncope		Wellbutrin	PS	Glaxosmithkline	ORAL
				Effexor	C		
UNKNOWN							

Date:11/07/03ISR Number: 4229745-3Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398364A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Halo Vision		Lamictal	C	Glaxosmithkline	
		Insomnia		Prozac	C		

Date:11/07/03ISR Number: 4229746-5Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0398385A

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Bezoar		Wellbutrin	PS	Glaxosmithkline	ORAL
22.5G Single							
dose		Overdose					

Date:11/07/03ISR Number: 4229747-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0398387A  
 Age: Gender: I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Muscle Twitching		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229748-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0398454A  
 Age: Gender:Male I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Gingival Pain Tooth Disorder		Wellbutrin Crystal Meth	PS C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229749-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0398459A  
 Age:30 YR Gender:Female I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Unknown	1 WK			Wellbutrin	SS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229750-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398479A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Syncope		Wellbutrin Effexor	PS C	Glaxosmithkline	ORAL
UNKNOWN							

Date:11/07/03ISR Number: 4229751-9Report Type:Periodic  
Age:2 MON Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0398538A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arrhythmia Neonatal Drug Exposure Via Breast Milk		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229752-0Report Type:Periodic  
Age:15 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398541A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229753-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398542A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Tachycardia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229754-4Report Type:Periodic  
Age:46 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0398614A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



150MG Twice	Liver Function Test	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	Abnormal MON	Professional				
			Alcohol Tarka	SS C		ORAL ORAL
2 WK			Pepcid Zantac Tylenol Clonidine	C C C C	Glaxosmithkline Glaxosmithkline	ORAL
.2U At night			Norvasc	C		ORAL

Date:11/07/03ISR Number: 4229756-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0398664A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229757-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0398856A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression Suicidal Incoherent		Wellbutrin Ambien Prinzide	PS C C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229758-1Report Type:Periodic  
Age:38 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0398858A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Dizziness	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3 WK	Drug Interaction	Professional				
UNKNOWN				Sudafed	SS	Glaxosmithkline	

Date:11/07/03ISR Number: 4229759-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398873A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 WK	Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
				Unknown Medication	C		

Date:11/07/03ISR Number: 4229760-XReport Type:Periodic  
Age:37 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398928A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Blood Testosterone		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1 YR	Decreased		Prozac	C		

Date:11/07/03ISR Number: 4229761-1Report Type:Periodic  
Age:18 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398949A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day	3 WK	Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
				Adderall	C		
				Zoloft	C		

Date:11/07/03ISR Number: 4229762-3Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399030A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Medication Error	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	6	MON	Migraine				
				Topamax	C		
				Klonopin	C		

Date:11/07/03ISR Number: 4229763-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399031A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Unknown			Pharmaceutical Product	Wellbutrin	PS	Glaxosmithkline	ORAL
			Complaint				
			Stool Analysis Abnormal				

Date:11/07/03ISR Number: 4229764-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399033A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Tic	Wellbutrin	PS	Glaxosmithkline	ORAL
				Ssri	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229766-0Report Type:Periodic  
 Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399077A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Constipation					
per day	4	MON		Tiazac	C	Glaxosmithkline	
		Insomnia		Activella	C		
		Tremor		Colace	C		

Date:11/07/03ISR Number: 4229767-2Report Type:Periodic  
 Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399079A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Hypoacusis					
400MG See		Tinnitus		Prozac	C		
dosage text	4	MON		Klonopin	C		
				Atarax	C		
				Zanaflex	C		

Date:11/07/03ISR Number: 4229768-4Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399080A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Ineffective					
per day				Ranitidine	C	Glaxosmithkline	
				Lipitor	C		
				Prinivil	C		
				Atenolol	C		
				Plavix	C		
				Celexa	C		

Date:11/07/03ISR Number: 4229769-6Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399095A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Twice			Abdominal Discomfort	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2	YR	Nausea				
			Vomiting	Creon	C		
				Duragesic	C		
				Klonopin	C		

Date:11/07/03ISR Number: 4229770-2Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399271A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Disturbance In Attention	Wellbutrin	PS	Glaxosmithkline	ORAL
			Dizziness	Sam E	C		
				Valerian Root	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229771-4Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0399279A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety Autonomic Nervous System Imbalance Drug Abuser Heart Rate Increased Intentional Misuse		Wellbutrin	PS	Glaxosmithkline	NASAL

Date:11/07/03ISR Number: 4229772-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399299A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	9 DAY	Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229773-8Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0399462A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Malaise		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229774-XReport Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399594A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice per day	2 MON	Drug Ineffective Insomnia Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229775-1Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0399605A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229776-3Report Type:Periodic  
Age:71 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399640A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Anxiety					
per day	7 DAY	Dry Mouth		Combivent	C		
RESPIRATORY		Insomnia					
(INHALATION)		Tremor					

Date:11/07/03ISR Number: 4229777-5Report Type:Periodic  
Age:52 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0399647A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cold Sweat		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Unknown	DAY	Dizziness		Zocor	C		
		Nervousness		Temazepam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229778-7Report Type:Periodic  
 Age:29 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399824A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin	PS	Glaxosmithkline	ORAL
350MG per day	5 MON	Chills		Allegra	C		
UNKNOWN		Dizziness		Benadryl	C	Glaxosmithkline	
25MG Unknown		Nausea					
		Sinusitis					
		Tremor					
		Vomiting					

Date:11/07/03ISR Number: 4229779-9Report Type:Periodic  
 Age:21 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399827A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Semen Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice				Adderall	C		
per day	2 YR						

Date:11/07/03ISR Number: 4229780-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399828A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Cytomel	C	Glaxosmithkline	
per day							

Date:11/07/03ISR Number: 4229781-7Report Type:Periodic  
 Age:59 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399835A



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day	MON	Headache		Zoloft	C		
100MG Per day							

Date:11/07/03ISR Number: 4229782-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0399872A  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cold Sweat		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Paraesthesia					
per day	11 DAY	Peripheral Coldness					

Date:11/07/03ISR Number: 4229783-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0399947A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229784-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0399967A  
 Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Hot		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Pruritus Generalised					
		Rash					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229785-4Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399969A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Unknown			Stool Analysis Abnormal	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229786-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399971A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day			Insomnia	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229787-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399972A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 WK			Pruritus	Wellbutrin	PS	Glaxosmithkline	ORAL
			Rash Urticaria	Zoloft	C		

Date:11/07/03ISR Number: 4229790-8Report Type:Periodic  
 Age:70 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0056952A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	11 DAY		Anxiety	Wellbutrin	PS	Glaxosmithkline	ORAL
			Bruxism	Elavil	C	Glaxosmithkline	
			Feeling Jittery	Zoloft	C		
			Nervousness	Paxil	C	Glaxosmithkline	
				Pepcid	C		
				Antacids	C		
				Tylenol	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229802-1Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0365399A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Feeling Hot					

Date:11/07/03ISR Number: 4229803-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0371708A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2 WK			Celexa	C		

Date:11/07/03ISR Number: 4229804-5Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0381524A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	1 YR			Oral Contraceptive	C		ORAL
				Asthma Medication	C		

UNKNOWN

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229806-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386565A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229807-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386583A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice per day							

Date:11/07/03ISR Number: 4229808-2Report Type:Periodic  
Age:54 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386605A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cough		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day							

Date:11/07/03ISR Number: 4229809-4Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386640A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
2 YR							
		Dizziness		Neurontin	C		
				Dilantin	C		
				Phenobarb	C		
				Triamterene	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229810-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0386654A  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Three times per day 6 YR							

Date:11/07/03ISR Number: 4229811-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0386664A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day WK							
		Anxiety					
		Drug Ineffective		Depakote	C		
		Hyperhidrosis					

Date:11/07/03ISR Number: 4229812-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0386699A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Estrogenic Effect		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229813-6Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386719A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	6 WK	Dysgeusia					
				Desogen	C		
				Mvi	C		
INTRAVENOUS							

Date:11/07/03ISR Number: 4229814-8Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0386720A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 WK	Aphthous Stomatitis					
		Dry Mouth		Androgel	C		
		Erythema					
		Pruritus					
		Rash					
		Swelling Face					
		Tongue Exfoliation					
		Weight Decreased					

Date:11/07/03ISR Number: 4229815-XReport Type:Periodic  
Age:47 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0386757A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Pancreatitis		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
Initial or Prolonged							
per day							
				Eskalith	C	Glaxosmithkline	
450MG Twice							
per day							

10MG Twice			Inderal		C		ORAL
per day							
20MG Per day			Geodon		C		ORAL

Date:11/07/03ISR Number: 4229816-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0386761A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Withdrawal Syndrome		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/07/03ISR Number: 4229817-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0386765A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Thrombocytopenia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229818-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0386766A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day		Drug Withdrawal Syndrome					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229819-7Report Type:Periodic  
Age:54 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0386768A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 4500MG Single Initial or Prolonged dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
		Convulsion					
		Hypertension					
		Intentional Misuse					
		Overdose					
		Rhabdomyolysis					
		Tachycardia					

Date:11/07/03ISR Number: 4229820-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386770A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400MG per day		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Cognitive Disorder					

Date:11/07/03ISR Number: 4229821-5Report Type:Periodic  
Age:16 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386903A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
		Anger					
		Convulsion					
		Dry Mouth					
		Feeling Jittery					
		Nausea					
		Overdose					
		Vomiting					

Date:11/07/03ISR Number: 4229822-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0386939A



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Lipitor	C		

Date:11/07/03ISR Number: 4229823-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0386947A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL
1 WK							

Date:11/07/03ISR Number: 4229824-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0387013A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice		Food Craving					
per day		Headache					
		Insomnia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229826-4Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387107A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4	MON					

Date:11/07/03ISR Number: 4229827-6Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387119A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2	MON					
		Drug Ineffective		Vioxx	C		
				Anaplex	C		

Date:11/07/03ISR Number: 4229828-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387122A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Myalgia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day		YR					

Date:11/07/03ISR Number: 4229829-XReport Type:Periodic  
Age:27 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387123A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		YR					
		Pharmaceutical Product		Unspecified			
		Complaint		Medication	C		
TOPICAL							

Date:11/07/03ISR Number: 4229830-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0387212A  
Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Nausea					
per day	2 DAY			Lexapro	C		

Date:11/07/03ISR Number: 4229831-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0387241A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accidental Overdose	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
600MG Unknown	1 MON	Aggression	Professional	Alcohol	SS		ORAL
		Alcohol Use		Prozac	C		
		Convulsion		Ambien	C		
		Medication Error					
		Postictal State					

Date:11/07/03ISR Number: 4229832-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0387252A  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	8 MON			Synthroid	C	Glaxosmithkline	
1MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229833-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387309A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	MON			Ativan	C		

Date:11/07/03ISR Number: 4229834-3Report Type:Periodic  
Age:54 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0387322A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Tinnitus					
per day		Urticaria		Altace	C		

Date:11/07/03ISR Number: 4229835-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387340A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Paraesthesia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229836-7Report Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387373A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharmaceutical Product		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Complaint					
per day		Therapeutic Response Decreased		Protonix	C		
				Reglan	C	Glaxosmithkline	
				Pravachol	C		

Date:11/07/03ISR Number: 4229837-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387376A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229838-0Report Type:Periodic  
Age:23 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387389A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disturbance In Attention		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Increased Appetite					
		Lethargy					

Date:11/07/03ISR Number: 4229839-2Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387425A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chromaturia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Faeces Discoloured		Nicotine Polacrilex	SS	Glaxosmithkline	ORAL
4 DAY				Aspirin	C	Glaxosmithkline	
				Vitamins	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229840-9Report Type:Periodic  
Age:53 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0387468A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Anxiety Drug Ineffective Insomnia Weight Decreased		Wellbutrin  Ambien	PS  C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229841-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387492A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety Obsessive-Compulsive Disorder Sleep Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229842-2Report Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387563A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG See dosage text	3 WK	Feeling Hot Flushing		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229843-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387585A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	3 WK	Dry Mouth Tongue Blistering		Wellbutrin	PS	Glaxosmithkline	ORAL

Androgel	C	
Baby Aspirin	C	Glaxosmithkline
Unknown Medication	C	

Date:11/07/03ISR Number: 4229844-6Report Type:Periodic  
 Age:42 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387622A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypotrichosis		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day				Avonex	C		

Date:11/07/03ISR Number: 4229845-8Report Type:Periodic  
 Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387645A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Crying		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK	Emotional Disorder		Pravachol	C		
				Prempro	C		

Date:11/07/03ISR Number: 4229846-XReport Type:Periodic  
 Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387646A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1 MON						

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vitamin C  
Sonata C

Date:11/07/03ISR Number: 4229847-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387651A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229848-3Report Type:Periodic  
Age:41 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387673A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 WK		Anorexia Crying Depression Difficulty In Walking Dry Mouth Impatience Insomnia Irritability Muscle Fatigue Muscle Tightness Nervousness Somnolence Tinnitus Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229849-5Report Type:Periodic  
Age:40 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0387678A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice per day	18 MON	Disturbance In Attention Grand Mal Convulsion Memory Impairment		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:11/07/03ISR Number: 4229850-1Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387683A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG			Drug Withdrawal Syndrome	Wellbutrin	PS	Glaxosmithkline	ORAL
Alternate			Headache				
days	YR		Psychomotor Hyperactivity				
			Tension				

Date:11/07/03ISR Number: 4229858-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387695A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Urine Odour Abnormal	Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229859-8Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0387702A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	4 WK	Altered Visual Depth Perception		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229860-4Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0387730A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		Abdominal Pain Upper Agitation		Bupropion	PS	Glaxosmithkline	ORAL
UNKNOWN		Anxiety		Depakote	C		
UNKNOWN		Anxiety		Geodon	C		
UNKNOWN		Confusional State		Prevacid	C		
UNKNOWN		Delusion		Lexapro	C		
UNKNOWN		Disturbance In Attention		Fluoxetine	C		
UNKNOWN		Dizziness		Nexium	C		
		Hallucination					
		Hypertension					
		Nausea					
		Paranoia					
		Psychotic Disorder					
		Tremor					

Date:11/07/03ISR Number: 4229861-6Report Type:Periodic  
 Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387754A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Contusion Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229862-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387797A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin Unknown	PS C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229863-XReport Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0387807A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Twice		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3 WK	Nausea					

Date:11/07/03ISR Number: 4229864-1Report Type:Periodic  
Age:51 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0387813A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
Hospitalization -		Eye Rolling					
per day							
Initial or Prolonged		Tonic Clonic Movements					
Other							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229865-3Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387821A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
3	DAY						

Date:11/07/03ISR Number: 4229866-5Report Type:Periodic  
Age:78 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387954A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cough		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	DAY	Tinnitus		Vioxx	C		
				Atacand	C		
				Motrin	C	Glaxosmithkline	
				Synthroid	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229867-7Report Type:Periodic  
Age:45 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0387972A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -		Incoherent		Celexa	C		ORAL
200MG Per day 2	WK						
Initial or Prolonged							
Other							

Date:11/07/03ISR Number: 4229868-9Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387973A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Contractions		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 8	MON	Involuntary					
		Mydriasis					

Date:11/07/03ISR Number: 4229869-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387975A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day	1	YR					
20MG Per day	1	YR		Paxil	SS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229870-7Report Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0387993A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	4	MON					

Date:11/07/03ISR Number: 4229871-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0388006A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
YR							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229872-0Report Type:Periodic  
Age:10 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0388019A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
50MG Per day	4	MON		No Concurrent Medication	C		

Date:11/07/03ISR Number: 4229873-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0388022A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/07/03ISR Number: 4229874-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0388025A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	1	MON					

Date:11/07/03ISR Number: 4229875-6Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0388127A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229876-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0388177A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Serotonin Syndrome		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229877-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0388515A  
 Age:30 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	29 DAY						
UNKNOWN	30MG Three			Armour	C		
times per day							
500MG Twice				Glucophage Xr	C		ORAL
per day							
				Allegra	C		ORAL
				Vicodin	C		ORAL

Date:11/07/03ISR Number: 4229878-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0388516A  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Affect Lability		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	16 DAY						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229880-XReport Type:Periodic  
 Age:45 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0388523A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Blood Pressure Diastolic		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	MON	Increased					
150MG Twice		Blood Pressure Systolic		Serzone	C		ORAL
per day		Increased					
UNKNOWN		Flushing		Avonex	C		
				Ibuprofen	C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229881-1Report Type:Periodic  
 Age:42 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0388561A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Chest Pain	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
Hospitalization -		Hypertension	Professional				
per day				Adderall	SS		
Initial or Prolonged							
UNKNOWN	20MG Unknown		WK				
Other							

Date:11/07/03ISR Number: 4229882-3Report Type:Periodic  
 Age:32 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0388589A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Three		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
times per day							
				Buspar	C		
				Propecia	C		
				Doxycycline	C		
100MG Twice							



per day

Advair C Glaxosmithkline  
Nasonex C

Date:11/07/03ISR Number: 4229883-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0388592A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	3 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
		Insomnia					

Date:11/07/03ISR Number: 4229884-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0388597A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Breast Pain Hypertrophy Breast					

Date:11/07/03ISR Number: 4229885-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0388600A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Twice			Wellbutrin	PS	Glaxosmithkline	ORAL
	per day	MON		Elavil	C	Glaxosmithkline	
		Glossodynia Tongue Blistering					



150MG Twice	Hyperhidrosis	Wellbutrin	PS	Glaxosmithkline	ORAL
per day					
300MG per day		Effexor	C		
		Klonopin	C		

Date:11/07/03ISR Number: 4229890-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0388647A  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG per day	4 WK	Drug Ineffective Tremor					

Date:11/07/03ISR Number: 4229891-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0388878A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
2 YR		Hyperventilation		Allegra	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229892-6Report Type:Periodic  
Age:65 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0388970A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	7 DAY	Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
		Urinary Hesitation		Xanax	C		
				Blood Pressure Medication	C		

Date:11/07/03ISR Number: 4229893-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0388971A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	100MG Per day	Swollen Tongue		Wellbutrin	PS	Glaxosmithkline	ORAL
	2 WK	Vomiting		Ritalin	C		
				Effexor Xr	C		

Date:11/07/03ISR Number: 4229894-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0388974A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Mania		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229895-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0388980A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229896-3Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0388984A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lymphadenopathy		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1	MON		Synthroid	C	Glaxosmithkline	
				Lantus	C		

Date:11/07/03ISR Number: 4229897-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0389047A  
 Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day	1	MON		Diazepam	C		
				Provigil	C		
				Fluoxetine	C		

Date:11/07/03ISR Number: 4229898-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0389054A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Screen Positive		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day				Zyprexa	C		
UNKNOWN	10MG Per day						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229899-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0389090A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Agitation					
per day		Flat Affect					
		Tinnitus					

Date:11/07/03ISR Number: 4229900-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0389186A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -		Intentional Misuse					
Initial or Prolonged							

Date:11/07/03ISR Number: 4229902-6Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0389197A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
400MG Per day	7 YR	Drug Ineffective					
UNKNOWN		Drug Interaction		Tegretol	SS		
UNKNOWN				Risperdal	SS		
UNKNOWN				Klonopin	C		

Date:11/07/03ISR Number: 4229904-XReport Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0389225A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 MON	Blood Pressure Increased					

INTRAVENOUS Sinus Headache Lupron C

Date:11/07/03ISR Number: 4229906-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0389226A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Antisocial Behaviour		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Unknown	6 MON			Lithium	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229907-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0389461A  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Flushing					
per day	5 DAY	Nausea		Pamelor	C		
		Road Traffic Accident		Daypro	C		
				Premarin	C		

Date:11/07/03ISR Number: 4229908-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0389471A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229909-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0389508A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Swelling Face		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229910-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0389515A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety		Wellbutrin	PS	Glaxosmithkline	
RESPIRATORY (INHALATION)		Drug Abuser 36 HR Intentional Misuse Medication Error		Beer	C		

Date:11/07/03ISR Number: 4229911-7Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0389524A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Abuser		Wellbutrin	PS	Glaxosmithkline	
RESPIRATORY (INHALATION)		Intentional Misuse 4 MON Medication Error		No Concurrent Medications	C		

Date:11/07/03ISR Number: 4229912-9Report Type:Periodic  
Age:52 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0389628A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	YR	Aggression Anger Chest Pain		Wellbutrin Effexor Xr	PS SS	Glaxosmithkline	ORAL



Flushing  
Insomnia  
Nausea  
Retching  
Stress

Date:11/07/03ISR Number: 4229920-8Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0389668A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Single		Vomiting					
dose				Singulair	C		
				Aerobid	C		
				Maxair	C		
				Oxygen	C		
				Valium	C		
				Flexeril	C		
				Vitamin	C		
				Aspirin	C	Glaxosmithkline	
				Vioxx	C		
				Nebulizer	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229921-XReport Type:Periodic  
Age:19 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0389672A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	121 DAY						

Date:11/07/03ISR Number: 4229922-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0389676A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Enuresis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229923-3Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0389687A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Weight Increased					
per day	7 MON						

Date:11/07/03ISR Number: 4229924-5Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0389688A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Migraine		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Zomig	C		
per day	3 DAY						

Date:11/07/03ISR Number: 4229925-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0389735A

Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	1 YR		Influenza Like Illness	Wellbutrin	PS	Glaxosmithkline	ORAL
			Medication Error				

Date:11/07/03ISR Number: 4229926-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0389736A

Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 WK			Insomnia	Wellbutrin	PS	Glaxosmithkline	ORAL
			Psychomotor Hyperactivity	Prozac	C		

Date:11/07/03ISR Number: 4229927-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0389809A

Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Alopecia	Wellbutrin	PS	Glaxosmithkline	ORAL
				Unknown Medication	C		

Date:11/07/03ISR Number: 4229928-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0389810A

Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Constipation	Wellbutrin	PS	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229929-4Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0389828A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	5 WK			Toprol Xl	C		

Date:11/07/03ISR Number: 4229930-0Report Type:Periodic  
 Age:55 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0389836A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day				Spiroinolactone	SS		
UNKNOWN							

Date:11/07/03ISR Number: 4229932-4Report Type:Periodic  
 Age:49 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0389957A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice		Pharmaceutical Product		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Complaint					
		Stool Analysis Abnormal		Trazodone	C		
				Alprazolam	C		

Date:11/07/03ISR Number: 4229933-6Report Type:Periodic  
 Age:36 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0389961A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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150MG Per day Pruritus Wellbutrin PS Glaxosmithkline ORAL

Date:11/07/03ISR Number: 4229934-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0389965A  
 Age:61 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	3 MON	Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Ineffective					
		Pharmaceutical Product Complaint					

Date:11/07/03ISR Number: 4229935-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0389967A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229936-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0389995A  
 Age:20 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 400MG Per day		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229937-3Report Type:Periodic  
Age:60 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390032A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day							
				Albuterol	C	Glaxosmithkline	
				Advair	C	Glaxosmithkline	
				Zocor	C		
20MG Per day							
				Aciphex	C		
20MG Twice							
per day							
				Ecotrin	C	Glaxosmithkline	
				Bc	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229938-5Report Type:Periodic  
Age:16 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390037A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation Dissociation	Consumer	Wellbutrin Marijuana	PS SS	Glaxosmithkline	ORAL
UNKNOWN		Flat Affect Panic Attack Paranoia					

Date:11/07/03ISR Number: 4229939-7Report Type:Periodic  
Age:59 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390152A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day	4 MON						
				Lipitor	C		

Date:11/07/03ISR Number: 4229940-3Report Type:Periodic  
Age:49 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0390175A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Parosmia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Pharmaceutical Product					
per day	4	MON	Complaint				
			Urine Odour Abnormal				

Date:11/07/03ISR Number: 4229941-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390180A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Hypersensitivity					
per day	3	WK	Rash				
			Therapeutic Response				
			Unexpected				
			Throat Tightness				
			Urticaria				
			Weight Decreased				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229942-7Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0390231A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229943-9Report Type:Periodic  
Age:33 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0390234A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	2 YR	Eructation Flatulence		Wellbutrin	PS	Glaxosmithkline	ORAL
				Seroquel Celexa	C C		

Date:11/07/03ISR Number: 4229944-0Report Type:Periodic  
Age:62 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0390247A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Disorientation Illusion Keratoconjunctivitis Sicca Muscle Twitching		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229946-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390250A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3 DAY	Insomnia Tension		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:11/07/03ISR Number: 4229947-6Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390254A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/07/03ISR Number: 4229948-8Report Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390339A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Stomach Discomfort		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Tinnitus					
per day	14 DAY						

Date:11/07/03ISR Number: 4229949-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390353A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Lethargy					
per day		Obsessive-Compulsive		Zoloft	C		
50MG Per day		Personality Disorder					
		Somnolence					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229950-6Report Type:Periodic  
Age:70 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390357A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Glossodynia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day				Evista	C		
				Levoxyl	C	Glaxosmithkline	
				Glyburide	C		

Date:11/07/03ISR Number: 4229951-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390370A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mouth Ulceration		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1 YR			Serzone	C		
150MG Three							
times per day				Ritalin	C		
150MG Per day	7 YR						

Date:11/07/03ISR Number: 4229952-XReport Type:Periodic  
Age:20 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390380A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
150MG Twice							
per day							

Date:11/07/03ISR Number: 4229953-1Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390388A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Smoker		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4	MON		Klonopin	C		
				Buspar	C		
				Librium	C		

Date:11/07/03ISR Number: 4229954-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0390431A  
 Age:9 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Personality Change					
per day							

Date:11/07/03ISR Number: 4229955-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0390910A  
 Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Contusion		Remeron	C		ORAL
46 DAY		Glossoptosis		Keppra	C		ORAL
750MG Twice		Grand Mal Convulsion					
per day		Tongue Biting		Oxycontin	C		ORAL
80MG Three							
times per day				Olanzapine	C		ORAL
5MG Twice per							
day				Hydroxyzine Pamoate	C		ORAL
50MG Three							
times per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229956-7Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390914A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
WK		Dry Mouth Nervousness					

Date:11/07/03ISR Number: 4229958-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390940A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lymphadenopathy		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day	1 MON	Pruritus					

Date:11/07/03ISR Number: 4229959-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390952A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Migraine		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229960-9Report Type:Periodic  
Age:35 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390954A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sensitivity Of Teeth		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	1 MON			Claritin	C		
				Vitamin	C		

Date:11/07/03ISR Number: 4229961-0Report Type:Periodic  
Age:9 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390959A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG In the morning	2	MON	Weight Increased	Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN	3MG per day			Risperidal	SS		

Date:11/07/03ISR Number: 4229962-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0390970A  
 Age:30 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	54	DAY	Muscle Contractions Involuntary	Wellbutrin	PS	Glaxosmithkline	ORAL
				Prevacid Ambien	C C		ORAL ORAL

Date:11/07/03ISR Number: 4229963-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0391191A  
 Age:32 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	9	MON	Drug Ineffective	Wellbutrin	PS	Glaxosmithkline	ORAL
50MG Three times per day			Feeling Abnormal Feeling Hot Therapeutic Response Unexpected Vomiting	Multivitamins Zoloft	C C		ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229964-6Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391199A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Nausea					
per day	1 WK			Propranolol	C		
				Triamterene	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229965-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391209A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	
Other				Topamax	C		
				Depakote	C		

Date:11/07/03ISR Number: 4229966-XReport Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391264A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
		Nausea					

Date:11/07/03ISR Number: 4229967-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391298A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229968-3Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391301A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Rash					
per day	2	MON					
		Urticaria					

Date:11/07/03ISR Number: 4229969-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0391456A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229970-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0391461A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Affective Disorder Depression Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229971-3Report Type:Periodic  
 Age:59 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391466A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Panic Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	WK						
UNKNOWN				Klonopin	C		
UNKNOWN				Prilosec	C		
UNKNOWN				Lipitor	C		
UNKNOWN				Esculine	C		
UNKNOWN				Soma	C		
UNKNOWN				Ultracet	C		
UNKNOWN				Vicodin	C		
UNKNOWN				Allegra	C		
UNKNOWN				Estrace	C		

Date:11/07/03ISR Number: 4229972-5Report Type:Periodic  
 Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391480A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 YR						

Date:11/07/03ISR Number: 4229973-7Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391484A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229974-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391488A



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
250MG Per day							

Date:11/07/03ISR Number: 4229975-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0391518A  
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anaphylactic Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							
				Percocet	C		
				Flexeril	C		
				Depo Provera	C		

Date:11/07/03ISR Number: 4229976-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0391652A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							

Date:11/07/03ISR Number: 4229977-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0391653A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flat Affect		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day 7 MON							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229978-6Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391654A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	4	MON					
UNKNOWN				Imitrex	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229979-8Report Type:Periodic  
Age:36 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0391667A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Muscle Contractions		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	15	DAY					
UNKNOWN		Involuntary					
	20MG Per day	8	YR	Prozac	C		

Date:11/07/03ISR Number: 4229980-4Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391730A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	17	DAY					
		Rash					
		Urticaria		Lexapro	C		

Date:11/07/03ISR Number: 4229981-6Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0391742A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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6	MON	Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Dyspepsia		Buspar	C		
		Male Sexual Dysfunction		Prozac	C		

Date:11/07/03ISR Number: 4229982-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0391747A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Cogwheel Rigidity		Wellbutrin	PS	Glaxosmithkline	ORAL
400MG Per day							

Date:11/07/03ISR Number: 4229983-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0391748A  
 Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY	Insomnia					

Date:11/07/03ISR Number: 4229984-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0391749A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229985-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391781A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229986-5Report Type:Periodic  
Age:66 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391850A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	4 DAY	Diarrhoea		Atacand	C		

Date:11/07/03ISR Number: 4229987-7Report Type:Periodic  
Age:67 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391859A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Lipitor	C		
per day	WK			Altace	C		
				Toprol	C		
				Coenzyme Q10	C		
				Baby Aspirin	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229988-9Report Type:Periodic  
Age:68 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391861A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Concerta	C		
per day	1 MON						

Buspar	C	
Synthroid	C	Glaxosmithkline
Baby Aspirin	C	Glaxosmithkline
Coenzyme Q10	C	

Date:11/07/03ISR Number: 4229989-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0391867A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3 YR						

Date:11/07/03ISR Number: 4229990-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0391886A  
 Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Insomnia		Buspar	C		
				Zocor	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229992-0Report Type:Periodic  
 Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392081A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Psychomotor Hyperactivity		Milk Thistle	C		
				Daily Vitamin	C		
				Vitamin C	C	Glaxosmithkline	
				Calcium Citrate	C		
				Folic Acid	C		
				Birth Control Pill	C		

Date:11/07/03ISR Number: 4229993-2Report Type:Periodic  
 Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392085A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthma		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1	MON		Advair	C	Glaxosmithkline	
RESPIRATORY							
(INHALATION)							
				Singulair	C		
				Zithromax	C		
				Medrol Dose Pak	C		
				Prednisone	C		
UNKNOWN	40MG Per day						

Date:11/07/03ISR Number: 4229994-4Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392100A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	MON	Stomach Discomfort		Valtrex	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229995-6Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392105A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
300MG Per day	DAY			Albuterol	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4229996-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392158A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4229997-XReport Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392192A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice				Neurontin	C		
per day	4 WK			Lamictal	C	Glaxosmithkline	
				Clonazepam	C		
				Trazodone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4229998-1Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392194A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day				Celexa	C		

Date:11/07/03ISR Number: 4229999-3Report Type:Periodic  
Age:80 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392195A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Avapro	C		
per day	3	MON					

Date:11/07/03ISR Number: 4230000-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392365A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2	MON					

Date:11/07/03ISR Number: 4230001-8Report Type:Periodic  
Age:56 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0392411A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oral Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	5	DAY					



Date:11/07/03ISR Number: 4230002-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0392420A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
MON				Zoloft	C		
				Surmontil	C		

Date:11/07/03ISR Number: 4230003-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0392428A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day							
UNKNOWN				Lithium	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4230005-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0392510A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Flushing Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230006-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392580A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction		Wellbutrin	PS	Glaxosmithkline	ORAL
		Hyperhidrosis		Zoloft	C		
				Effexor	C		

Date:11/07/03ISR Number: 4230007-9Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392597A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Haematuria		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG per day	8 YR	Proteinuria		Luvox	C		
				Lipitor	C		
				Aleve	C		
3	YR						

Date:11/07/03ISR Number: 4230008-0Report Type:Periodic  
Age:26 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0392748A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1 MON						

Date:11/07/03ISR Number: 4230009-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392757A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230010-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392808A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230011-0Report Type:Periodic  
Age:14 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392915A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Irritability		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Rash Generalised					
per day	10 MON			Eskalith	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4230012-2Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392916A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Burning Sensation		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Confusional State					
per day		Dizziness		Imitrex	C	Glaxosmithkline	
UNKNOWN		Myalgia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230014-6Report Type:Periodic  
Age:35 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0392934A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 WK		Crohn'S Disease		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230015-8Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0392935A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
YR		Drug Interaction		Wellbutrin	PS	Glaxosmithkline	ORAL
YR		Liver Function Test		Ritalin	SS		
		Abnormal Road Traffic Accident					

Date:11/07/03ISR Number: 4230016-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392938A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230017-1Report Type:Periodic  
Age:76 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0392955A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3 DAY	Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
25MG Per day				Metoprolol	C		
				Antacid	C		

Date:11/07/03ISR Number: 4230018-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0392960A  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Deafness Neurosensory		Wellbutrin	PS	Glaxosmithkline	ORAL
400MG Per day	5 MON						
Other				Prozac	C		
80MG Per day							

Date:11/07/03ISR Number: 4230019-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0392964A  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
7 WK							
				Zocor	C		
				Enalapril Maleate	C		

Date:11/07/03ISR Number: 4230020-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0392969A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Three		Insomnia					
times per day	10 DAY	Nervousness		Amitriptyline	SS		
UNKNOWN		YR					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230021-3Report Type:Periodic  
Age:37 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392994A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Per day	7 MON		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN	60MG	Unknown		Celexa	C		
UNKNOWN	15MG	Unknown		Buspar	C		
				Hydroxyzine	C		

Date:11/07/03ISR Number: 4230022-5Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393111A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	3 WK			Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230023-7Report Type:Periodic  
Age:23 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393124A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	200MG Twice			Wellbutrin	PS	Glaxosmithkline	ORAL
	per day	3 MON					

Date:11/07/03ISR Number: 4230024-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393192A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urinary Retention		Wellbutrin Zoloft	PS C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230025-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0393193A  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
250MG Per day	1 MON	Drug Ineffective Pharmaceutical Product Complaint Somnolence					

Date:11/07/03ISR Number: 4230026-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0393197A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other	400MG Per day						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230027-4Report Type:Periodic  
 Age:60 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393201A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	3 WK	Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
				Flomax	C		
				Zoloft	C		

Date:11/07/03ISR Number: 4230028-6Report Type:Periodic  
 Age:15 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393205A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Alternate days	1 YR	Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230029-8Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0393208A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Per day		Pharmaceutical Product		Wellbutrin	PS	Glaxosmithkline	ORAL
		Complaint Stool Analysis Abnormal		Paroxetine	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4230030-4Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393210A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:11/07/03ISR Number: 4230032-8Report Type:Periodic  
Age:17 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407166A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
350MG per day	6	MON		Seroquel	C		
				Adderall	C		

Date:11/07/03ISR Number: 4230033-XReport Type:Periodic  
Age:46 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0407314A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Blister		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Erythema					
per day	1	WK		Metoprolol	C		
		Limb Discomfort		Flexeril	C		
		Lip Blister		Oxycodone	C		
		Oral Mucosal Blistering		Naproxen	C		
		Oral Pain					
		Rash					
		Swollen Tongue					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230042-0Report Type:Periodic  
Age:41 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0407315A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Twice		Drug Screen False		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Positive					
				Synthroid	C	Glaxosmithkline	
				Motrin	C	Glaxosmithkline	
				Aspirin	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4230043-2Report Type:Periodic  
Age:25 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0407349A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
UNKNOWN	150MG Per day	Urine Odour Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
				Effexor	C		

Date:11/07/03ISR Number: 4230044-4Report Type:Periodic  
Age:8 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407350A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG In the		Dyspepsia		Wellbutrin	PS	Glaxosmithkline	ORAL
morning				Trileptal	C		

Date:11/07/03ISR Number: 4230045-6Report Type:Periodic  
Age:40 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0407384A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Thrombocytopenia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Unknown							
Hospitalization -							

Initial or Prolonged  
Other

Date:11/07/03ISR Number: 4230046-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407397A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Screen Positive		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230047-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407502A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rectal Haemorrhage		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK						

Date:11/07/03ISR Number: 4230048-1Report Type:Periodic  
Age:16 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0407503A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharmaceutical Product		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day	MON	Complaint Reading Disorder Speech Disorder		Trazodone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230049-3Report Type:Periodic  
 Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407542A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 WK	Arthralgia Joint Swelling		Wellbutrin	PS	Glaxosmithkline	ORAL
				Tamoxifen	C		
				Nicoderm	C	Glaxosmithkline	
				Vitamin E	C		
				Calcium	C		

Date:11/07/03ISR Number: 4230050-XReport Type:Periodic  
 Age:25 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407546A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG per day	31 DAY	Rash Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
				Protein Drink	C		

Date:11/07/03ISR Number: 4230051-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407550A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	1 YR	Headache		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230052-3Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407554A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Vomiting		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230053-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407639A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vomiting		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230054-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407640A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aphasia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230055-9Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407650A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Verapamil	C		
per day				Furosemide	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230056-0Report Type:Periodic  
Age:55 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0407653A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	MON	Arthralgia Chills Connective Tissue Disorder Myalgia Pain Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230057-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407654A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230058-4Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407704A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 MON	Dizziness Headache Nausea		Wellbutrin Protonix	PS C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230059-6Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407726A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Back Pain Burning Sensation Dry Mouth		Wellbutrin Diovan Hct Synthroid	PS C C	Glaxosmithkline Glaxosmithkline	ORAL

Tremor

Date:11/07/03ISR Number: 4230060-2Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407730A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Celexa C

Date:11/07/03ISR Number: 4230061-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407802A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230062-6Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407803A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	2 YR						

Ambien C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Trazodone C  
 Bextra C  
 Hypoglycemic C

Date:11/07/03ISR Number: 4230063-8Report Type:Periodic  
 Age:21 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407808A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Swollen Tongue					
per day	12 DAY			Depakote	C		

Date:11/07/03ISR Number: 4230064-XReport Type:Periodic  
 Age:58 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0407840A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1 MON			Prozac	C		

Date:11/07/03ISR Number: 4230065-1Report Type:Periodic  
 Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407849A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2 WK						

Date:11/07/03ISR Number: 4230066-3Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407877A



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mania		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day							

Date:11/07/03ISR Number: 4230067-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0407946A  
 Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Screen False		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Three		Positive					
times per day				Celexa	C		
				Motrin	C	Glaxosmithkline	
				Allegra	C		

Date:11/07/03ISR Number: 4230068-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0407953A  
 Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Depressed Mood					
per day	3 WK	Eye Irritation		Restoril	C		
		Yawning					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230069-9Report Type:Periodic  
Age:19 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0408231A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Somnolence		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG per day				Clarinet	C		
				Propecia	C		

Date:11/07/03ISR Number: 4230070-5Report Type:Periodic  
Age:27 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408232A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Dyspnoea					
per day	16 DAY						

Date:11/07/03ISR Number: 4230071-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0408249A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Three							
times per day	1 YR						

Date:11/07/03ISR Number: 4230072-9Report Type:Periodic  
Age:38 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0408262A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cyanosis		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	YR	Heart Rate Increased		Allegra	C		
		Hyperventilation		Synthroid	C	Glaxosmithkline	
		Palpitations		Coffee	C		
		Panic Reaction					

Stress

Date:11/07/03ISR Number: 4230074-2Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408351A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230075-4Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408352A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230076-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408405A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Stevens-Johnson Syndrome		Bupropion	PS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230078-XReport Type:Periodic  
 Age:21 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0408554A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	28	DAY					
		Grand Mal Convulsion					
		Loss Of Consciousness		Alcohol	C		ORAL
		Postictal State		Ativan	C		ORAL
3MG Per day							
		Urinary Incontinence		Norco	C		ORAL
				Effexor	C		

Date:11/07/03ISR Number: 4230083-3Report Type:Periodic  
 Age:15 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408564A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Contusion		Wellbutrin	PS	Glaxosmithkline	ORAL
per day							
		Nail Bed Tenderness					
				Antibiotic	C		

Date:11/07/03ISR Number: 4230084-5Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408565A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230085-7Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0408622A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
		Stool Analysis Abnormal		Fiber Laxative	C		

Date:11/07/03ISR Number: 4230086-9Report Type:Periodic  
Age:27 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408624A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Myalgia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	DAY			Contraceptive	C		ORAL

Date:11/07/03ISR Number: 4230087-0Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0408625A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
MON							

Date:11/07/03ISR Number: 4230088-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408644A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nocturia		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230089-4Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408647A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
1 WK				Atenolol	C		
				Progestin	C		
				Estrogen	C		
				Imitrex	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4230090-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408665A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Health	Bupropion	PS	Glaxosmithkline	ORAL
Other			Professional	Alcohol	SS		
250MG per day							

Date:11/07/03ISR Number: 4230091-2Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0408671A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Health	Bupropion	PS	Glaxosmithkline	ORAL
Other			Professional	Benzodiazepine	SS		
300MG per day				Effexor	C		

Date:11/07/03ISR Number: 4230092-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408683A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230093-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0408696A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:11/07/03ISR Number: 4230094-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0408698A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
6 WK							

Date:11/07/03ISR Number: 4230095-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0408814A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urinary Retention		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230096-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0408815A  
Age:48 YR Gender:Female I/FU:F

Outcome  
PT  
Dry Mouth  
Dysuria

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Urinary Retention

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	MON		Wellbutrin	PS	Glaxosmithkline	ORAL
40MG In the morning	1 YR		Nexium	C		ORAL
10MG At night	MON		Lipitor	C		ORAL
1MG In the morning	MON		Arimidex	C		ORAL
600MG Three times per day	1 YR		Neurontin	C		ORAL
.5MG Four times per day	MON		Klonopin	C		ORAL

Date:11/07/03ISR Number: 4230097-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0408816A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Interaction		Detrol La	SS		ORAL

Date:11/07/03ISR Number: 4230098-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0408916A  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Photosensitivity Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL
2 WK							



Date:11/07/03ISR Number: 4230099-7Report Type:Periodic  
Age:9 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408921A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
1	WK						

Date:11/07/03ISR Number: 4230100-0Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408925A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Anxiety					
per day	119 DAY	Demented Relative		Klonopin	C		

Date:11/07/03ISR Number: 4230101-2Report Type:Periodic  
Age:52 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0409018A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Bupropion	PS	Glaxosmithkline	ORAL
Hospitalization -							
150MG Twice							
Initial or Prolonged							
per day				Keppra	C		
				Lexapro	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230102-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0409040A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Bupropion	PS	Glaxosmithkline	ORAL
150MG Twice		Medication Error					
per day							

Date:11/07/03ISR Number: 4230103-6Report Type:Periodic  
Age:26 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0409058A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Bupropion	PS	Glaxosmithkline	ORAL
150MG Three		Loss Of Consciousness					
times per day	270 DAY	Postictal State					
		Simple Partial Seizures					

Date:11/07/03ISR Number: 4230104-8Report Type:Periodic  
Age:69 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0409190A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	2 MON			Lexapro	C		

Date:11/07/03ISR Number: 4230105-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0409191A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day	2 WK						

Date:11/07/03ISR Number: 4230106-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0409192A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3 YR	Drug Ineffective		Prevacid	C		
				Lodine	C		

Date:11/07/03ISR Number: 4230107-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0409193A  
 Age:36 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3 MON			Synthroid	C	Glaxosmithkline	
				Phentermine	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4230108-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0409305A  
 Age: Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Unknown	3 WK	Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230109-7Report Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0409466A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	6	MON					

Date:11/07/03ISR Number: 4230110-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0409468A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Loss Of Consciousness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1	WK					

Date:11/07/03ISR Number: 4230111-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0409469A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flatulence		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3	DAY					
		Insomnia					

Date:11/07/03ISR Number: 4230112-7Report Type:Periodic  
Age:50 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0409470A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day	1	YR					
UNKNOWN	10MG	Unknown		Paxil	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4230113-9Report Type:Periodic  
Age:39 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0409471A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Unknown	2 YR		Dermatitis Allergic	Wellbutrin	PS	Glaxosmithkline	ORAL
				Topamax	C		
				Lexapro	C		
				Provigil	C		

Date:11/07/03ISR Number: 4230114-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0409487A  
Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG per day	5 YR		Feeling Abnormal	Wellbutrin	PS	Glaxosmithkline	ORAL
			Insomnia				
			Pollakiuria				
			Tinnitus				

Date:11/07/03ISR Number: 4230116-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0409517A  
Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice	3 MON		Weight Increased	Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230117-6Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0409562A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Bupropion	PS	Glaxosmithkline	ORAL
200MG Twice per day 75 DAY							

Date:11/07/03ISR Number: 4230118-8Report Type:Periodic  
Age:35 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0409568A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Electroencephalogram		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230119-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0409636A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230127-9Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0409638A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Flushing		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230128-0Report Type:Periodic  
Age:53 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0409640A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day							
		Anxiety	Health	Wellbutrin	PS	Glaxosmithkline	ORAL

5	YR	Depression	Professional	Zoloft	SS	ORAL
		Drug Ineffective		Ansaid	C	
	100MG Twice	Fatigue				
	per day	Fibromyalgia				
		Hyperhidrosis				
		Insomnia				
		Muscle Spasms				
		Myalgia				
		Sedation				
		Yawning				

Date:11/07/03ISR Number: 4230129-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0409943A  
 Age:62 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Amnesia	Health	Bupropion	PS	Glaxosmithkline	ORAL
2 YR						
Initial or Prolonged	Convulsion	Professional	Alcohol	SS		ORAL
Other	Tongue Biting		Xanax	C		
UNKNOWN	.5MG As					
required						
UNKNOWN	10MG Unknown		Ambien	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230130-9Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0409944A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK			Motrin	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4230131-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0410041A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230132-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0410042A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tooth Discolouration		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2 WK						

Date:11/07/03ISR Number: 4230133-4Report Type:Periodic  
Age:30 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0410087A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Convulsion	Health	Bupropion	PS	Glaxosmithkline	ORAL
150MG Twice							
Initial or Prolonged		Head Injury	Professional				
per day	1 YR						
Other		Stress					

Date:11/07/03ISR Number: 4230134-6Report Type:Periodic  
Age:47 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0410159A



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Headache		Bupropion	PS	Glaxosmithkline	ORAL
150MG Twice							
Other		Hypertension					
per day	8 DAY			Diuretic	C		
				Actonel	C		

Date:11/07/03ISR Number: 4230135-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0410172A  
 Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Discomfort		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230136-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0410290A  
 Age:14 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Per day	40 DAY	Priapism		Wellbutrin	PS	Glaxosmithkline	ORAL
		Spontaneous Penile Erection					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230141-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0410291A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
		Thinking Abnormal					

Date:11/07/03ISR Number: 4230142-5Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0410304A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG per day	2 WK	Rash		Birth Control	C		
		Urticaria					

Date:11/07/03ISR Number: 4230143-7Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0410323A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/07/03ISR Number: 4230144-9Report Type:Periodic  
Age:11 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0410333A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Insomnia					
per day	6 MON						

Date:11/07/03ISR Number: 4230145-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0410335A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/07/03ISR Number: 4230146-2Report Type:Periodic  
Age:80 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0410339A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Disorientation					
per day	6 WK	Fatigue		Zocor	C		
				Lexapro	C		
				Xalatan	C		
				Lorazepam	C		
.5MG At night				Folic Acid	C		
				Vitamin E	C		
				Imodium	C		
				Aspirin	C	Glaxosmithkline	
				Multivitamin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230147-4Report Type:Periodic  
Age:55 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0410381A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache Keratoconjunctivitis Sicca Nasal Dryness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230148-6Report Type:Periodic  
Age:45 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0410500A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Unknown							

Date:11/07/03ISR Number: 4230149-8Report Type:Periodic  
Age:48 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0410501A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anaemia		Wellbutrin	PS	Glaxosmithkline	ORAL
Other 150MG See dosage text							
		Chest Pain					
		Dyspepsia		Gaviscon	C	Glaxosmithkline	
		Gastric Disorder		Black Cohosh	C	Glaxosmithkline	
		Gastrooesophageal Reflux Disease		Echinacea	C		
				Aloe Vera	C		
				Tylenol Arthritis	C	Glaxosmithkline	
				Calcium	C		
				Aspirin	C	Glaxosmithkline	
				Amitriptyline	C		
				Ginger Root	C		
				Acidophilus	C		
				Hawthorn Berry	C		
				Kava-Kava	C		

Date:11/07/03ISR Number: 4230154-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0410502A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sleep Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	5 DAY						

Date:11/07/03ISR Number: 4230155-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0410590A  
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Pruritic		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Urticaria					
per day	19 DAY						

Date:11/07/03ISR Number: 4230156-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0410618A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4 DAY						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230157-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0410623A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230158-9Report Type:Periodic  
Age:72 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0410743A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gastrointestinal		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Haemorrhage		Lithium	C	Glaxosmithkline	
150MG Per day	1 WK			Risperdal	C		

Date:11/07/03ISR Number: 4230159-0Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0410781A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Tremor					
per day	1 YR			Prilosec	C		
UNKNOWN				Estrogen Replacement	C		
UNKNOWN							

Date:11/07/03ISR Number: 4230160-7Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0410838A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	13 DAY	Flatulence		Aspirin	C	Glaxosmithkline	
UNKNOWN							

UNKNOWN

Vitamins C

UNKNOWN

Folic Acid C

Date:11/07/03ISR Number: 4230161-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0410842A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus Thermal Burn		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230162-0Report Type:Periodic  
Age:16 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0410843A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Life-Threatening 1200MG Single Hospitalization - dose 1 DAY Initial or Prolonged 12TAB Single Other dose 1 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
		Overdose		Celexa	SS		ORAL

Date:11/07/03ISR Number: 4230163-2Report Type:Periodic  
Age:47 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0411022A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hospitalization - 150MG Single Initial or Prolonged dose 1 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
		Anxiety		No Concurrent Medications	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230164-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0411050A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hyperhidrosis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230165-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0411187A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
		Dyspepsia		Paxil	SS	Glaxosmithkline	ORAL
		Headache		Sarafem	SS		
UNKNOWN		Pruritus					
		Urticaria					

Date:11/07/03ISR Number: 4230166-8Report Type:Periodic  
Age:68 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0411257A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Crying		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice		Dry Mouth					
per day	1 MON	Tremor		Lexapro	C		
UNKNOWN				Diovan	C		
UNKNOWN				Triavil	C		
UNKNOWN							

Date:11/07/03ISR Number: 4230167-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0411262A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:11/07/03ISR Number: 4230168-1Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0411264A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230169-3Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0411266A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Contusion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230170-XReport Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0411371A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Premature Ejaculation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	6	WK					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230171-1Report Type:Periodic  
Age:59 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0411372A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Asthenia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1 MON	Dizziness					
		Tremor		Tussin With Codeine	C		
		Urine Flow Decreased		Oxygen Therapy	C		

Date:11/07/03ISR Number: 4230172-3Report Type:Periodic  
Age:65 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0411373A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Unknown	YR	Diarrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
2 YR		Gastroenteritis Bacterial		Singulair	C		
UNKNOWN		Palpitations		Paxil	C	Glaxosmithkline	
		Tremor					
		Weight Decreased					

Date:11/07/03ISR Number: 4230173-5Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0411374A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Musculoskeletal Stiffness		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 WK	Pharyngolaryngeal Pain					

Date:11/07/03ISR Number: 4230179-6Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0411375A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Medication Error Wellbutrin PS Glaxosmithkline ORAL

100MG Twice  
per day 1 MON

Date:11/07/03ISR Number: 4230180-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0411376A  
Age: Gender:Female I/FU:I

Outcome PT Report Source Product Role Manufacturer Route  
Dose Duration Oedema Peripheral Wellbutrin PS Glaxosmithkline ORAL  
150MG Twice  
per day Pharyngolaryngeal Pain  
Pruritus  
Urticaria

Date:11/07/03ISR Number: 4230182-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0411377A  
Age: Gender:Male I/FU:I

Outcome PT Report Source Product Role Manufacturer Route  
Dose Duration Hallucination Wellbutrin PS Glaxosmithkline ORAL  
200MG Per day  
2 YR Lamictal C Glaxosmithkline ORAL

Date:11/07/03ISR Number: 4230184-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0411401A  
Age: Gender: I/FU:I

Outcome PT Report Source Product Role Manufacturer Route  
Dose Duration Suicidal Ideation Wellbutrin PS Glaxosmithkline ORAL



150MG Per day	10	DAY	Anorgasmia	Wellbutrin	PS	Glaxosmithkline	ORAL
			Libido Decreased	Klonopin	C		

Date:11/07/03ISR Number: 4230190-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0411555A  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Elevated Mood		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	26	Euphoric Mood					
UNKNOWN	37.5MG	Feeling Drunk		Maxzide	C	Glaxosmithkline	
Unknown		Somnolence					
				Atenolol	C		
				Synthroid	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4230191-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0411556A  
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
400MG Per day		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230192-9Report Type:Periodic  
Age:59 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0411557A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	WK		Wellbutrin	PS	Glaxosmithkline	ORAL
		Feeling Abnormal		Premarin	C		
				Vitamins	C		
				Ditropan	C		

Date:11/07/03ISR Number: 4230193-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0411558A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Urine Amphetamine Positive					

Date:11/07/03ISR Number: 4230194-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0411635A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion					

Date:11/07/03ISR Number: 4230195-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0411780A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Unknown			Wellbutrin	PS	Glaxosmithkline	ORAL
		Hyperaesthesia					

Date:11/07/03ISR Number: 4230196-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0411933A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Twice	Headache	Wellbutrin	PS	Glaxosmithkline	ORAL	
per day	1 WK					
Date:11/07/03ISR Number: 4230197-8Report Type:Periodic		Company Report #US-GLAXOSMITHKLINE-A0412081A				
Age:	Gender:Female	I/FU:I				
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Csf Pressure Increased		Bupropion	PS	Glaxosmithkline	ORAL
Date:11/07/03ISR Number: 4230198-XReport Type:Periodic		Company Report #US-GLAXOSMITHKLINE-A0412085A				
Age:45 YR	Gender:Male	I/FU:F				
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Dyspepsia		Bupropion	PS	Glaxosmithkline	ORAL
100MG Three						
times per day	3 YR		Seroquel	C		
Date:11/07/03ISR Number: 4230199-1Report Type:Periodic		Company Report #US-GLAXOSMITHKLINE-A0412092A				
Age:41 YR	Gender:Female	I/FU:F				
Outcome	PT					
Other	Cold Sweat					
	Delusion					
19-Aug-2005	12:44 PM					
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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	8 YR	Dizziness Dysgeusia Syncope Weight Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN				Vitamins	C		
UNKNOWN				Acidophilis	C		
UNKNOWN				Ranitidine	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4230200-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0412105A  
 Age:62 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	1 YR		Crying Depressed Mood		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230201-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0412107A  
 Age:46 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Twice per day		Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL
					Celexa	C		
					Prilosec	C		

Date:11/07/03ISR Number: 4230202-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0412214A  
 Age:50 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	100MG Twice		Back Pain		Wellbutrin	PS	Glaxosmithkline	ORAL



per day	2	WK	Dry Mouth				
			Local Swelling	Prevacid	C		
			Micturition Disorder	Saw Palmetto	C		
			Rash	Flonase	C	Glaxosmithkline	NASAL
				Glutamine	C		
				Mega Vitamins	C		
				Melatonin	C		
				5-Htp	C		
				Vitamin B	C		
				Gingko	C		
				Ativan	C		
				Kelp	C		

Date:11/07/03ISR Number: 4230203-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0412246A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Energy Increased	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	5	DAY					
		Insomnia					
		Therapeutic Response		Periostat	C	Glaxosmithkline	
		Unexpected		Norvasc	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230204-2Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0412257A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day							

Date:11/07/03ISR Number: 4230205-4Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0412277A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230206-6Report Type:Periodic  
 Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0412281A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice per day							
	2	MON		Birth Control Pills	C		
		Dysgeusia		Lomotil	C		

Date:11/07/03ISR Number: 4230207-8Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0412410A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharmaceutical Product		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice per day							
	6	WK		Complaint			
		Stool Analysis Abnormal					

Date:11/07/03ISR Number: 4230208-XReport Type:Periodic  
Age:37 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0412411A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Screen Positive		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	9 YR			Clarinet	C		
				Valium	C		
				Albuterol	C	Glaxosmithkline	

RESPIRATORY  
(INHALATION)

Date:11/07/03ISR Number: 4230209-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0412412A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Dyspnoea					
per day	2 WK	Hyperhidrosis		Synthroid	C	Glaxosmithkline	
		Nausea		Norvasc	C		
				Atacand	C		

Date:11/07/03ISR Number: 4230210-8Report Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0412413A

Outcome PT  
Abdominal Discomfort  
Fatigue

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Insomnia				
		Weight Increased				
Dose	Duration		Report Source	Product	Role	Manufacturer
150MG	Three			Wellbutrin	PS	Glaxosmithkline
times per day	1	YR				ORAL
				Buspar	C	
				Prilosec	C	
				Seroquel	C	
				Didanosine	C	

Date:11/07/03ISR Number: 4230211-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0412414A  
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline
100MG	Twice	Chest Pain				ORAL
per day	1	MON		Acid Reflux Med.	C	
UNKNOWN		Hunger		Calcium	C	

Date:11/07/03ISR Number: 4230212-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0412416A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Rash		Wellbutrin	PS	Glaxosmithkline
150MG	Unknown					ORAL

Date:11/07/03ISR Number: 4230213-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0412487A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline
						ORAL

Date:11/07/03ISR Number: 4230214-5Report Type:Periodic  
Age:59 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0412617A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Twice		Dry Mouth		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2	Tremor					
		MON					
UNKNOWN				Effexor	SS		

Date:11/07/03ISR Number: 4230215-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0412618A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
450MG Per day		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230216-9Report Type:Periodic  
Age:18 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0412619A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Syncope		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
Initial or Prolonged							
per day	1	YR					
Other				Lithium	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230217-0Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0412628A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
		Weight Decreased					

Date:11/07/03ISR Number: 4230218-2Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0412678A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Obsessive-Compulsive		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice		Disorder					
per day	7	MON		Zoloft	C		ORAL
200MG Unknown							

Date:11/07/03ISR Number: 4230219-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0412812A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/07/03ISR Number: 4230220-0Report Type:Periodic  
Age:15 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0412813A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	18	DAY					
		Difficulty In Walking					
		Joint Swelling					
		Urticaria					

Date:11/07/03ISR Number: 4230221-2Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0412814A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG See							
dosage text							
				Lamictal	C	Glaxosmithkline	
				Ambien	C		

Date:11/07/03ISR Number: 4230222-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413098A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/07/03ISR Number: 4230223-6Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0413139A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
MON							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230224-8Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413266A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Gingival Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Tooth Abscess					
		Tooth Disorder		Antibiotic	C		

Date:11/07/03ISR Number: 4230225-XReport Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413269A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day				Synthroid	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4230226-1Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413270A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG per day	2 YR	Keratoconjunctivitis		Wellbutrin	PS	Glaxosmithkline	ORAL
		Sicca		Clonopin	C		

Date:11/07/03ISR Number: 4230227-3Report Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413276A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 DAY		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
		Rash		No Concurrent Medication	C		



Date:11/07/03ISR Number: 4230228-5Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413292A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Unknown	3 DAY			Quinine	C		ORAL
250MG Per day				Xanax	C		
				Vitamins	C		

Date:11/07/03ISR Number: 4230229-7Report Type:Periodic  
Age:17 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413368A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	1 DAY	Vomiting					

Date:11/07/03ISR Number: 4230230-3Report Type:Periodic  
Age:42 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0413382A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Palpitations		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230232-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413540A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Fatigue					
per day	10 DAY	Mood Swings					

Date:11/07/03ISR Number: 4230233-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413754A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Disturbance In Attention		Coreg	C	Glaxosmithkline	
		Memory Impairment		Acyclovir	C	Glaxosmithkline	
		Vision Blurred		Vitamin C	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4230234-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413755A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Headache					

Date:11/07/03ISR Number: 4230235-2Report Type:Periodic  
Age:39 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0413756A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -		Abdominal Pain					
150MG Twice		Anorexia					
Initial or Prolonged							
per day	1 DAY	Liver Function Test					
Other		Abnormal					
		Nausea					
		Vomiting					

Date:11/07/03ISR Number: 4230236-4Report Type:Periodic  
Age:16 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0413757A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dysphemia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1	MON		Minocycline	C		

Date:11/07/03ISR Number: 4230237-6Report Type:Periodic  
Age:65 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413758A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hyperhidrosis		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Vomiting					
per day	2	WK		Zoloft	C		
				Lisinopril	C		
				Omeprazole	C		
				Lescol Xl	C		
				Toprol Xl	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230238-8Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413768A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bruxism		Wellbutrin Unspecified Medications	PS  C	Glaxosmithkline	ORAL
UNKNOWN							

Date:11/07/03ISR Number: 4230239-XReport Type:Periodic  
 Age:42 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0413792A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gingival Bleeding		Wellbutrin	PS	Glaxosmithkline	ORAL
350MG Per day	8 MON	Insomnia		Restoril	C		

Date:11/07/03ISR Number: 4230240-6Report Type:Periodic  
 Age:45 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413803A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Depressed Level Of Consciousness Feeling Abnormal Gait Disturbance Memory Impairment Somnolence		Prilosec	C		
150MG Per day	5 YR						

Date:11/07/03ISR Number: 4230241-8Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0413935A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							

per day	Agitation					
UNKNOWN	Hepatitis C			Peg-Intron	SS	
UNKNOWN	Impulsive Behaviour			Rebetol	SS	
600MG Per day				Neurontin	C	
				Amitriptyline	C	

Date:11/07/03ISR Number: 4230242-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0413937A  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice		Feeling Abnormal					
per day	1 YR	Irritability		Lexapro	SS		
UNKNOWN	10MG Per day			Ambien	C		
UNKNOWN	10MG At night						

Date:11/07/03ISR Number: 4230243-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0413938A  
 Age:19 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Photosensitivity Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL
225MG Per day	2 WK			Sunscreen	C		
TOPICAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230244-3Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413982A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
				Premarin	C		

Date:11/07/03ISR Number: 4230245-5Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413999A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230246-7Report Type:Periodic  
Age:45 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0414010A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 MON	Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN		Pharmaceutical Product Complaint		Synthroid	C	Glaxosmithkline	
		Retching					

Date:11/07/03ISR Number: 4230250-9Report Type:Periodic  
Age:27 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0414034A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 MON	Amenorrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230251-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0414186A  
Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Three		Blood Creatine		Wellbutrin	PS	Glaxosmithkline	ORAL
times per day 1	1	YR					
		Phosphokinase Increased					
		Myoglobin Blood Increased					

Date:11/07/03ISR Number: 4230252-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0414187A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Ageusia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		WK					

Date:11/07/03ISR Number: 4230253-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0414332A  
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
37.5MG Twice		Daydreaming		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	10	DAY					
		Insomnia					
		Somnolence		Potaba	C		
		Yawning		Prevacid	C		
				Oxycodone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230254-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0414333A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	600MG per day	Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
		Overdose					

Date:11/07/03ISR Number: 4230255-8Report Type:Periodic  
Age:25 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0414353A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	1TAB Per day	Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
	1 YR	Fatigue					
		Nausea					
		Pharmaceutical Product Complaint					

Date:11/07/03ISR Number: 4230256-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0414402A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Twice per day	Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
	7 WK						

Date:11/07/03ISR Number: 4230258-3Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0414409A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	Asthenia		Wellbutrin	PS	Glaxosmithkline	ORAL
	4 MON						



Date:11/07/03ISR Number: 4230259-5Report Type:Periodic  
Age:45 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0414532A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1	YR					

Date:11/07/03ISR Number: 4230260-1Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0414540A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sunburn		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1	YR					

Date:11/07/03ISR Number: 4230261-3Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0414553A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day	3	DAY		Fiorinal	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230262-5Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0414554A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 WK	Dry Mouth Dysgeusia Gingival Infection Oral Candidiasis Oral Fungal Infection Tongue Discolouration		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230263-7Report Type:Periodic  
Age:21 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0414569A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice per day		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230264-9Report Type:Periodic  
Age:9 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0414592A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day	3 MON	Difficulty In Walking		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN		Dyskinesia		Zyrtec	C	Glaxosmithkline	
		Dystonia Eye Movement Disorder Muscle Spasms					

Date:11/07/03ISR Number: 4230265-0Report Type:Periodic  
Age:61 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0414617A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	200MG per day	9 MON	Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
			Dysgeusia		Wellbutrin	SS	Glaxosmithkline	ORAL
			Reaction To Azo-Dyes					

Date:11/07/03ISR Number: 4230266-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0414623A  
 Age:54 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Twice	per day	Hypoglycaemia		Wellbutrin	PS	Glaxosmithkline	ORAL
		2 YR			Sinus Medication	C		
					Arthritis Medication	C		

Date:11/07/03ISR Number: 4230268-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0414627A  
 Age: Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230269-8Report Type:Periodic  
Age:47 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0414628A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1 YR	Fibromyalgia		Armour	C		
15MG Three				Klonopin	C		
times per day				Seroquel	C		
.05MG Three				Trazodone	C		
times per day				Lamictal	C	Glaxosmithkline	
200MG Per day				Claritin D	C		
300MG Per day				Nasonex	C		
200MG Twice				Allergy Shots	C		
per day				Dhea	C		
25MG Per day				Multivitamin	C		
				Selenium	C		
				Magnesium	C		
				Folic Acid	C		
				Calcium	C		
				Miralax	C		
				Tylenol	C	Glaxosmithkline	
				Biotin	C		
				Primrose Oil	C		
				Flax Seed Oil	C		

Date:11/07/03ISR Number: 4230270-4Report Type:Periodic  
Age:32 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0414923A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Weight Increased					
per day	1	MON		Contraceptive	C		

Date:11/07/03ISR Number: 4230271-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0415187A  
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Libido Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day				Viagra	C		
				Lipitor	C		
				Nexium	C		
				Methocarbamol	C		
				Vitamins	C		
				Rabeprazole	C		

Date:11/07/03ISR Number: 4230272-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0415214A  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Insomnia					
per day	8	DAY		Mvi	C		
		Paranoia		Lipitor	C		
		Vision Blurred		Aspirin	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nexium	C
Altace	C
Nadolol	C
Folic Acid	C
Ditropan Xl	C
Plavix	C
Flexeril	C

Date:11/07/03ISR Number: 4230273-XReport Type:Periodic  
Age:26 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0415223A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Diarrhoea	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1	WK	Medication Residue				
			Stool Analysis Abnormal				
			Urticaria				

Date:11/07/03ISR Number: 4230274-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0415232A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Somnolence	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4230275-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399975A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		YR	Dry Mouth	Wellbutrin	PS	Glaxosmithkline	ORAL
				Elavil	C	Glaxosmithkline	
				Effexor	C		

Date:11/07/03ISR Number: 4230276-5Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399976A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3	WK		Birth Control	C		

Date:11/07/03ISR Number: 4230277-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0399981A  
Age: Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
		Nausea		Zestril	C		
				Glucophage	C		

Date:11/07/03ISR Number: 4230283-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0399988A  
Age:64 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1	MON		Klonopin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4230284-4Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399991A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mydriasis		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Unknown	8 DAY					
		Nervousness					

Date:11/07/03ISR Number: 4230285-6Report Type:Periodic  
Age:35 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399999A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation		Wellbutrin	PS	Glaxosmithkline	ORAL
2	WK						
		Fatigue					

Date:11/07/03ISR Number: 4230286-8Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400003A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aptyalism		Wellbutrin	PS	Glaxosmithkline	ORAL
		Dental Caries					
		Depression					
		Dry Mouth					
		Pain					
		Suicidal Ideation					
		Tooth Loss					

Date:11/07/03ISR Number: 4230287-XReport Type:Periodic  
Age:68 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0400008A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Coma		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -							
150MG Twice							
Initial or Prolonged		Convulsion					
per day	3	MON					



Other Paxil C Glaxosmithkline ORAL  
 10MG Per day

Date:11/07/03ISR Number: 4230288-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0400012A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Serotonin Syndrome		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN	40MG	Unknown		Prozac	C		

Date:11/07/03ISR Number: 4230289-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0400014A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Prozac	C		

Date:11/07/03ISR Number: 4230290-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0400019A  
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150VA See		Arthralgia					
dosage text	6 MON	Insomnia		Avandia	SS	Glaxosmithkline	ORAL
		Tinnitus		Multivitamin	C		

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150MG Twice			Drug Use For Unknown	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	31	DAY	Indication				
			Rash	Nasonex	C		
				Calcium	C		

Date:11/07/03ISR Number: 4231709-0Report Type:Direct Company Report #CTU 205561  
 Age:43 YR Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Headache		Bupropion Sr 150 Mg	PS		
150 MG QD							

Date:11/07/03ISR Number: 4231714-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0415242A  
 Age:14 YR Gender:Male I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Extrasystoles		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Hyperacusis					
per day	6	WK					

Date:11/07/03ISR Number: 4231715-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0415787A  
 Age:38 YR Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Coumadin C Glaxosmithkline

Date:11/07/03ISR Number: 4231716-8Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0415799A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 WK	Euphoric Mood Urticaria Weight Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231721-1Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0415801A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
per day				Quinine Alprazolam	C C		

Date:11/07/03ISR Number: 4231722-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0415802A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Feeling Hot		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1 YR			Clonopin	C		

Date:11/07/03ISR Number: 4231723-5Report Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0415803A

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Disturbance In Attention		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
		Drug Ineffective					
per day	1	MON					

Date:11/07/03ISR Number: 4231724-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0415805A  
 Age: Gender: I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Unknown							
		Nightmare					

Date:11/07/03ISR Number: 4231725-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0415806A  
 Age:11 YR Gender: I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Tic		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231726-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0415807A  
 Age: Gender: I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231727-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0415808A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Halitosis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231728-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0415809A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	4 YR	Drug Ineffective		Diovan Hct	C		
				Lipitor	C		

Date:11/07/03ISR Number: 4231729-6Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0415810A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fluid Retention		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	8 WK	Photosensitivity Reaction		Celebrex	C		
		Weight Increased		Zoloft	C		

Date:11/07/03ISR Number: 4231730-2Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0415811A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Unknown	6 WK	Fibromyalgia		Lexapro	C		
UNKNOWN	20MG Per day	Headache		Synthroid	C	Glaxosmithkline	
		Tremor		Estratest	C		
				Lamisil	C	Glaxosmithkline	
				Bextra	C		
				Vicodin	C		

Date:11/07/03ISR Number: 4231731-4Report Type:Periodic  
Age:71 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0415812A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Ageusia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	140 DAY	Dry Mouth					
UNKNOWN	.5MG Twice			Klonopin	C		
per day	69 DAY						
UNKNOWN	.5MG At night			Risperdal	C		
UNKNOWN	10MG Three			Ambien	C		
times per day							

Date:11/07/03ISR Number: 4231732-6Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0415813A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 WK						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231733-8Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0415815A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Fatigue Irritability Libido Decreased Weight Increased		Wellbutrin Mircette	PS C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231734-XReport Type:Periodic  
Age:54 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0416549A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3 YR	Skin Odour Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231735-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0416591A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231736-3Report Type:Periodic  
Age:55 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0416592A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG per day	5 YR	Hunger Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:11/07/03ISR Number: 4231737-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0416593A  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Elevated Mood		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3 WK	Therapeutic Response					
		Decreased					

Date:11/07/03ISR Number: 4231738-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0416654A  
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							
Initial or Prolonged		Grand Mal Convulsion		Prozac	C		
Other		Incontinence		Elavil	C	Glaxosmithkline	
UNKNOWN		Tongue Biting					

Date:11/07/03ISR Number: 4231739-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0416671A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
30 DAY		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231740-5Report Type:Periodic  
Age:33 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0416679A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	7 WK	Muscle Twitching Pruritus Generalised		Wellbutrin Prozac	PS C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231741-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0416860A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety Panic Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231742-9Report Type:Periodic  
Age:59 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0416864A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	10 DAY	Feeling Abnormal Gait Disturbance Pain In Extremity		Wellbutrin Blood Pressure Medication Xanax	PS C C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231743-0Report Type:Periodic  
Age:73 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0416865A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	3 MON	Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

4	YR			Singulair	C
4	YR			Theodur	C
4	YR			Q-Bid La	C
4	YR			Ativan	C

Date:11/07/03ISR Number: 4231744-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0416870A  
 Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Transient Ischaemic		Bupropion	PS	Glaxosmithkline	ORAL
200MG Twice		Attack					
per day	1	YR		Nefazodone	C		
				Gabapentin	C		
				Diltiazem	C	Glaxosmithkline	
				Albuterol Inhaler	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4231745-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0417152A  
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	8	MON		Loestrin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231746-6Report Type:Periodic  
Age:43 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0417153A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Per day	2 WK	Chapped Lips		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN	15MG	Cheilitis		Buspirone	C		
times per day	Three	Dry Mouth					
UNKNOWN		Oral Pain		Lipitor	C		
UNKNOWN		Stomatitis		Toprol Xl	C		
UNKNOWN		Swollen Tongue		Hctz	C		
UNKNOWN		Tongue Ulceration		Amitriptyline	C		

Date:11/07/03ISR Number: 4231747-8Report Type:Periodic  
Age:48 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0417154A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3 WK	Tinnitus					
150MG Twice		Tremor		Wellbutrin	SS	Glaxosmithkline	ORAL
per day				Quinine	C		
				Alprazolam	C		

Date:11/07/03ISR Number: 4231749-1Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0417156A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Asthenopia		Wellbutrin	PS	Glaxosmithkline	ORAL

per day 1 MON Feeling Abnormal  
Nausea Birth Control C ORAL  
Photophobia  
Vomiting

Date:11/07/03ISR Number: 4231750-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0417157A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231751-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0417158A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cognitive Disorder Confusional State Disturbance In Attention Feeling Abnormal Memory Impairment Speech Disorder Developmental Thinking Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231752-1Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0417159A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Unknown		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
		Urticaria					

Date:11/07/03ISR Number: 4231753-3Report Type:Periodic  
Age:25 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0417248A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysphagia		Bupropion	PS	Glaxosmithkline	ORAL
150MG Twice		Joint Swelling					
per day		Musculoskeletal Stiffness					
		Pain					
		Stevens-Johnson Syndrome					
		Urticaria					

Date:11/07/03ISR Number: 4231757-0Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0417461A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Per day		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
		Urticaria					

Date:11/07/03ISR Number: 4231758-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0417479A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Twice		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3 WK	Urticaria					

Date:11/07/03ISR Number: 4231759-4Report Type:Periodic  
Age:27 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0417626A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	7 MON	Rash					

Date:11/07/03ISR Number: 4231760-0Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0417628A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
400MG per day		Therapeutic Response Decreased		Prilosec Blood Pressure Medication	C C		

Date:11/07/03ISR Number: 4231761-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0417649A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hyperglycaemia		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231762-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0417824A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Lymphadenopathy		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231763-6Report Type:Periodic  
Age:29 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0417843A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	DAY	Eye Movement Disorder					
		Nausea					

Date:11/07/03ISR Number: 4231765-XReport Type:Periodic  
Age: Gender:I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0417845A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Unknown		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231766-1Report Type:Periodic  
Age:63 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0417846A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Twice		Gastrooesophageal Reflux		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Disease					
				Klonopin	C		
				Trazodone	C		
				Prilosec	C		



Date:11/07/03ISR Number: 4231767-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0417847A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin Prozac	PS C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231768-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0417848A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Concerta	PS C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231769-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0417849A  
Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day	3 DAY	Disorientation					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231770-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0417850A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	1 WK	Abdominal Pain Upper Chest Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
		Dizziness Flatulence Gastrooesophageal Reflux Disease					

Date:11/07/03ISR Number: 4231771-5Report Type:Periodic  
Age:52 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0417864A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day	WK	Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN		Vertigo		Neurontin	SS		
				Tagamet	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4231772-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418008A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Blood Alcohol Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN				Prozac	SS		
7 HR				Wine	SS		ORAL

Date:11/07/03ISR Number: 4231773-9Report Type:Periodic  
Age:57 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418010A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
		Dysgeusia					
	WK	Energy Increased		Hctz	C		
				Accupril	C		
				Prevacid	C		
				Aspirin	C	Glaxosmithkline	
				Multivitamins	C		
				Activella	C		

Date:11/07/03ISR Number: 4231774-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0418050A  
 Age:46 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Rash		Bupropion	PS	Glaxosmithkline	ORAL
	1 MON			No Concurrent Medication	C		

Date:11/07/03ISR Number: 4231775-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0418051A  
 Age:50 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Myalgia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Zantac	C	Glaxosmithkline	
				Levothyroxine	C	Glaxosmithkline	

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Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231776-4Report Type:Periodic  
 Age:32 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418069A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	11 MON	Somnolence					
RESPIRATORY				Albuterol	C	Glaxosmithkline	
(INHALATION)							

Date:11/07/03ISR Number: 4231777-6Report Type:Periodic  
 Age:40 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418071A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG In the		Diarrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
morning		Disturbance In Attention					
		Dizziness		Prevacid	C		
		Fatigue		Caffeine	C		
		Feeling Abnormal					
		Paraesthesia					
		Temperature Intolerance					
		Tinnitus					
		Tremor					

Date:11/07/03ISR Number: 4231778-8Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418188A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Personality Change					

Date:11/07/03ISR Number: 4231779-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0418232A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 WK		Heart Rate Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
		Heart Rate Irregular Influenza Like Illness Tinnitus					

Date:11/07/03ISR Number: 4231780-6Report Type:Periodic  
Age:72 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0418253A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
4 WK		Abdominal Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
				Trileptal	C		
				Neurontin	C		
				Cozar	C		
				Toprol	C		
				Zyprexa	C		
				Aricept	C		

Date:11/07/03ISR Number: 4231781-8Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0418420A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
5 WK		Hypoaesthesia Oral		Wellbutrin	PS	Glaxosmithkline	ORAL
				No Concurrent			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Medication C

Date:11/07/03ISR Number: 4231784-3Report Type:Periodic  
Age:32 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418470A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Ineffective					
per day	2 WK	Dry Mouth		Vitamins	C		
		Insomnia					
		Somnolence					
		Weight Decreased					

Date:11/07/03ISR Number: 4231785-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0418601A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Abnormal Dreams					

Date:11/07/03ISR Number: 4231786-7Report Type:Periodic  
Age:51 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0418602A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1 WK	Photosensitivity Reaction					
		Rash		Testosterone	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4231787-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418603A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Transferrin Increased					

Date:11/07/03ISR Number: 4231788-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0418787A  
Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2	MON					

Date:11/07/03ISR Number: 4231789-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0418788A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Breast Tenderness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231790-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0418790A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oligomenorrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	MON			Effexor	C		
				Ritalin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231791-0Report Type:Periodic  
Age:69 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0418791A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	4 DAY	Asthenia Hyperhidrosis International Normalised Ratio Increased Muscle Spasms		Wellbutrin	PS	Glaxosmithkline	ORAL
				Avandia	C	Glaxosmithkline	
				Coumadin	C	Glaxosmithkline	
				Digoxin	C	Glaxosmithkline	
				Pravachol	C		
				Niaspan	C		
				Metformin	C		
				Zetia	C		
				Avapro	C		
				Amaryl	C		
				Prevacid	C		
				Coreg	C	Glaxosmithkline	
				Lantus	C		
				Endocet	C		

Date:11/07/03ISR Number: 4231792-2Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418799A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache Migraine Therapeutic Response Unexpected		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231793-4Report Type:Periodic  
Age:18 YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418801A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	YR	Vision Blurred		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:11/07/03ISR Number: 4231794-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0418826A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral		Wellbutrin Flexeril	PS C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231795-8Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0418956A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG In the morning	14 DAY			Methotrexate Vioxx Folic Acid Imitrex Prevacid Prozac Estrogen Testosterone	C C C C C C C C	Glaxosmithkline Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231797-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0418957A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231798-3Report Type:Periodic  
Age:51 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418958A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Single		Dysgeusia					
dose		Gingival Pain		Hydrocodone	C		
		Headache		Klonopin	C		
		Tooth Injury		Zantac	C	Glaxosmithkline	
		Toothache		Zocor	C		
				Baby Aspirin	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4231799-5Report Type:Periodic  
Age:81 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0419152A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
5 DAY				Neurontin	C		
				Remeron	C		
				Celexa	C		

Date:11/07/03ISR Number: 4231800-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0419168A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							

per day 3 MON

Adderall C  
Zelnorm C  
Nexium C  
Lexapro C

Date:11/07/03ISR Number: 4231801-0Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0419169A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown	2 WK	Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231802-2Report Type:Periodic  
Age:34 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0419170A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 WK		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231803-4Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0419172A

Outcome PT  
Crying  
Lip Ulceration

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Swelling Face

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
			Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231804-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0419173A  
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Unknown	7 MON			Nexium	SS		
UNKNOWN		16 MON					

Date:11/07/03ISR Number: 4231805-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0419174A  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Erythema					
per day	16 DAY	Eyelid Oedema					
		Inflammation					
		Ocular Hyperaemia					
		Swelling Face					
		Urticaria					

Date:11/07/03ISR Number: 4231806-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0419175A  
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	WK	Increased Appetite		Lotrel	C		
		Weight Increased					

Date:11/07/03ISR Number: 4231807-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0419176A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2	WK	Constipation	Wellbutrin	PS	Glaxosmithkline	ORAL
			Dizziness				
			Dry Mouth				
			Headache				
			Insomnia				
			Nausea				

Date:11/07/03ISR Number: 4231808-3Report Type:Periodic  
Age:60 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0419177A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Chest Pain	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	10	DAY	Dyspnoea				
			Paraesthesia	Verapamil	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231809-5Report Type:Periodic  
 Age:27 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0419181A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	30 DAY			Metabolife	C		

Date:11/07/03ISR Number: 4231810-1Report Type:Periodic  
 Age:44 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0419182A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG In the morning		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN	40MG In the morning			Celexa	C		

Date:11/07/03ISR Number: 4231811-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0419231A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG per day		Crying		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231814-9Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0419317A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Vaginismus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231815-0Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0419708A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3	WK					

Date:11/07/03ISR Number: 4231816-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0419709A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Jittery		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	2	WK					

Date:11/07/03ISR Number: 4231817-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0419710A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
YR							

Date:11/07/03ISR Number: 4231818-6Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0419711A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ageusia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	MON						

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231819-8Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0419712A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	2 MON		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Menorrhagia		Inderal	C		
				Tenormin	C		
				Contraceptive	C		ORAL

Date:11/07/03ISR Number: 4231820-4Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0419761A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG In the			Wellbutrin	PS	Glaxosmithkline	ORAL
Other	morning	2 WK					
		Mouth Injury					

Date:11/07/03ISR Number: 4231821-6Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0419917A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	1TAB Twice			Wellbutrin	PS	Glaxosmithkline	ORAL
	per day	7 DAY		Loestrin	C		
				Zantac	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4231822-8Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0419933A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Rash					



Date:11/07/03ISR Number: 4231823-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0419949A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/07/03ISR Number: 4231824-1Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0419977A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
1 WK							

Date:11/07/03ISR Number: 4231826-5Report Type:Periodic  
Age:20 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420005A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Postictal State					
150MG Twice							
per day	4 MON			Lexapro	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231827-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420007A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Keratoconjunctivitis Sicca		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231828-9Report Type:Periodic  
 Age:62 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420195A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	19	MON		Wellbutrin	PS	Glaxosmithkline	ORAL
25MG Per day	4	DAY		Zoloft	SS		
		Anxiety Arthritis Confusional State Dyskinesia Mental Status Changes Musculoskeletal Stiffness Nervousness Tension Vision Blurred					

Date:11/07/03ISR Number: 4231829-0Report Type:Periodic  
 Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420307A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Libido Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231830-7Report Type:Periodic  
 Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420314A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3	DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
		Cough Pain					

Pharyngolaryngeal Pain  
Rhinorrhoea  
Sneezing

Date:11/07/03ISR Number: 4231831-9Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420315A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	1	YR		Geodon	C		

Date:11/07/03ISR Number: 4231832-0Report Type:Periodic  
Age:19 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420527A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Joint Swelling		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Pyrexia					
200MG Per day	2	Serum Sickness					
		Urticaria					

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Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231833-2Report Type:Periodic  
Age:16 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420602A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Urine Amphetamine Positive		Wellbutrin	PS	Glaxosmithkline	ORAL
				Pseudoephedrine Hcl	SS	Glaxosmithkline	

Date:11/07/03ISR Number: 4231834-4Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420610A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG		Asthma		Wellbutrin	PS	Glaxosmithkline	ORAL
Variable dose	6 WK	Constipation Feeling Abnormal Suicidal Ideation		Advair	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4231835-6Report Type:Periodic  
Age:18 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420629A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 WK	Hypotrichosis		Wellbutrin	PS	Glaxosmithkline	ORAL
				Klonopin	C		

Date:11/07/03ISR Number: 4231836-8Report Type:Periodic  
Age:16 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0420632A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice per day	3 MON	Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

80MG Twice  
 per day  
 1MG As  
 required  
 150MG Twice  
 per day

Geodon C  
 Ativan C  
 Trileptal C

Date:11/07/03ISR Number: 4231837-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0420633A  
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Paraesthesia		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day				Birth Control	C		
				Amoxicillin	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4231838-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0420638A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							

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Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231839-3Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0420646A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200MG per day	3 WK	Dizziness Hyperhidrosis Overdose Palpitations		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231840-XReport Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420648A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	10 YR	Dizziness Malaise Nausea Pharmaceutical Product Complaint		Wellbutrin  Lipitor	PS  C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231841-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420760A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Unknown	3 MON	Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231842-3Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0420761A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Single		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL

dose  
Pallor  
Palpitations

Date:11/07/03ISR Number: 4231843-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0420762A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vaginal Haemorrhage		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				Premarin	SS		
UNKNOWN				Provera	SS		
UNKNOWN				Zoloft	C		
UNKNOWN	200MG Unknown			Buspar	C		
UNKNOWN	20MG Unknown						

Date:11/07/03ISR Number: 4231844-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0420763A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 YR	Foreign Body Trauma		Unspecified Medication	C		
		Insomnia					
		Pharmaceutical Product Complaint					

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Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231845-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420765A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
6	YR						

Date:11/07/03ISR Number: 4231846-0Report Type:Periodic  
 Age:48 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420809A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	5	DAY		Morphine Sulphate	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4231847-2Report Type:Periodic  
 Age:70 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420810A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3	YR					
		Myalgia					

Date:11/07/03ISR Number: 4231848-4Report Type:Periodic  
 Age:57 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420821A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1	WK		Monopril	C		
				Hydrochlorothiazide	C		



Date:11/07/03ISR Number: 4231849-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0420827A  
Age:6 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Retching		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	4	MON		Ritalin	C		

Date:11/07/03ISR Number: 4231850-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0420847A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ecchymosis		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4	DAY					

Date:11/07/03ISR Number: 4231851-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0420850A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day							

Date:11/07/03ISR Number: 4231852-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0420888A  
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

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Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231853-8Report Type:Periodic  
Age:49 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0420920A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	YR	Chest Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
50MG Per day	YR	Pain In Jaw		Wellbutrin	SS	Glaxosmithkline	ORAL
		Weight Increased		Xanax	C		

Date:11/07/03ISR Number: 4231854-XReport Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420960A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Chapped Lips		Wellbutrin	PS	Glaxosmithkline	ORAL
		Cheilitis		Seroquel	SS		
		Erythema		Celexa	SS		
		Lip Dry		Zoloft	SS		
		Overdose					

Date:11/07/03ISR Number: 4231855-1Report Type:Periodic  
Age:6 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420963A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
50MG Per day	4 MON	Retching		Wellbutrin	PS	Glaxosmithkline	ORAL
		Vomiting		Ritalin	C		

Date:11/07/03ISR Number: 4231856-3Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420975A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	1 MON	Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
		Vision Blurred					
				Estrogen Patch	C		

Date:11/07/03ISR Number: 4231857-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0421226A  
 Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231858-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0421244A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Transient Ischaemic		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Attack		Nortriptyline	C		
1TAB Twice				Ibuprofen	C	Glaxosmithkline	
per day							

Date:11/07/03ISR Number: 4231859-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0421256A  
 Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Panic Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

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Date:11/07/03ISR Number: 4231860-5Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0421259A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Elevated Mood Feeling Abnormal Feeling Jittery		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231861-7Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0421260A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Difficulty In Walking Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231862-9Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422161A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231863-0Report Type:Periodic  
 Age:40 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422164A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day	YR	Dry Mouth Tremor		Ritalin	C		

Date:11/07/03ISR Number: 4231864-2Report Type:Periodic  
 Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422178A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Chest Discomfort		Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Per day		Dyspnoea		Allegra Prometrium	C C		

Date:11/07/03ISR Number: 4231865-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0422180A  
 Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Swollen Tongue		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2 WK						

Date:11/07/03ISR Number: 4231866-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0422182A  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4 DAY			Xanax	C		
				Thyroid	C		
				Aciphex	C		
				Estrogen Cream	C		

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Date:11/07/03ISR Number: 4231867-8Report Type:Periodic  
 Age:25 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422190A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Migraine		Wellbutrin	PS	Glaxosmithkline	ORAL
3 YR				Oral Contraceptives	C		

Date:11/07/03ISR Number: 4231868-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422210A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2 WK						

Date:11/07/03ISR Number: 4231869-1Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422221A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Chest Discomfort					
per day	2 WK	Pruritus		Zoloft	C		
		Rash		Depakote	C		
		Restlessness					

Date:11/07/03ISR Number: 4231870-8Report Type:Periodic  
 Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422300A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	10 DAY	Insomnia		Klonopin	C		

Date:11/07/03ISR Number: 4231871-XReport Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422309A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Drug Ineffective	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3	MON	Increased Appetite				
			Libido Decreased				

Date:11/07/03ISR Number: 4231872-1Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422318A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Headache	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	7	MON		Cardura	C		
				Lisinopril	C		

Date:11/07/03ISR Number: 4231873-3Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422352A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Nausea	Wellbutrin	PS	Glaxosmithkline	

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Date:11/07/03ISR Number: 4231874-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422483A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accidental Overdose		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice per day							

Date:11/07/03ISR Number: 4231875-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422484A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphemia		Wellbutrin	PS	Glaxosmithkline	ORAL
3 MON							

Date:11/07/03ISR Number: 4231876-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422617A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day							
MON Pharmaceutical Product Complaint							

Date:11/07/03ISR Number: 4231877-0Report Type:Periodic  
Age: YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0422634A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
4 DAY							

Date:11/07/03ISR Number: 4231878-2Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422655A



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Screen Positive		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day				Zyprexa	C		

Date:11/07/03ISR Number: 4231879-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0422656A  
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day		Hypervigilance		Protonix	C		
		Insomnia		Guiafenesin-Dm	C		
		Onychorrhaxis					

Date:11/07/03ISR Number: 4231880-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0422682A  
 Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4 DAY	Impatience		Synthroid	C	Glaxosmithkline	
		Tremor		Reglan	C	Glaxosmithkline	
				Ambien	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231881-2Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0422687A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Twice		Therapeutic Response					
per day		Decreased					

Date:11/07/03ISR Number: 4231882-4Report Type:Periodic  
Age:16 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0422718A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -		Convulsion					
10 MON							
Initial or Prolonged							

Date:11/07/03ISR Number: 4231883-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422723A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Grand Mal Convulsion					
300MG Unknown 3 WK							

Date:11/07/03ISR Number: 4231884-8Report Type:Periodic  
Age:27 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422724A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
3 WK		Rash					
				Claritin	C		
				Albuterol	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4231885-XReport Type:Periodic  
Age:72 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0422730A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	6 DAY	Depression					
		Dyspnoea		Advair	C	Glaxosmithkline	
		Irritability		Combivent	C		
		Tremor					

Date:11/07/03ISR Number: 4231886-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0422745A  
 Age:53 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 MON						

Date:11/07/03ISR Number: 4231887-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0422746A  
 Age:41 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice		Pupils Unequal		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	5 DAY			Celexa	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231888-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422843A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3 MON	Anorexia Anxiety Asthenia Gastrointestinal Disorder Headache Nausea Weight Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231889-7Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0422845A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Belligerence		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231890-3Report Type:Periodic  
 Age:79 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422910A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Variable dose		Balance Disorder Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231891-5Report Type:Periodic  
 Age:27 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0422927A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1TAB Twice per day	2 WK	Dermatitis Allergic Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231896-4Report Type:Periodic  
Age:27 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422929A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1 DAY			Wellbutrin	PS	Glaxosmithkline	ORAL
		Headache					
		Heart Rate Irregular					
		Insomnia					

Date:11/07/03ISR Number: 4231897-6Report Type:Periodic  
Age:35 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0423092A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Per day	1 WK			Wellbutrin	PS	Glaxosmithkline	ORAL
		Abdominal Distension					
10MG At night	3 YR			Flexeril	C		ORAL
		Weight Increased					
200MG Per day	1 YR			Celebrex	C		ORAL

Date:11/07/03ISR Number: 4231898-8Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0423154A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening							
150MG Twice				Wellbutrin	PS	Glaxosmithkline	ORAL
Disability		Convulsion					
per day	12 DAY						
Other				Aspirin	C	Glaxosmithkline	
				Tums	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231899-XReport Type:Periodic  
 Age: YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0423369A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Anorexia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	5 DAY	Drug Ineffective					
3 YR		Headache		Paxil	C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231900-3Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0423412A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 WK	Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
		Rash Papular		Flax Seed Oil	C		
				Birth Control Pills	C		

Date:11/07/03ISR Number: 4231901-5Report Type:Periodic  
 Age:15 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0423418A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day		Drug Screen False		Wellbutrin	PS	Glaxosmithkline	ORAL
		Positive		None	C		

Date:11/07/03ISR Number: 4231902-7Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0423421A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
per day				Effexor Xr	C		

Xanax	C	
Lamictal	C	Glaxosmithkline
Synthroid	C	Glaxosmithkline

Date:11/07/03ISR Number: 4231903-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0423601A  
 Age:22 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Sexual Dysfunction		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3 YR						

Date:11/07/03ISR Number: 4231904-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0423620A  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Twice		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Feeling Abnormal					
				Effexor	C		
				Neurontin	C		
				Lamictal	C	Glaxosmithkline	
				Synthroid	C	Glaxosmithkline	
				Xanax Sr	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231905-2Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0423629A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Dry Mouth					
per day		Dysgeusia		Effexor	C		
		Hyperhidrosis		Triphasil	C		
		Tremor					

Date:11/07/03ISR Number: 4231906-4Report Type:Periodic  
Age:50 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0423633A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3 YR	Hypersensitivity		Buspar	C		
		Rash		Sonata	C		
		Rash Pruritic		Ambien	C		
UNKNOWN				Ativan	C		
UNKNOWN				Klonopin	C		
UNKNOWN				Glucophage	C		
UNKNOWN	1G Twice per						
day				Actos	C		
UNKNOWN	30MG Per day			Tricor	C		
UNKNOWN	160MG Per day			Lipitor	C		
UNKNOWN	10MG Per day			Levaquin	C		OTHER
				Flagyl	C	Glaxosmithkline	OTHER

Date:11/07/03ISR Number: 4231907-6Report Type:Periodic  
Age:25 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0423635A



Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	6	Gastrooesophageal Reflux Disease		Risperdal	C		
		Muscle Twitching		Ativan	C		
				Eskalith	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4231915-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0423698A  
 Age: Gender: I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231916-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0423709A  
 Age:20 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Benign Soft Tissue		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Neoplasm					
150MG Per day	1	Dizziness		Effexor Xr	C		ORAL
		Painful Response To Normal Stimuli					
		Skin Nodule					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231917-9Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0423713A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Three times per day	3 YR	Cough		Wellbutrin	PS	Glaxosmithkline	ORAL
				Clonazepam	C		
				Primatene Mist	C		

Date:11/07/03ISR Number: 4231918-0Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0423717A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Activities Of Daily Living Impaired		Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Ineffective Emotional Distress		Neurontin	C		

Date:11/07/03ISR Number: 4231922-2Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0423720A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75MG Per day	3 YR	Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
				Lexapro	C		
				Toprol	C		

Date:11/07/03ISR Number: 4231923-4Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0423740A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Pain Nausea		Wellbutrin	PS	Glaxosmithkline	

Date:11/07/03ISR Number: 4231924-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0423839A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Irritability		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231925-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0423840A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231926-XReport Type:Periodic  
Age:20 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0423982A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other	300MG Per day 9 WK						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231927-1Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0423999A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	4 DAY	Dry Mouth Flushing Headache		Wellbutrin  Amitriptyline	PS  C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231928-3Report Type:Periodic  
Age:73 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0424013A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin Testomax	PS SS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231929-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424031A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Chest Discomfort Yawning		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231930-1Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424098A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day	8 DAY	Anorgasmia		Wellbutrin  Birth Control Pills	PS  C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231931-3Report Type:Periodic  
Age:76 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424099A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day				Lipitor	C		
				Zoloft	C		
				Vitamins	C		

Date:11/07/03ISR Number: 4231932-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0424100A  
 Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Myalgia		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day							
				Synthroid	C	Glaxosmithkline	
				Toprol	C		
				Lotrel	C		

Date:11/07/03ISR Number: 4231933-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0424112A  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Social Avoidant Behaviour		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	21 MON			Amitriptyline	C		
				Seroquel	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Fioricet C

Date:11/07/03ISR Number: 4231934-9Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424118A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Platelet Count Abnormal					
200MG Per day		Platelet Function Test Abnormal		Ambien	C		

Date:11/07/03ISR Number: 4231935-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424127A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Depression					
per day		Drug Ineffective					
		Stress					

Date:11/07/03ISR Number: 4231936-2Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424141A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Alopecia					
per day							

Date:11/07/03ISR Number: 4231937-4Report Type:Periodic  
Age:37 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0424301A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

100MG Unknown 1 YR Abdominal Pain Wellbutrin PS Glaxosmithkline ORAL  
Diarrhoea  
Nausea

Date:11/07/03ISR Number: 4231938-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0424305A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoaesthesia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Lip Dry					
per day	1 YR			Zestril	C		

Date:11/07/03ISR Number: 4231941-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0424336A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	26 WK			Weight Loss Medication	C		
				Multi Vitamin + Minerals	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231942-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424377A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Distractibility Euphoric Mood Headache Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231943-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424519A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		Liver Function Test Abnormal		Wellbutrin Adderall Ortho Tri Cyclen	PS C C	Glaxosmithkline	

Date:11/07/03ISR Number: 4231944-1Report Type:Periodic  
Age: YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0424523A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231945-3Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424688A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Dizziness Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231946-5Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424698A



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	1 WK	Apathy		Paxil Cr	C	Glaxosmithkline	
		Asthenia		Naproxen	C		

Date:11/07/03ISR Number: 4231947-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0424721A  
 Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK	Insomnia					
		Psychomotor Hyperactivity					

Date:11/07/03ISR Number: 4231948-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0424739A  
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG		Liver Function Test					
Variable dose		Abnormal		Avandamet	SS	Glaxosmithkline	ORAL
4MG Per day		Memory Impairment		Aspirin	C	Glaxosmithkline	
		Therapeutic Response		Prozac	C		
		Unexpected		Lipitor	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Ativan C  
Tetracycline C

Date:11/07/03ISR Number: 4231949-0Report Type:Periodic  
Age:74 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0424742A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Abnormal Behaviour		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 YR	Tinnitus					

Date:11/07/03ISR Number: 4231950-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424789A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Level Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
				Elavil	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4231951-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424875A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day		Muscle Twitching		Wellbutrin	SS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231952-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424876A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

Muscle Twitching

Wellbutrin SS Glaxosmithkline ORAL

100MG Per day

Date:11/07/03ISR Number: 4231953-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0424950A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	5 YR						

Date:11/07/03ISR Number: 4231954-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0425034A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK						

Date:11/07/03ISR Number: 4231955-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0425050A  
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hangover		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK			Adipex	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231956-8Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425067A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	7 DAY	Dizziness Dry Mouth Feeling Jittery		Wellbutrin	PS	Glaxosmithkline	

Date:11/07/03ISR Number: 4231957-XReport Type:Periodic  
 Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425082A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1 DAY	Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
				Dapson	C		

Date:11/07/03ISR Number: 4231958-1Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0425087A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 per day	3 WK	Drug Interaction Headache Somnolence Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
				Xanax	SS		
				Alcohol	SS		ORAL

Date:11/07/03ISR Number: 4231959-3Report Type:Periodic  
 Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425133A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Anxiety Headache Paraesthesia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Synthroid	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4231960-XReport Type:Periodic  
Age:52 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425163A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice		Fatigue					
per day	4 MON	Insomnia		Synthroid	C	Glaxosmithkline	
		Neuropathy		Reglan	C	Glaxosmithkline	
		Sinusitis		Celebrex	C		
		Weight Decreased		Zyprexa	C		
				Prevacid	C		
				Allegra	C		

Date:11/07/03ISR Number: 4231961-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425362A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Energy Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Fatigue					
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231962-3Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425516A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2	WK	Abdominal Pain Upper	Wellbutrin	PS	Glaxosmithkline	ORAL
			Headache				

Date:11/07/03ISR Number: 4231963-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425583A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Tremor	Bupropion	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231964-7Report Type:Periodic  
Age:86 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0425675A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Per day	24	DAY	Anxiety	Wellbutrin	PS	Glaxosmithkline	ORAL
			Drug Ineffective	Plendil	C		
UNKNOWN			Tremor	Atenolol	C		
				Zestril	C		

Date:11/07/03ISR Number: 4231965-9Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425683A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Unknown	1	DAY		Dilantin	C		

Date:11/07/03ISR Number: 4231967-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0425696A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 WK		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231968-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0425710A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Arthralgia Myalgia Pain Pain In Extremity		Wellbutrin  Synthroid Atacand Atenolol Lipitor Singulair	PS  C C C C C	Glaxosmithkline  Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231969-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0425926A  
 Age:42 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - per day Initial or Prolonged	60 WK	Thrombocytopenia		Wellbutrin  Effexor Xr Adderall Xr	PS  C C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231970-2Report Type:Periodic  
Age:60 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425944A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1	YR					
				Duragesic	C		
				Neurontin	C		
				Xanax	C		
				Tylenol #3	C		

Date:11/07/03ISR Number: 4231971-4Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425949A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -							
Initial or Prolonged							

Date:11/07/03ISR Number: 4231972-6Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425972A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1	MON					
				Ultram	C		
				Vicodin	C		

Date:11/07/03ISR Number: 4231973-8Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425973A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Memory Impairment		Wellbutrin	PS	Glaxosmithkline	ORAL
1 per day	1	WK					



Sluggishness  
Stress

Serzone C  
Haldol C  
Depakote C  
Lipitor C  
Glucotrol C

Date:11/07/03ISR Number: 4231974-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425979A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
WK							

Date:11/07/03ISR Number: 4231975-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425980A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Dyspepsia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Nausea					
		Vomiting					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231976-3Report Type:Periodic  
Age:60 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426108A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	9	MON		Proscar	C		
				Fosamax	C		
				Prinivil	C		

Date:11/07/03ISR Number: 4231977-5Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426123A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 YR		Gynaecomastia		Wellbutrin	PS	Glaxosmithkline	

Date:11/07/03ISR Number: 4231978-7Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426136A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1	MON		Vit. B	C		
				Vit C	C	Glaxosmithkline	
				Vit E	C		
				Flax Seed Oil	C		

Date:11/07/03ISR Number: 4231979-9Report Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426138A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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150MG Twice	Amnesia	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	Asthenia				
	Cognitive Disorder	Protonix	C		
	Dissociation	Zyban	C	Glaxosmithkline	ORAL
	Drug Ineffective				
	Dyspepsia				
	Early Morning Awakening				
	Fatigue				
	Muscle Twitching				
	Nausea				
	Reading Disorder				
	Sleep Disorder				

Date:11/07/03ISR Number: 4231987-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0426186A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Liver Function Test		Wellbutrin	PS	Glaxosmithkline	ORAL
		Abnormal		Strattera	SS		
UNKNOWN							

Date:11/07/03ISR Number: 4231988-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0426452A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
per day							

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Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231989-1Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0426572A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day							

Date:11/07/03ISR Number: 4231990-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426577A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vulvovaginal Dryness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231991-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426578A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vulvovaginal Dryness		Wellbutrin Depakote	PS C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231992-1Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426587A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Libido Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 1 YR							

Date:11/07/03ISR Number: 4231993-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426594A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

100MG Twice		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2	MON					
				Celexa	C		
				Calcium	C		

Date:11/07/03ISR Number: 4231994-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0426599A  
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day	1	MON					
				Eye Drop	C		
				Topical Cream	C		

Date:11/07/03ISR Number: 4231995-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0426619A  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3	WK					
		Tension					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231996-9Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0426882A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:11/07/03ISR Number: 4231997-0Report Type:Periodic  
Age:55 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0426885A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day 3 DAY							
				Lexapro	C		
				Geodon	C		

Date:11/07/03ISR Number: 4231998-2Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426893A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4231999-4Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426902A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day 3 YR							

Date:11/07/03ISR Number: 4232000-9Report Type:Periodic  
Age:10 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426924A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	7 DAY			Effexor	C		

Date:11/07/03ISR Number: 4232001-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0426925A  
 Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Celexa	C		
per day	3 YR			Synthroid	C	Glaxosmithkline	
				Oral Contraceptives	C		
				Eye Drops	C		

Date:11/07/03ISR Number: 4232002-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0426935A  
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Palpitations		Zoloft	C		
per day	5 YR			Doxepin	C		
				Chlorazepate			
				Dipotassium	C		
				Geodon	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Premarin	C	
Vit E	C	
Vit C	C	Glaxosmithkline
Calcium	C	
Vit D	C	
Multivitamin	C	
Evening Primrose Oil	C	

Date:11/07/03ISR Number: 4232003-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0426938A  
 Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice per day		Anxiety					
		Disturbance In Attention		Lithium	C	Glaxosmithkline	
		Insomnia		6mp	C	Glaxosmithkline	
				Blood Pressure Medication	C		
				Unspecified Medication	C		
				Paxil	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4232004-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0427014A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pruritus Generalised		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4232005-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0427015A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:11/07/03ISR Number: 4232006-XReport Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427016A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 YR	Flushing		Paxil	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4232007-1Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427019A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Feeling Jittery					
per day	4 WK	Nausea					

Date:11/07/03ISR Number: 4232008-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427032A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Irritability		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Tinnitus					
per day	1 YR						

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4232009-5Report Type:Periodic  
 Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427035A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Male Sexual Dysfunction		Wellbutrin	PS	Glaxosmithkline	

Date:11/07/03ISR Number: 4232010-1Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427054A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Road Traffic Accident		Methadone	C	Glaxosmithkline	
5 WK							

Date:11/07/03ISR Number: 4232011-3Report Type:Periodic  
 Age:34 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427055A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day	7 MON			Zoloft	C		

Date:11/07/03ISR Number: 4232012-5Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427058A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Nausea					
per day	4 MON						

Date:11/07/03ISR Number: 4232013-7Report Type:Periodic  
 Age:50 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0427084A

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG	Per day 19 DAY						

Date:11/07/03ISR Number: 4232014-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0427216A  
 Age:32 YR Gender:Male I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Twice per day						

Date:11/07/03ISR Number: 4232015-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0427336A  
 Age: YR Gender:Male I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Per day 1 WK			Zyprexa	C		

Date:11/07/03ISR Number: 4232016-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0427349A  
 Age:43 YR Gender:Male I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
75MG	Twice per day 3 MON			Lamictal	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4232017-4Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427350A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 WK		Anxiety Crying Irritability Nicotine Dependence		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4232018-6Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427359A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	13 WK	Dry Mouth Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4232019-8Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427369A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 MON	Dehydration Dizziness Palpitations		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4232020-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0427391A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
4 YR		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4232021-6Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427403A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4232022-8Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427512A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Visual Acuity Reduced		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	5	MON		Estratest	C		
				Prilosec	C		
				Soma	C		
				Vicodin	C		
				Ibuprofen	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4232024-1Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427514A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blister		Wellbutrin	PS	Glaxosmithkline	ORAL
3	WK	Pain		None	C		
		Pain In Extremity					
		Pruritus					
		Tongue Blistering					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4232025-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A042753A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4232026-5Report Type:Periodic  
Age:50 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0427679A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day 10 DAY Other		Angioneurotic Oedema Swelling Face		Wellbutrin	PS	Glaxosmithkline	ORAL
				Nicotrol Inhaler	SS	Glaxosmithkline	
				Monopril	C		
				Allopurinol	C	Glaxosmithkline	
				Norvasc	C		
				Prevacid	C		
				Lipitor	C		
				Xanax	C		
.25MG Weekly				Advair	C	Glaxosmithkline	
				Combivent	C		
2PUFF Four times per day				Ecotrin	C	Glaxosmithkline	
325MG Per day							

Date:11/07/03ISR Number: 4232027-7Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427694A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fungal Infection		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 6 MON				Aciphex	C		
				Vioxx	C		
				Flexeril	C		
				Hydrochlorothiazide	C		

Date:11/07/03ISR Number: 4232028-9Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0427703A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Renal Impairment		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4232029-0Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0427726A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4232030-7Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427844A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Weight Decreased					
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4232031-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427869A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	8 DAY						

Date:11/07/03ISR Number: 4232032-0Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427878A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Muscle Tightness					
per day	4 DAY						

Date:11/07/03ISR Number: 4232033-2Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427898A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Drunk		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4 DAY						

Date:11/07/03ISR Number: 4232034-4Report Type:Periodic  
Age: YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0427902A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							

Date:11/07/03ISR Number: 4232035-6Report Type:Periodic  
Age:14 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428051A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



100MG Twice  
per day 3 MON  
Anger  
Irritability

Wellbutrin PS Glaxosmithkline ORAL

Date:11/07/03ISR Number: 4232036-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0428052A  
Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	9 DAY	Mood Swings		Vitamins	C		
		Weight Decreased					

Date:11/07/03ISR Number: 4232037-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0428057A  
Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Feeling Abnormal					
per day	2 WK	Insomnia					
		Joint Stiffness					
		Muscle Tightness					
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4232038-1Report Type:Periodic  
Age:17 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0428093A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4232039-3Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0428109A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4232040-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428115A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4232047-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0432608A  
Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
1 MON		Nightmare		Bupropion	SS	Glaxosmithkline	
		Urinary Retention		Heart Medication	C		
				Antihypertensives	C		
				Lopressor	C		

Date:11/07/03ISR Number: 4232048-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0432624A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Date:11/07/03ISR Number: 4232054-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0296280A

Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Blood Creatine	Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	3 WK		Phosphokinase Increased	Soprol	C		ORAL
Initial or Prolonged							
.5TAB Per day							
Other			Dermatomyositis	Microval	C		ORAL
			Erythema				
			Myalgia				
			Skin Ulcer				

Date:11/07/03ISR Number: 4232057-5Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0309991A

Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening			Face Injury	Zyban	PS	Glaxosmithkline	ORAL
1TAB Per day	13 DAY		Facial Bones Fracture	Atorvastatin	C		ORAL
Other							
10MG Per day			Grand Mal Convulsion	Ketotifen	C		ORAL
2MG Per day							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4232060-5Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0311375A  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anorexia		Zyban	PS	Glaxosmithkline	ORAL
3	MON						
		Feeling Abnormal					
		Insomnia					
		Palpitations					
		Tremor					
		Weight Decreased					

Date:11/07/03ISR Number: 4232061-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0311639A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Back Pain		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	3 DAY						
		Cardio-Respiratory Arrest					
		Myocardial Infarction					
		Nausea					
		Sudden Death					

Date:11/07/03ISR Number: 4232062-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0311918A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain		Zyban	PS	Glaxosmithkline	ORAL
29	DAY						
		Chest X-Ray Abnormal					
		Constipation					
		Hepatic Enzyme Increased					
		Hepatic Neoplasm					
		Hepatic Pain					
		Macrocytosis					
		Nausea					
		Vomiting					
		Weight Decreased					

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Bupropion	PS	Glaxosmithkline	
UNKNOWN		Sleep Walking					

Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	18 MON	Lethargy					
				Premarin	C		
				Synthroid	C	Glaxosmithkline	
				Cholesterol Reducing			
				Agent Name Not Known	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233288-0Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400181A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice				Wellbutrin	PS	Glaxosmithkline	ORAL
per day		WK					
75MG Twice				Wellbutrin	SS	Glaxosmithkline	ORAL
per day		WK					

Date:11/07/03ISR Number: 4233289-2Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400193A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other				Bupropion	PS	Glaxosmithkline	ORAL
				Prozac	C		

Date:11/07/03ISR Number: 4233290-9Report Type:Periodic  
 Age:70 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400207A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Three				Wellbutrin	PS	Glaxosmithkline	ORAL
times per day 5		YR					
				Multivitamin	C		
				Zocor	C		
				Atenolol	C		

Date:11/07/03ISR Number: 4233291-0Report Type:Periodic  
 Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400247A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Twice	Headache	Wellbutrin	PS	Glaxosmithkline	ORAL
per day					
		Cenestin	C		
		Albuterol	C	Glaxosmithkline	
		Norvasc	C		
		Celebrex	C		
		Singulair	C		
		Maxzide	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4233292-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0400248A  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Feeling Jittery		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 YR			Caffeine	C		ORAL
6PACK Per day							

Date:11/07/03ISR Number: 4233293-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0400275A  
 Age:12 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
1 YR							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233294-6Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0400277A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
6 WK		Palpitations Pollakiuria Tachycardia					

Date:11/07/03ISR Number: 4233295-8Report Type:Periodic  
Age: Gender:I I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400400A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Liver Function Test Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
				Depakote	C		
				Zyprexa	C		
				Remeron	C		

Date:11/07/03ISR Number: 4233296-XReport Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400438A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 YR	Drug Interaction					
UNKNOWN				Delsym	SS	Glaxosmithkline	
				No Concurrent Medication	C		

Date:11/07/03ISR Number: 4233297-1Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400576A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



150MG Per day	MON	Food Craving	Wellbutrin	PS	Glaxosmithkline	ORAL
		Increased Appetite	Celexa	C		
		Weight Increased	Thyrolar	C		

Date:11/07/03ISR Number: 4233298-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0400581A  
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Dizziness					
per day	45 DAY	Dyspnoea					
		Insomnia					

Date:11/07/03ISR Number: 4233299-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0400603A  
 Age:24 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	6 MON	Muscle Twitching		Flonase	C	Glaxosmithkline	NASAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233300-9Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400622A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Distension		Wellbutrin	PS	Glaxosmithkline	ORAL
5 DAY		Anxiety Irritability Paranoia					

Date:11/07/03ISR Number: 4233301-0Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400668A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Twice		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	4 MON			Zoloft Lithobid Enema	C C C	Glaxosmithkline	

Date:11/07/03ISR Number: 4233302-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400669A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Unknown		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233303-4Report Type:Periodic  
Age:41 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400717A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Twice		Abnormal Sensation In Eye		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1 MON	Corneal Reflex Decreased					

Dehydration  
Increased Insulin  
Requirement  
Insomnia

Insulin C  
Ambien C  
Valium C  
Xanax C  
Alcohol C

ORAL

Date:11/07/03ISR Number: 4233304-6Report Type:Periodic  
Age:14 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0400721A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	536 DAY	Convulsion Decreased Appetite Nausea		Wellbutrin  Concerta Verapamil Zyrtec-D Nasonex Advair	PS  C C C C C	Glaxosmithkline    Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233305-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400727A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 150MG Per day	5 YR	Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233306-XReport Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400776A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Vomiting		Premarin	C		

Date:11/07/03ISR Number: 4233307-1Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400780A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG per day	3 WK	Headache		Paxil Cr	C	Glaxosmithkline	
		Social Avoidant Behaviour		Xanax	C		

Date:11/07/03ISR Number: 4233308-3Report Type:Periodic  
Age:2 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400781A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accidental Exposure		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - 300MG Single Initial or Prolonged dose	1 DAY	Tachycardia					
1 DAY				Diet Medication	SS		ORAL

Date:11/07/03ISR Number: 4233309-5Report Type:Periodic  
Age:41 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0400966A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day	2 WK	Anger					

3CAP Twice			Dry Mouth	Fat Burner	SS	ORAL
per day	1	DAY	Dyspnoea			
			Energy Increased	Stamina Rx	SS	ORAL
			Irritability			
			Nervousness			
			Palpitations			
			Tachycardia			
			Tremor			

Date:11/07/03ISR Number: 4233310-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0401020A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233311-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0401029A  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Therapeutic Response		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Decreased					
per day	2	YR					



per day

Date:11/07/03ISR Number: 4233316-2Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0401286A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Unknown	11 DAY	Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
		Fatigue					
		Nausea					

Date:11/07/03ISR Number: 4233317-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0401328A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233318-6Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0401506A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Unknown	19 DAY	Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
		Urticaria		Prednisone	C		
				Fiorinal	C		
				Vitamins	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233319-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0401524A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
4 MON		Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
		Hunger		Klonopin	C		
		Increased Appetite					
		Nightmare					
		Parosmia					
		Polydipsia					
		Restless Legs Syndrome					
		Smoker					

Date:11/07/03ISR Number: 4233320-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0401539A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypersensitivity		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233321-6Report Type:Periodic  
 Age:16 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0401549A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 DAY		Rash Macular		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233322-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0401805A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Transmission Of Drug Via Semen		Wellbutrin	PS	Glaxosmithkline	OTHER
		Vaginal Infection		Birth Control	C		



Date:11/07/03ISR Number: 4233323-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0401814A  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
15 DAY				Birth Control Pills	C		

Date:11/07/03ISR Number: 4233324-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0401821A  
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	4	MON					
80MG per day				Prozac	C		
5MG per day				Zyprexa	C		

Date:11/07/03ISR Number: 4233325-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0401827A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dermatitis Allergic		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233326-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0401831A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 MON	Feeling Jittery		Wellbutrin	PS	Glaxosmithkline	ORAL
				Strattera	SS		

Date:11/07/03ISR Number: 4233327-7Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402003A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Neurontin	C		
				Lexapro	C		

Date:11/07/03ISR Number: 4233328-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402006A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Pancytopenia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233329-0Report Type:Periodic  
Age:65 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0402007A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Per day		Musculoskeletal Stiffness		Wellbutrin	PS	Glaxosmithkline	ORAL
				Tramadol	C		
				Lotensin	C	Glaxosmithkline	
				Vicodin	C		
				Synthroid	C	Glaxosmithkline	
				Toprol	C		

Date:11/07/03ISR Number: 4233330-7Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402008A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
3 DAY		Swollen Tongue		Prometrium	C		
				Clonazepam	C		

Date:11/07/03ISR Number: 4233331-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402013A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day							

Date:11/07/03ISR Number: 4233332-0Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402026A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day				Alprazolam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233333-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402041A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Bruxism		Wellbutrin	PS	Glaxosmithkline	ORAL
		Facial Pain		Prozac	SS		
UNKNOWN		Nervousness					

Date:11/07/03ISR Number: 4233334-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402168A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day		Weight Decreased		Celexa	C		
UNKNOWN	10MG Per day						

Date:11/07/03ISR Number: 4233335-6Report Type:Periodic  
Age:58 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0402173A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Eye Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Hypoaesthesia					
per day	4 MON	Keratoconjunctivitis		Atenolol	C		
		Sicca		Ambien	C		
UNKNOWN		Vision Blurred		Ketoprofen	C		
UNKNOWN				Methocarbamol	C		
UNKNOWN				Imitrex	C	Glaxosmithkline	
UNKNOWN				Meclizine	C		

Date:11/07/03ISR Number: 4233336-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0402176A  
Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	23	DAY		Risperidal	C		
				Ativan	C		
				Neurontin	C		

Date:11/07/03ISR Number: 4233337-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0402207A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Muscle Tightness					
per day	5	MON		Anfebutamona	SS	Glaxosmithkline	
		Nervousness					

Date:11/07/03ISR Number: 4233338-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0402279A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Memory Impairment		Wellbutrin	PS	Glaxosmithkline	ORAL
375MG per day	2	YR					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233339-3Report Type:Periodic  
Age:2 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0402291A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accidental Exposure		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233340-XReport Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402305A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Skin Odour Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice		Weight Decreased					
per day	1 YR			Advil	C	Glaxosmithkline	
				Midol	C		

Date:11/07/03ISR Number: 4233341-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402362A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lethargy		Wellbutrin	PS	Glaxosmithkline	ORAL
				Ssri	C		

Date:11/07/03ISR Number: 4233342-3Report Type:Periodic  
Age:48 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0402414A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG In the		Vomiting					
morning	11 YR			Trazodone	C		
				Relafen	C	Glaxosmithkline	
				Birth Control	C		
				Prilosec	C		
				Vitamins	C		

Date:11/07/03ISR Number: 4233343-5Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402431A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Prozac	C		
				Topamax	C		

Date:11/07/03ISR Number: 4233344-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402541A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
3 WK		Dizziness					
		Insomnia					

Date:11/07/03ISR Number: 4233345-9Report Type:Periodic  
Age:71 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0402543A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Essential Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	12 YR						

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

5MG Twice per day	Hctz	C	ORAL
	Hytrin	C	ORAL
50MG Twice per day	Toprol	C	ORAL
	Cozaar	C	ORAL
	Norvasc	C	ORAL
	Wine	C	

Date:11/07/03ISR Number: 4233346-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0402567A  
 Age:32 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice Initial or Prolonged per day	Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233347-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0402569A  
 Age:40 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration 91 DAY	Blood Creatine Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
			Risperdal	C		

Date:11/07/03ISR Number: 4233348-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0402571A  
 Age:53 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration 150MG Per day 18 DAY	Abdominal Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
	Diarrhoea		Methadone	C	Glaxosmithkline	
	Headache		Hydrocodone	C		



Nausea

Excedrin

C

Date:11/07/03ISR Number: 4233349-6Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402589A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY						

Date:11/07/03ISR Number: 4233350-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402591A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Medication Error					
per day							

Date:11/07/03ISR Number: 4233351-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0402593A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menstruation Irregular		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Oral Contraceptive Pills	C		
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233352-6Report Type:Periodic  
Age:49 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0402600A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	20 DAY	Tinnitus Yawning		Wellbutrin	PS	Glaxosmithkline	ORAL
				Blood Pressure Medication	C		

Date:11/07/03ISR Number: 4233353-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402660A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233354-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402666A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day		Grand Mal Convulsion Intentional Misuse		Wellbutrin	PS	Glaxosmithkline	ORAL
				Strattera	SS		

Date:11/07/03ISR Number: 4233355-1Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0402672A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Unknown		Pharmaceutical Product Complaint Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233356-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402677A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Rash					
per day		Urticaria					

Date:11/07/03ISR Number: 4233357-5Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402894A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
400MG Per day 3	MON	Crying		Estrogen	C		
INTRADERMAL		Depressed Mood		Klonopin	C		
		Irritability		Zyrtec	C	Glaxosmithkline	
		Palpitations					

Date:11/07/03ISR Number: 4233358-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0402896A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Burning Sensation		Wellbutrin	PS	Glaxosmithkline	ORAL
		Dermatitis Exfoliative		Prevacid	C		
		Rash		Levaquin	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Advair	C	Glaxosmithkline
Solumedrol	C	
Atrovent	C	
Imdur	C	
Klonopin	C	
Prednisone	C	
Baby Aspirin	C	Glaxosmithkline
Albuterol	C	Glaxosmithkline

Date:11/07/03ISR Number: 4233359-9Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0402904A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Contusion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Fall					
per day	3	MON					
60MG Per day	973	DAY	Peroneal Nerve Palsy	Prozac	SS		ORAL

Date:11/07/03ISR Number: 4233360-5Report Type:Periodic  
 Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0403048A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Screen False		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Positive					
per day	1	YR					
				Paxil	C	Glaxosmithkline	
				Depakote	C		
				Seroquel	C		

Date:11/07/03ISR Number: 4233361-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0403091A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		DAY					

UNKNOWN                      Diarrhoea                      Prozac                      C  
 20MG Unknown  
 Feeling Abnormal  
 Headache  
 Insomnia  
 Nausea

Date:11/07/03ISR Number: 4233362-9Report Type:Periodic                      Company Report #US-GLAXOSMITHKLINE-A0403098A  
 Age:                      Gender:                      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233363-0Report Type:Periodic                      Company Report #US-GLAXOSMITHKLINE-A0403105A  
 Age:26 YR                      Gender:Female                      I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Generalised		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Advair	C	Glaxosmithkline	
per day				Albuterol	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233364-2Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0403109A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Polymyalgia		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day	3 WK			Valproic Acid	C		

Date:11/07/03ISR Number: 4233365-4Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0403118A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Cheilitis					
per day	1 MON	Dysgeusia					
		Flushing					
		Gingival Pain					
		Hypersensitivity					
		Nervousness					
		Pruritus					
		Stomatitis					

Date:11/07/03ISR Number: 4233366-6Report Type:Periodic  
Age:24 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0403313A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diplopia		Wellbutrin	PS	Glaxosmithkline	ORAL
75MG Three		Keratoconjunctivitis					
times per day	2 WK	Sicca					
		Photophobia					

Date:11/07/03ISR Number: 4233367-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0403314A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233368-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0403425A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Somnolence		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233369-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0403441A  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day	3 MON	Headache					

Date:11/07/03ISR Number: 4233370-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0403495A  
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Rash		Wellbutrin	SS	Glaxosmithkline	ORAL
100MG Per day	1 DAY						
5 DAY							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Remeron	C
Prozac	C
Xanax	C
Seroquel	C

Date:11/07/03ISR Number: 4233371-XReport Type:Periodic  
 Age:58 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0403566A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Dizziness					
per day	3	DAY					
		Dry Mouth					
		Tinnitus					

Date:11/07/03ISR Number: 4233372-1Report Type:Periodic  
 Age:22 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0403680A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	42	DAY		Insulin	C		

Date:11/07/03ISR Number: 4233373-3Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0403687A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accidental Overdose		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion		Concerta	C		

Date:11/07/03ISR Number: 4233374-5Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0403734A



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233375-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0403736A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Screen False		Wellbutrin	PS	Glaxosmithkline	ORAL
		Positive		Paxil	SS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233376-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0403749A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day							

Date:11/07/03ISR Number: 4233377-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0403781A  
Age:42 YR Gender:Female I/FU:I

Outcome	PT
	Drug Ineffective
	Energy Increased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Feeling Jittery Insomnia Nervousness	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	6 MON	Therapeutic Response Unexpected		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233378-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0403786A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
350MG Unknown		Vision Blurred Weight Increased		Wellbutrin Concurrent Medications	PS C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233379-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0404002A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Per day	DAY	Disturbance In Attention Dizziness Dry Mouth Food Craving Insomnia Judgement Impaired Somnolence		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233380-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0404042A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
2 YR		Depression Drug Ineffective		Wellbutrin Hormone	PS C	Glaxosmithkline	ORAL

Fatigue  
Laziness

Date:11/07/03ISR Number: 4233381-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0404142A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anaphylactic Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Pruritus					
per day		Rash					
		Urticaria					

Date:11/07/03ISR Number: 4233382-4Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0404166A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oligomenorrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
400MG per day	2 MON			Estrogen Patch	C		
				Klonopin	C		



150MG Twice		Thyroxine Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2	WK		Motrin	C	Glaxosmithkline	
				Levothyroxine	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4233388-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0404375A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233389-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0404379A  
 Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Screen Positive		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day	3	YR		Imitrex	C	Glaxosmithkline	
				Flonase	C	Glaxosmithkline	NASAL
				Celexa	C		
				Allegra	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233390-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0404387A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Allodynia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233391-5Report Type:Periodic  
Age:31 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0404406A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day	3 WK	Headache Musculoskeletal Stiffness Neck Pain Nightmare		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233392-7Report Type:Periodic  
Age:59 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0404407A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG At night	DAY	Impaired Work Ability Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
75MG Per day				Plavix	C		ORAL
10MG Per day				Potassium	C		ORAL
10MG Unknown				Prednisone	C		ORAL
.75U Per day				Synthroid	C	Glaxosmithkline	ORAL
100MG Twice per day				Toprol	C		ORAL
RESPIRATORY (INHALATION)	1PUFF Twice			Advair	C	Glaxosmithkline	

per day

Date:11/07/03ISR Number: 4233393-9Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0404408A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Stool Analysis Abnormal	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	MON			Zoloft	C		

Date:11/07/03ISR Number: 4233394-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0404439A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Stool Analysis Abnormal	Wellbutrin	PS	Glaxosmithkline	ORAL
6 MON				Prevacid	C		
				Prempro	C		
				Serzone	C		

Date:11/07/03ISR Number: 4233395-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0404622A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Personality Change	Wellbutrin	PS	Glaxosmithkline	ORAL
10 DAY							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233396-4Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0404626A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	1 WK	Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233397-6Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0404680A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
5 YR				Klonopin	C		ORAL
				Marijuana	C		

Date:11/07/03ISR Number: 4233398-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0404801A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Weight Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
				Depakote	C		
				Zyprexa	C		

Date:11/07/03ISR Number: 4233399-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0404802A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day		Nightmare		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:11/07/03ISR Number: 4233400-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0404813A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Screen Positive		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233401-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0404858A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG per day	3	MON					

Date:11/07/03ISR Number: 4233402-7Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0404871A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash Generalised		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day							

Date:11/07/03ISR Number: 4233403-9Report Type:Periodic  
Age:23 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0404872A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Oedema Peripheral					
per day	2	WK					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233404-0Report Type:Periodic  
Age:56 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0405013A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	17 DAY	Urticaria					

Date:11/07/03ISR Number: 4233405-2Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0405032A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG							
Variable dose	WK			Prevacid	C		
				Topomax	C		
				Clonazepam	C		

Date:11/07/03ISR Number: 4233406-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0405044A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Zoloft	C		
				Trileptal	C		

Date:11/07/03ISR Number: 4233407-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0405144A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Skin Odour Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Three							
times per day	WK						

Date:11/07/03ISR Number: 4233408-8Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0405146A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Menstruation Irregular		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233409-XReport Type:Periodic  
Age:59 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0405153A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	40 DAY	Back Pain		Zyprexa	C		

Date:11/07/03ISR Number: 4233410-6Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0405155A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Screen Positive		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day				Otc Products	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233411-8Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLIN-A0405156A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG At night		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
	MON	Dyspnoea		Synthroid	C	Glaxosmithkline	
				Lipitor	C		

Date:11/07/03ISR Number: 4233412-XReport Type:Periodic  
Age:36 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLIN-A0405176A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
	WK	Feeling Abnormal					

Date:11/07/03ISR Number: 4233416-7Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLIN-A0405300A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	WK	Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
		Heart Rate Increased		Allegra D	C		
		Hyperhidrosis		Contraceptives	C		ORAL
		Nausea					
		Sleep Disorder					
		Tremor					

Date:11/07/03ISR Number: 4233417-9Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLIN-A0405302A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypotension		Wellbutrin	PS	Glaxosmithkline	ORAL

UNKNOWN 100MG Unknown

Amitriptyline

SS

Date:11/07/03ISR Number: 4233418-0Report Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0405326A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice per day	3 YR	Obsessive-Compulsive Disorder Paraphilia		Wellbutrin Lithium	PS C	Glaxosmithkline Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233419-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0405331A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	DAY	Dizziness Heart Rate Increased Middle Insomnia Migraine Paraesthesia Vision Blurred		Wellbutrin Trazodone	PS C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233420-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0405366A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Myoclonus		Wellbutrin Concurrent Medications	PS  C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233421-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0405392A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Liver Function Test Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233422-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0405431A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
10 DAY		Anxiety Irritability Logorrhoea Pharyngolaryngeal Pain		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233423-4Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0405447A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	4 DAY	Agitation Limb Injury Sleep Walking		Wellbutrin  Levoxyl Rhinocort Lipitor Diovan	PS  C C C C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233424-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0405461A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin Minocycline	PS C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233425-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0405489A  
 Age: Gender:I I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233426-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0405735A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Bupropion Diflucan	PS C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233427-1Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0405757A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Twice per day	Agitation Insomnia Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
				Ativan	C		
				Zantac	C	Glaxosmithkline	
				Prozac	C		
				Menest	C	Glaxosmithkline	
				Azmacort	C		
				Intal	C		
				Flonase	C	Glaxosmithkline	
				Medroxyprogesterone	C		

Date:11/07/03ISR Number: 4233428-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0405786A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	100MG Per day	Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
		Feeling Abnormal		Wellbutrin	SS	Glaxosmithkline	ORAL
		Feeling Jittery		Lorazepam	C		
		Headache					
		Impaired Driving Ability					
		Loss Of Consciousness					

Date:11/07/03ISR Number: 4233429-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0405800A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Unknown	Hypersensitivity		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233430-1Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0405813A



Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness Nausea		Wellbutrin Ambien	PS C	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233435-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0405933A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
5 WK		Drug Ineffective Muscle Tightness Nausea Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233436-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0405964A  
 Age:53 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Drug Screen Positive		Wellbutrin	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233437-4Report Type:Periodic  
Age:76 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406022A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3 YR			Lithium Unspecified Medication	C	Glaxosmithkline	
					C		

Date:11/07/03ISR Number: 4233438-6Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406045A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Insomnia Nausea Tinnitus					

Date:11/07/03ISR Number: 4233439-8Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406059A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Bupropion	PS	Glaxosmithkline	ORAL
Other		Drug Ineffective					
150MG Twice							
per day							

Date:11/07/03ISR Number: 4233440-4Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406070A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hair Growth Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							

per day 2 YR

Date:11/07/03ISR Number: 4233441-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406073A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233442-8Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406086A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Diarrhoea					
per day		Muscle Spasms		No Concurrent			
		Pharmaceutical Product		Medications	C		
		Complaint					

Date:11/07/03ISR Number: 4233443-XReport Type:Periodic  
Age:75 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0406262A

Outcome  
Life-Threatening  
Hospitalization -  
Initial or Prolonged

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Freedom Of Information (FOI) Report

Disability  
Other

Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
			Seroquel	C		

Date:11/07/03ISR Number: 4233444-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0406286A  
Age: Gender: I/FU:I

Duration	PT	Report Source	Product	Role	Manufacturer	Route
	Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233445-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0406293A  
Age:14 YR Gender:Male I/FU:I

Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	Tic		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233446-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0406304A  
Age: Gender:Female I/FU:I

Duration	PT	Report Source	Product	Role	Manufacturer	Route
	Aggression Anger		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233447-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0406363A  
Age:39 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1	MON		Prometrium	C		
				Clomiphene	C		
				Metformin	C		

Date:11/07/03ISR Number: 4233448-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0406401A  
 Age:41 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Anorexia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	4	Anxiety					
		Dissociation		Paxil	C	Glaxosmithkline	
		Nausea					

Date:11/07/03ISR Number: 4233449-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0406406A  
 Age:65 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice		Disturbance In Attention		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3	Dizziness		Aspirin	C	Glaxosmithkline	
				Lupron	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233450-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406437A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tachycardia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Unknown						

Date:11/07/03ISR Number: 4233451-9Report Type:Periodic  
 Age:25 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0406454A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Twice	Anger					
per day	3 WK	Depression		Birth Control	C		

Date:11/07/03ISR Number: 4233452-0Report Type:Periodic  
 Age:53 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0406571A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Emotional Distress		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG	Twice	Tinnitus					
per day	66 DAY			Ambien	C		

Date:11/07/03ISR Number: 4233453-2Report Type:Periodic  
 Age:60 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406688A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyskinesia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Myoclonus					

Date:11/07/03ISR Number: 4233454-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0406714A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Interaction		Cardizem	SS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233455-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0406770A  
Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoglycaemia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	8 DAY			Glyburide	C		
				Metformin	C		

Date:11/07/03ISR Number: 4233456-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0406852A  
Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Head Injury		Zoloft	C		
100MG Per day		Loss Of Consciousness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233457-XReport Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406863A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	9	MON		Wellbutrin	PS	Glaxosmithkline	ORAL
				Glucosamine/Chondroitin	SS		
10	DAY						

Date:11/07/03ISR Number: 4233460-XReport Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406875A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Per day		YR		Wellbutrin	PS	Glaxosmithkline	ORAL
				Synthroid	C	Glaxosmithkline	
				Lotrel	C		
				Estratest	C		
				Prozac	C		
				Skelaxin	C		
				Relafen	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4233461-1Report Type:Periodic  
Age:13 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0406886A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Twice				Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3	YR					
				Strattera	C		ORAL
				Concerta	C		ORAL
36MG Unknown							
SUBCUTANEOUS				Gonadotropin	C		



Date:11/07/03ISR Number: 4233462-3Report Type:Periodic  
Age:17 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0406898A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menorrhagia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day				Trazodone	C		

Date:11/07/03ISR Number: 4233463-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406904A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233464-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406907A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Spasms		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Unknown							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233465-9Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406918A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Ineffective					
per day		Somnolence		Synthroid	C	Glaxosmithkline	
				Vioxx	C		

Date:11/07/03ISR Number: 4233466-0Report Type:Periodic  
Age:19 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0406920A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day	5 DAY	Agitation					
		Belligerence		Thyroid Medication	C		
		Dissociation					

Date:11/07/03ISR Number: 4233467-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406927A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion					

Date:11/07/03ISR Number: 4233468-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406929A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Amenorrhoea					

Date:11/07/03ISR Number: 4233469-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406987A

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Convulsion		Bupropion	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233470-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0407007A  
Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233471-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0407022A  
Age: Gender:I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

400MG per day

Date:11/07/03ISR Number: 4233472-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0407068A  
Age: Gender:I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4233473-8Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407069A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
450MG per day	1 YR	Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Chest Pain		Effexor	C		
				Vitamins	C		

Date:11/07/03ISR Number: 4233474-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407074A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Electroencephalogram Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233475-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407080A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Twice per day		Abnormal Dreams		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233476-3Report Type:Periodic  
Age:47 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0407089A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice per day		Hallucinations, Mixed Urine Amphetamine Positive		Wellbutrin	PS	Glaxosmithkline	ORAL
				Provigil	SS		
				Albuterol	C	Glaxosmithkline	
				Viagra	C		

Date:11/07/03ISR Number: 4233477-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407104A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Nonspecific Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/07/03ISR Number: 4233478-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407127A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG per day	1 YR			Effexor	C		
225MG per day	1 YR						

Date:11/07/03ISR Number: 4233479-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0407131A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
3 DAY							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/10/03ISR Number: 4231010-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0433145A

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Ageusia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Anorexia					
per day	22 WK	Dysgeusia		Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day		Fatigue		Oxycontin	SS		
		Irregular Sleep Phase		Remeron	SS		
		Middle Insomnia					
		Sleep Disorder					

Date:11/10/03ISR Number: 4231012-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0433261A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
		Depression					
		Fatigue					
		Loss Of Consciousness					
		Myalgia					
		Vomiting					

Date:11/10/03ISR Number: 4231013-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0433263A

Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Per day		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Medication Error		Wellbutrin	SS	Glaxosmithkline	ORAL
per day				Xanax	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Required 1 1 PER DAY Intervention to ORAL Prevent Permanent Impairment/Damage		Abdominal Pain Upper Agitation  Anxiety  Completed Suicide Depression Diarrhoea Flatulence Headache Insomnia Nausea Paranoia Psychotic Disorder Pyrexia Toothache Tremor Vomiting		Wellbutrin Sr 100 Mg Glaxo	PS	Glaxo	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/12/03ISR Number: 4243148-7Report Type:Direct  
 Age: Gender: I/FU:I

Company Report #USP 56117

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin Sr	PS	Glaxosmithkline	
TABLET, EXTENDED RELEASE, 150 MG 150 MG							
TABLET, EXTENDED RELEASE							
				Wellbutrin Xl	SS	Glaxosmithkline	

Date:11/13/03ISR Number: 4234347-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0433006A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							
150MG Per day							
				Wellbutrin	SS	Glaxosmithkline	ORAL

Date:11/13/03ISR Number: 4234348-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0438844A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cerebrovascular Spasm		Wellbutrin	PS	Glaxosmithkline	



Date:11/13/03ISR Number: 4234376-5Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0314554A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Oropharyngeal Swelling		Zyban	PS	Glaxosmithkline	ORAL
1TAB See							

dosage text

Date:11/14/03ISR Number: 4235305-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0354550A

Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Discomfort
Initial or Prolonged	Aggression
Other	Agitation
	Anger
	Anorexia
	Anxiety
	Asthenia
	Confusional State
	Coordination Abnormal
	Depression
	Disturbance In Attention
	Dizziness
	Drug Ineffective
	Drug Withdrawal Syndrome
	Fatigue
	Gait Disturbance
	Homicidal Ideation
	Influenza
	Intentional Self-Injury
	Irritability

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
20MG Per day			Paxil	PS	Glaxosmithkline	ORAL
150MG Twice per day			Wellbutrin	SS	Glaxosmithkline	ORAL
			Ritalin	C		
			Amoxicillin	C	Glaxosmithkline	

Lethargy  
 Malaise  
 Memory Impairment  
 Myalgia  
 Nausea  
 Nightmare  
 Obsessive Thoughts  
 Obsessive-Compulsive Disorder  
 Paraesthesia  
 Paranoia  
 Pyrexia  
 Suicidal Ideation  
 Suicide Attempt  
 Tremor  
 Vertigo

Date:11/14/03ISR Number: 4235313-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0438799A  
 Age: YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose			Wellbutrin	PS	Glaxosmithkline	ORAL
Other						
3 WK						

PT  
 Convulsion  
 Fall  
 Skin Laceration

Date:11/14/03ISR Number: 4235314-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0438824A  
 Age: Gender: I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose			Wellbutrin	PS	Glaxosmithkline	

PT  
 Medication Error

Date:11/14/03ISR Number: 4235330-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0313823A  
 Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Angioneurotic Oedema Arthralgia C-Reactive Protein Increased Erythema Feeling Hot Neutrophil Count Increased Oedema Peripheral Petechiae Pyrexia		Zyban	PS	Glaxosmithkline	ORAL

Date:11/14/03ISR Number: 4235332-3Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0314310A  
Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 14 DAY		Atrial Fibrillation		Zyban	PS	Glaxosmithkline	
				No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/17/03ISR Number: 4236118-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0433179A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly			Drug Exposure During Pregnancy	Wellbutrin	PS	Glaxosmithkline	
			Ventricular Septal Defect				

Date:11/17/03ISR Number: 4236119-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0438833A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Medication Error	Wellbutrin	PS	Glaxosmithkline	

Date:11/17/03ISR Number: 4236120-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439228A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Medication Error	Wellbutrin	PS	Glaxosmithkline	
			Pharmaceutical Product Complaint				

Date:11/17/03ISR Number: 4236121-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439423A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Medication Error	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/17/03ISR Number: 4236122-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439429A

Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	
		Insomnia		Wellbutrin	SS	Glaxosmithkline	ORAL
		Medication Error		Lorazepam	C		

Date:11/17/03ISR Number: 4236123-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439431A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:11/17/03ISR Number: 4236124-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439627A  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	
		Pharmaceutical Product		Wellbutrin	SS	Glaxosmithkline	ORAL
450MG Per day	5 WK	Complaint		Thyroid	C		
				Provigil	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/17/03ISR Number: 4236135-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0313883A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Zyban	PS	Glaxosmithkline	ORAL
150MG See							
dosage text	14	DAY					
		Crying					
		Insomnia					
		Petit Mal Epilepsy					
		Visual Disturbance					

Date:11/17/03ISR Number: 4236136-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0314049A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Psoriasis		Zyban	PS	Glaxosmithkline	ORAL
150MG See							
dosage text	34	DAY					

Date:11/17/03ISR Number: 4236140-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0314404A  
Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eczema		Zyban	PS	Glaxosmithkline	ORAL
150MG See							
dosage text	13	DAY					
UNKNOWN	75MG per day			Aspirin	C	Glaxosmithkline	
UNKNOWN	40MG per day			Pravastatin	C		
UNKNOWN	625MG per day	7	DAY	Co-Amoxiclav	C	Glaxosmithkline	
UNKNOWN	50MG per day	7	DAY	Nitrofurantoin	C	Glaxosmithkline	
UNKNOWN	500MG per day	7	DAY	Cefadroxil	C		

UNKNOWN 4MG per day

Perindopril

C

Date:11/17/03ISR Number: 4236141-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0314463A

Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2TAB Twice				Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	13 DAY		Rash Erythematous				

Date:11/17/03ISR Number: 4236250-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439877A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Accident Convulsion Loss Of Consciousness	Zyban	PS	Glaxosmithkline	ORAL

Date:11/17/03ISR Number: 4236559-7Report Type:Direct

Company Report #CTU 206230

Age:40 YR Gender:Female I/FU:I

Outcome	PT
Disability	Abnormal Behaviour
Other	Attention Deficit/Hyperactivity Disorder Bulimia Nervosa Condition Aggravated Crying Depression

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Freedom Of Information (FOI) Report

Dose	Duration	Diabetes Mellitus Insulin-Dependent Drug Withdrawal Syndrome	Report Source	Product	Role	Manufacturer	Route
40 MG PER DAY		Morbid Thoughts Post-Traumatic Stress Disorder		Prozac 20 Mg Eli Lilley	PS	Eli Lilly	ORAL
ORAL		Relationship Breakdown		Wellbutrin 150 Mg	SS		
150 MG PER DAY							

Date:11/17/03ISR Number: 4236592-5Report Type:Direct Company Report #CTU 206217  
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 300 MG QD Required ORAL		Hepatitis Acute		Wellbutrin Xl 300 Mg	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage				Mircette Ocp Ecchinacea	C C		

Date:11/17/03ISR Number: 4237080-2Report Type:Expedited (15-DaCompany Report #200301196  
 Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
500 MG BID PO		Abdominal Pain Anxiety	Consumer Health	Extra Strength Tylenol Product	PS		ORAL
100 MG TID PO		Blood Creatinine Abnormal	Professional	Wellbutrin	SS		ORAL
1300 MG BID PO		Blood Urea Abnormal Dysphagia Hypokalaemia Renal Disorder	Other	Tylenol Arthritis Pain Er Caplet 650mg/Caplet	SS		ORAL



Atenolol C  
Hydrochlorothiazide C

Date:11/17/03ISR Number: 4237081-4Report Type:Expedited (15-DaCompany Report #200301196  
Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Pain Anxiety	Consumer Health	Extra Strength Tylenol Product	PS		ORAL
500 MG BID PO		Blood Creatinine Abnormal	Professional	Wellbutrin	SS		ORAL
100 MG TID PO		Blood Urea Abnormal Dysphagia Hyperkalaemia	Other	Tylenol Arthritis Pain Er Caplet 650mg/Caplet	SS		ORAL
1300 MG BID PO		Renal Disorder		Atenolol Hydrochlorothiazide	C C		

Date:11/18/03ISR Number: 4236726-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439007A  
Age:78 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1TAB Twice Initial or Prolonged per day	1 MON	Chills Condition Aggravated Tremor		Wellbutrin Flomax Cozaar Indapamide Duragesic	PS C C C C	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/18/03ISR Number: 4236727-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439046A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blindness		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2 DAY			Norvasc	C		
				Xanax	C		
				Zocor	C		

Date:11/18/03ISR Number: 4236728-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439062A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Doxipan	C	Glaxosmithkline	
		Medication Error		Sonata	C		

Date:11/18/03ISR Number: 4236759-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439007A

Age:78 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Chills	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Twice							
Initial or Prolonged		Condition Aggravated					
per day	1 MON			Flomax	C		
		Depression		Cozaar	C		
		Tremor		Indapamide	C		
				Duragesic	C		

Date:11/18/03ISR Number: 4236760-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439046A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other 300MG Per day 2 DAY	Blindness	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
		Professional	Norvasc	C		
			Xanax	C		
			Zocor	C		

Date:11/18/03ISR Number: 4236761-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439062A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Per day		Convulsion	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
		Medication Error	Professional	Doxipan Sonata	C C	Glaxosmithkline	

Date:11/18/03ISR Number: 4236780-8Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0312102A  
 Age:26 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 22 DAY		Arthralgia	Health	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged 1TAB Single dose		Body Temperature Increased Eyelid Oedema Face Oedema Myalgia Oedema Peripheral Urticaria	Professional	Methylprednisolone	C		

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Freedom Of Information (FOI) Report

Date:11/18/03ISR Number: 4236781-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0312169A  
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Vision Blurred	Health	Zyban	PS	Glaxosmithkline	ORAL
		Visual Acuity Reduced	Professional				

Date:11/18/03ISR Number: 4236783-3Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0041959A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anxiety		Zyban	PS	Glaxosmithkline	ORAL
1TAB Per day	1 DAY						
Initial or Prolonged		Cardiovascular Disorder					

Date:11/19/03ISR Number: 4237439-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0379981A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Blood Ethanol Increased	Consumer	Bupropion	PS	Glaxosmithkline	ORAL
		Cardiomegaly		Carbamazepine	SS		
UNKNOWN		Coma		Olanzapine	SS		
UNKNOWN		Completed Suicide		Clonazepam	SS		
		Convulsion		Ethanol	SS		
		Depressed Level Of Consciousness		Aspirin	SS	Glaxosmithkline	
		Ecchymosis					
		Hepatic Steatosis					
		Intentional Misuse					
		Mydriasis					
		Ovarian Cyst					
		Pulmonary Congestion					
		Pulmonary Oedema					
		Pupil Fixed					

Date:11/19/03ISR Number: 4237440-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0379982A  
Age:36 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Death	Areflexia		Bupropion	PS	Glaxosmithkline	ORAL
Hospitalization -	Brain Death		Clonazepam	SS		
UNKNOWN						
Initial or Prolonged	Cardio-Respiratory Arrest		Trazodone	SS		
UNKNOWN						
	Completed Suicide		Gabapentin	SS		
	Depressed Level Of					
	Consciousness					
	Hyporeflexia					
	Intentional Misuse					
	Pupillary Reflex Impaired					
	Subarachnoid Haemorrhage					
	Tachycardia					

Date:11/19/03ISR Number: 4237441-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0379983A  
Age:51 YR Gender:Male I/FU:F

Outcome	PT
Death	Cardio-Respiratory Arrest
	Completed Suicide
	Coronary Artery
	Atherosclerosis

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		Hypertension Intentional Misuse Nausea	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Bupropion Ethanol	PS SS	Glaxosmithkline	ORAL ORAL

Date:11/19/03ISR Number: 4237446-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439829A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:11/19/03ISR Number: 4237447-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439996A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Insomnia Muscle Tightness					

Date:11/19/03ISR Number: 4237449-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0440188A  
Age: YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Disorder Convulsion Depression Joint Dislocation Renal Impairment		Wellbutrin	PS	Glaxosmithkline	
Other							

Date:11/19/03ISR Number: 4237463-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0379984A  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Circulatory Collapse	Consumer	Bupropion	PS	Glaxosmithkline	ORAL
		Completed Suicide		Ethylene Glycol	SS		ORAL
		Convulsion		Hypericum Perforatum	SS		ORAL
		Drug Toxicity		Nefazodone	SS		
		Fall		St. Johns Wort	C		
		Haemorrhage Intracranial		Lorazepam	C		
		Intentional Misuse					
		Mydriasis					
		Pupil Fixed					
		Ventricular Fibrillation					
		Ventricular Tachycardia					

Date:11/19/03ISR Number: 4237464-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0379985A  
Age:46 YR Gender:Female I/FU:F

Outcome	PT
Death	Aspiration
	Cardio-Respiratory Arrest
	Cerebellar Haemorrhage
	Cerebral Haemorrhage
	Coma
	Depressed Level Of

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Consciousness Drug Screen Positive Hypotension Mydriasis	Consumer	Bupropion Naproxen Cocaine	PS SS SS	Glaxosmithkline	ORAL ORAL
UNKNOWN		Tachycardia					

Date:11/19/03ISR Number: 4237465-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0379987A  
Age:35 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Acidosis	Consumer	Bupropion	PS	Glaxosmithkline	ORAL
Hospitalization - UNKNOWN			Cardiovascular Disorder		Venlafaxine	SS		
Initial or Prolonged UNKNOWN			Completed Suicide		Methylphenidate	SS		
			Convulsion		Quetiapine	SS		
			Depressed Level Of Consciousness		Clonazepam	SS		
			Hypotension		Trazodone	SS		
			Hypothermia		Gabapentin	SS		
			Intentional Misuse		Lansoprazole	SS		
			Respiratory Depression					

Date:11/19/03ISR Number: 4237605-7Report Type:Direct Company Report #CTU 206421  
Age:46 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	100 MG BID		Angioneurotic Oedema		Wellbutrin	PS		
Hospitalization - Initial or Prolonged			Rash Erythematous		Prednisone	C		
			Rash Papular		Lexapro	C		
					Nexium	C		
					Nadolol	C		



Date:11/19/03ISR Number: 4238278-XReport Type:Direct  
Age:39 YR Gender:Female I/FU:I

Company Report #CTU 206492

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Stevens-Johnson Syndrome		Wellbutrin 100 Mg Tablet	PS		ORAL

Date:11/19/03ISR Number: 4238962-8Report Type:Expedited (15-DaCompany Report #S03-USA-04440-01  
Age:17 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Gastrooesophageal Reflux Disease	Health Professional	Lexapro (Escitalopram)	PS		ORAL
10 MG QD PO		Grand Mal Convulsion Throat Irritation		Wellbutrin (Bupropion Hydrochloride)	SS		
200 MG BID				Wellbutrin (Bupropion Hydrochloride)	SS		
100 MG BID				Wellbutrin (Bupropion Hydrochloride)	SS		
100 MG QD							

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Freedom Of Information (FOI) Report

Neurontin  
(Gabapentin) C

Date:11/19/03ISR Number: 4239070-2Report Type:Expedited (15-DaCompany Report #2003-03773  
Age:50 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Literature Health Professional	Lorazepam (Watson Laboratories) (Lorazepam) Tablet Diphenhydramine (Watson Laboratories) (Diphenhydramine Hydrochloride) Bupropion (Bupropion)	PS    SS SS		

Date:11/20/03ISR Number: 4238031-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0379986A  
Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Arrhythmia Cardiac Arrest		Bupropion Olanzapine	PS SS	Glaxosmithkline	ORAL
UNKNOWN		Completed Suicide		Clonazepam	SS		
UNKNOWN		Convulsion Gastrointestinal Motility Disorder Hypokinesia Hypotension Intentional Misuse Mental Status Changes Overdose Pupillary Reflex Impaired Ventricular Fibrillation		Aspirin Acetaminophen Sertraline Trazodone Ibuprofen	SS SS SS SS SS	Glaxosmithkline Glaxosmithkline	

Date:11/20/03ISR Number: 4238032-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0379988A  
Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Acidosis		Bupropion	PS	Glaxosmithkline	ORAL
		Blood Creatine		Fluoxetine	SS		
UNKNOWN		Phosphokinase Increased		Acetaminophen +			
		Cardiac Arrest		Hydrocodone	SS		
UNKNOWN		Cardio-Respiratory Arrest					
		Dialysis					
		Hypotension					

Date:11/20/03ISR Number: 4238033-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0379989A  
Age:16 YR Gender: I/FU:F

Outcome	PT
Death	Acidosis
	Cardio-Respiratory Arrest
	Coma
	Completed Suicide
	Convulsion

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Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Depressed Level Of Consciousness Hypotension				
UNKNOWN		Intentional Misuse Multiple Drug Overdose	Bupropion Quetiapine	PS SS	Glaxosmithkline	ORAL
UNKNOWN		Mydriasis Pupil Fixed	Valproic Acid	SS		

Date:11/20/03ISR Number: 4238034-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0379990A  
Age:46 YR Gender: I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Death		Blood Pressure Decreased Cardio-Respiratory Arrest	Bupropion Alprazolam	PS SS	Glaxosmithkline	ORAL
UNKNOWN		Coma	Olanzapine	SS		
UNKNOWN		Completed Suicide Cyanosis Depressed Level Of Consciousness Heart Rate Increased Intentional Misuse Miosis Oxygen Saturation Decreased Respiratory Rate Increased	Gabapentin Cyanide	SS SS		

Date:11/20/03ISR Number: 4238037-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0380101A  
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Death		Blood Ph Decreased Completed Suicide	Bupropion Amitriptyline	PS SS	Glaxosmithkline	ORAL
UNKNOWN		Disorientation	Clonidine	SS		

Hypotension  
Intentional Misuse  
Loss Of Consciousness  
Vascular Resistance  
Systemic

Buspirone SS  
Clonazepam SS

Date:11/20/03ISR Number: 4238038-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0380102A  
Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Acidosis Cardio-Respiratory Arrest		Bupropion Carbamazepine	PS SS	Glaxosmithkline	ORAL
UNKNOWN		Completed Suicide		Ethanol	SS		
UNKNOWN		Convulsion Drug Ineffective Intentional Misuse Loss Of Consciousness Mydriasis Pupil Fixed Pyrexia		Benzodiazepine	C		

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Date:11/20/03ISR Number: 4238039-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383127A  
 Age:17 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest		Bupropion	PS	Glaxosmithkline	ORAL
UNKNOWN		Completed Suicide		Fluvoxamine	SS		
		Intentional Misuse					
		Self-Medication					

Date:11/20/03ISR Number: 4238040-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383129A  
 Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest		Bupropion	PS	Glaxosmithkline	ORAL
UNKNOWN		Coagulopathy		Isotretinoin	SS		
UNKNOWN		Completed Suicide		Diphenhydramine	SS		
		Intentional Misuse					
		Multi-Organ Failure					

Date:11/20/03ISR Number: 4238041-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383135A  
 Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aspiration		Bupropion	PS	Glaxosmithkline	ORAL
150MG Single		Cardio-Respiratory Arrest					
dose		Completed Suicide		Citalopram	SS		
UNKNOWN	30MG Single	Convulsion					
dose		Intentional Misuse		Dextroamphetamine	SS	Glaxosmithkline	
UNKNOWN	30MG Single	Somnolence					
dose				Lisinopril	SS		
				Trazodone	SS		

Date:11/20/03ISR Number: 4238042-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383137A  
Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Apnoea		Bupropion	PS	Glaxosmithkline	ORAL
Hospitalization -		Blood Pressure Decreased		Trazodone	SS		
UNKNOWN							
Initial or Prolonged		Cardio-Respiratory Arrest		Venlafaxine	SS		
UNKNOWN							
		Coma					
		Completed Suicide					
		Depressed Level Of					
		Consciousness					
		Electroencephalogram					
		Abnormal					
		Intentional Misuse					
		Pupil Fixed					

Date:11/20/03ISR Number: 4238043-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383140A  
Age:30 YR Gender:Male I/FU:F

Outcome	PT
Death	Completed Suicide
Hospitalization -	Convulsion
Initial or Prolonged	Disseminated
	Intravascular Coagulation
	Electroencephalogram
	Abnormal

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hepatic Failure Intentional Misuse Mydriasis					
		Nystagmus		Bupropion	PS	Glaxosmithkline	ORAL
		Posturing		Relafen	SS	Glaxosmithkline	ORAL
		Pyrexia		Acetaminophen + Diphenhydramine	SS		
UNKNOWN		Renal Failure					
UNKNOWN		Rhabdomyolysis		Venlafaxine	SS		
UNKNOWN		Tachycardia		Tylenol	SS	Glaxosmithkline	
UNKNOWN				Librium	SS		
UNKNOWN				Ambien	SS		
UNKNOWN				Skelaxin	SS		
UNKNOWN				Midol	SS		
UNKNOWN				Pyridium	SS		
UNKNOWN				Aleve	SS		

Date:11/20/03ISR Number: 4238044-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383143A

Age:19 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest		Bupropion Tramadol	PS SS	Glaxosmithkline	ORAL
UNKNOWN				Metoclopramide	SS	Glaxosmithkline	

Date:11/20/03ISR Number: 4238045-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383146A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest		Bupropion	PS	Glaxosmithkline	ORAL



UNKNOWN Completed Suicide Olanzapine SS  
Intentional Misuse  
Loss Of Consciousness

Date:11/20/03ISR Number: 4238046-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383147A  
Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Agitation		Bupropion	PS	Glaxosmithkline	ORAL
		Cardio-Respiratory Arrest		Quetiapine	SS		
UNKNOWN		Completed Suicide		Levofloxacin	SS		
		Intentional Misuse					

Date:11/20/03ISR Number: 4238047-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383148A  
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardio-Respiratory Arrest		Bupropion	PS	Glaxosmithkline	ORAL
		Completed Suicide		Quetiapine	SS		
UNKNOWN		Intentional Misuse					

Date:11/20/03ISR Number: 4238048-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383151A  
Age:52 YR Gender:Male I/FU:F

Outcome	PT
Death	Cardiac Arrest
	Completed Suicide

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Intentional Misuse

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
UNKNOWN			Bupropion	PS	Glaxosmithkline	ORAL
UNKNOWN			Carvedilol	SS	Glaxosmithkline	
UNKNOWN			Amlodipine + Benazepril	SS		
UNKNOWN			Oxycodone	SS		
UNKNOWN			Celecoxib	SS		

Date:11/20/03ISR Number: 4238051-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0421236A  
Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day	3 DAY	Agitation	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	25MG Per day	2 MON	Breast Pain	Paxil	SS	Glaxosmithkline	ORAL
150MG Twice			Depression	Wellbutrin	SS	Glaxosmithkline	ORAL
per day			Derealisation				
UNKNOWN			Disturbance In Attention	Xanax	C		
UNKNOWN			Dysgeusia	Bc Powder	C	Glaxosmithkline	
UNKNOWN			Dyspepsia	Estradiol	C		
UNKNOWN	.1MG	Unknown	Eye Pain	Levoxyl	C	Glaxosmithkline	
UNKNOWN			Eye Swelling	Medroxyprogesterone	C		
UNKNOWN	500MG	Unknown	Fatigue	Calcium	C		
UNKNOWN			Gastric Disorder	Thera M	C		
RESPIRATORY			Headache	Flovent	C	Glaxosmithkline	
(INHALATION)			Hunger				
			Hyperventilation	Albuterol	C	Glaxosmithkline	
			Insomnia	Citracal	C		
			Libido Increased	Multivitamins	C		

Nausea  
Negative Thoughts  
Paraesthesia  
Paraesthesia Oral  
Respiratory Sighs  
Restlessness  
Suicidal Ideation  
Tension

Date:11/20/03ISR Number: 4238107-4Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426613A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypoaesthesia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG per day		Paraesthesia		Accuretic	C		
				Celexa	C		
				Advair	C	Glaxosmithkline	

Date:11/20/03ISR Number: 4238108-6Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0426882A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							

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Date:11/20/03ISR Number: 4238109-8Report Type:Periodic  
Age:37 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0427548A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day	5 DAY	Hallucination, Auditory	Wellbutrin	PS	Glaxosmithkline	ORAL
Other	10MG At night		Psychotic Disorder	Ambien	C		
	.05MG Per day			Synthroid	C	Glaxosmithkline	
	600MG Per day			Tegretol	C		

Date:11/20/03ISR Number: 4238110-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427869A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day	8 DAY	Drug Ineffective	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238111-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427916A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Tinnitus	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238112-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428108A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Rash	Wellbutrin	PS	Glaxosmithkline	ORAL
				Wellbutrin	C	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238113-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428114A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238114-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0428266A  
 Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238115-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0428703A  
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	4 DAY			Vioxx	C		
				Lortab	C		
				Nexium	C		
				Synthroid	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/20/03ISR Number: 4238116-5Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428730A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Irritability		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK			Celexa	SS		

Date:11/20/03ISR Number: 4238117-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0428860A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Weight Decreased					

Date:11/20/03ISR Number: 4238118-9Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428958A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction		Wellbutrin	PS	Glaxosmithkline	
		Pyrexia		Sonata	SS		
		Urticaria		Unknown Medication	C		
		Vomiting					

Date:11/20/03ISR Number: 4238119-0Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428977A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
2 WK							

Date:11/20/03ISR Number: 4238120-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429120A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anger		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238121-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0429224A  
 Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthritis		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK	Rash		Zoloft	C		

Date:11/20/03ISR Number: 4238122-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0429225A  
 Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK	Rash					
		Rash Macular					
		Rash Papular					
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/20/03ISR Number: 4238123-2Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429662A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tension Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2 DAY						

Date:11/20/03ISR Number: 4238124-4Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429668A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4 DAY	Dry Mouth Myalgia					

Date:11/20/03ISR Number: 4238125-6Report Type:Periodic  
 Age:45 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429689A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY						

Date:11/20/03ISR Number: 4238126-8Report Type:Periodic  
 Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429690A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK						

Date:11/20/03ISR Number: 4238127-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429717A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Agitation  
Feeling Jittery  
Heart Rate Increased

Wellbutrin

PS

Glaxosmithkline

ORAL

Date:11/20/03ISR Number: 4238128-1Report Type:Periodic  
Age:17 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429813A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:11/20/03ISR Number: 4238129-3Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429818A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oral Mucosal Blistering		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238130-XReport Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430013A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	4 DAY	Headache		Nexium	C		
		Nausea					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/20/03ISR Number: 4238131-1Report Type:Periodic  
 Age:68 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0430054A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	7 DAY	Drug Ineffective	Wellbutrin	PS	Glaxosmithkline	ORAL
			Pharmaceutical Product Complaint	Verapamil	C		
			Stool Analysis Abnormal	Doxazosin	C		

Date:11/20/03ISR Number: 4238132-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430105A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Drug Interaction	Wellbutrin	PS	Glaxosmithkline	ORAL
				Alcohol	SS		ORAL

Date:11/20/03ISR Number: 4238133-5Report Type:Periodic  
 Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430282A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	1 DAY	Dizziness	Wellbutrin	PS	Glaxosmithkline	ORAL
				Toprol	C		

Date:11/20/03ISR Number: 4238134-7Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430486A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Hunger	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238135-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430602A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Headache		Effexor	C		

Date:11/20/03ISR Number: 4238136-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430642A  
 Age:15 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Joint Sprain		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 MON	Sleep Walking		Concerta	SS		ORAL
54MG Per day							

Date:11/20/03ISR Number: 4238137-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430643A  
 Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Lip Blister		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	6 DAY	Swelling Face		Hydroquinone	C		
				Ogen	C		
				Prometrium	C		
				Actonel	C		
				Buspar	C		
				Topomax	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/20/03ISR Number: 4238138-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430651A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
			Central Nervous System Stimulation Sleep Disorder				

Date:11/20/03ISR Number: 4238139-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430973A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK		Dizziness	Zyprexa Nembutal	C C		

Date:11/20/03ISR Number: 4238140-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430982A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day			Rash Pruritic	Lotrel	C		

Date:11/20/03ISR Number: 4238141-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431126A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
2 WK			Influenza Like Illness Myalgia				

Date:11/20/03ISR Number: 4238142-6Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431204A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:11/20/03ISR Number: 4238143-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431216A  
 Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
				Wellbutrin	C	Glaxosmithkline	ORAL
1	YR						

Date:11/20/03ISR Number: 4238144-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431217A  
 Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
4	DAY			Zoloft	C		



150MG Per day	3	DAY	Anxiety	Wellbutrin	PS	Glaxosmithkline	ORAL
			Depression	Multivitamin	C		
			Drug Withdrawal Syndrome	Vitamin E	C		
			Headache	Niacin	C		
			Muscle Tightness	Zocor	C		
				Flomax	C		
				Verapamil	C		
				Allopurinol	C	Glaxosmithkline	
				Altace	C		
				Avandamet	C	Glaxosmithkline	
				Celebrex	C		
				Xanax	C		

Date:11/20/03ISR Number: 4238150-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431396A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Flushing Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/20/03ISR Number: 4238151-7Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431413A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:11/20/03ISR Number: 4238152-9Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431421A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Heart Rate Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	10 DAY						
		Insomnia		Vivelle-Dot	C		

Date:11/20/03ISR Number: 4238153-0Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431428A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							
100MG Per day	8 MON			Thyroxine	C	Glaxosmithkline	

Date:11/20/03ISR Number: 4238154-2Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431587A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day							
300MG Per day				Wellbutrin	C	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238155-4Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431610A



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/20/03ISR Number: 4238156-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431613A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238157-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431615A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238158-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431623A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Unknown							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/20/03ISR Number: 4238159-1Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431624A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	DAY						

Date:11/20/03ISR Number: 4238160-8Report Type:Periodic  
Age: Gender:I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431625A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
				Risperdal	SS		
UNKNOWN							

Date:11/20/03ISR Number: 4238161-XReport Type:Periodic  
Age:52 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431798A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Chest Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Heart Rate Increased		Wellbutrin	SS	Glaxosmithkline	ORAL
1 MON							

Date:11/20/03ISR Number: 4238162-1Report Type:Periodic  
Age:12 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431810A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Chest Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY	Chills					
		Palpitations					

Date:11/20/03ISR Number: 4238163-3Report Type:Periodic  
Age: Gender:I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431830A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Mania		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238164-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431833A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Mania		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238165-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431834A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspepsia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238166-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432099A  
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 1 MON				Depakote	C		
				Abilify	C		
				Klonopin	C		
				Glucovance	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/20/03ISR Number: 4238167-0Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432100A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Urine Flow Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:11/20/03ISR Number: 4238168-2Report Type:Periodic  
Age:37 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432137A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Erectile Dysfunction		Wellbutrin	PS	Glaxosmithkline	ORAL
2 MON							
		Insomnia		Effexor	SS		
225MG Per day	2 MON						
		Libido Decreased					
		Tremor					

Date:11/20/03ISR Number: 4238169-4Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432144A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1 WK						
				Tylenol	C	Glaxosmithkline	

Date:11/20/03ISR Number: 4238170-0Report Type:Periodic  
Age:28 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432147A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Enuresis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238171-2Report Type:Periodic  
Age:34 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432149A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Per day	4 DAY						

Date:11/20/03ISR Number: 4238172-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432151A  
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK						
		Energy Increased		Ventolin Mdi	C	Glaxosmithkline	
		Heart Rate Increased		Synthroid	C	Glaxosmithkline	
				Trazodone	C		
				Depakote	C		

Date:11/20/03ISR Number: 4238173-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432152A  
 Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK						
		Rash Erythematous		Flexeril	C		
				Multivitamin	C		
				Darvocet	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/20/03ISR Number: 4238174-8Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432153A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	3 DAY	Agitation Nervousness Somnolence		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238175-XReport Type:Periodic  
Age:40 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0432263A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG Per day 9 DAY	Breast Discharge Galactorrhoea Paraesthesia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238176-1Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432266A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG Per day 2 WK	Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238177-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432296A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG Per day 3 WK	Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
				Toprol Xl	C		
				Ambien	C		

Date:11/20/03ISR Number: 4238178-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432305A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Premature Ejaculation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238179-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432306A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Palpitations		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG In the morning							
150MG Twice per day							
				Wellbutrin	C	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238180-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432348A  
Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vomiting		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 2 WK							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/20/03ISR Number: 4238181-5Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432393A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	10 DAY	Abdominal Distension	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238182-7Report Type:Periodic  
 Age:29 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432409A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238183-9Report Type:Periodic  
 Age: YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0432431A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Tinnitus	Wellbutrin	PS	Glaxosmithkline	

Date:11/20/03ISR Number: 4238184-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432438A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	3 WK	Dizziness	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/20/03ISR Number: 4238185-2Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432445A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Hallucination	Wellbutrin	PS	Glaxosmithkline	ORAL



Date:11/20/03ISR Number: 4238186-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0433118A  
Age: YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	

Date:11/21/03ISR Number: 4238845-3Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0440268A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs Drug Exposure During Pregnancy		Bupropion	PS	Glaxosmithkline	

Date:11/21/03ISR Number: 4238847-7Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0292359A  
Age:48 YR Gender:Male I/FU:F

Outcome	PT
Other	Blood Immunoglobulin E Increased Chest Pain Lymphopenia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN	150MG	As		Zyntabac	PS	Glaxosmithkline	
directed	9 DAY	Pruritus Generalised					
UNKNOWN		White Blood Cell Count		Aluminum Hydroxide	C		
UNKNOWN		Increased		Almagate	C		

Date:11/21/03ISR Number: 4238849-0Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0311911A  
 Age:39 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG	Twice	Acne		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		per day	Convulsion					
Other			Fall		No Concurrent Medication	C		
			Respiratory Arrest					
			Tooth Injury					

Date:11/21/03ISR Number: 4238853-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0312384A  
 Age:51 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG	See	Drug Withdrawal Syndrome		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization -		dosage text	Hepatic Trauma					
Initial or Prolonged		6 WK	Pelvic Fracture					
			Suicide Attempt					

Date:11/21/03ISR Number: 4238856-8Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0313519A  
 Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Disability		Convulsion Depressed Level Of Consciousness Overdose		Zyban	PS	Glaxosmithkline	ORAL

Date:11/21/03ISR Number: 4238857-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0314631A  
 Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged dosage text	11 DAY	Localised Oedema Throat Irritation Urticaria Generalised		Zyban  Tetrazepam	PS  C	Glaxosmithkline	ORAL  ORAL

Date:11/21/03ISR Number: 4238866-0Report Type:Expedited (15-DaCompany Report #SK-GLAXOSMITHKLINE-B0315050A  
 Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Single Initial or Prolonged dose	1 DAY	Angina Unstable Chest Pain Hypertension Tension Headache		Wellbutrin  Augmentin	PS  C	Glaxosmithkline	ORAL  ORAL
625MG Twice per day	6 DAY						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/21/03ISR Number: 4238877-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0315237A  
 Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day 22 DAY	Supraventricular		Zyban	PS	Glaxosmithkline	ORAL
Other		Tachycardia		Salbutamol	C	Glaxosmithkline	
3 YR				Beclomethasone Dipropionate	C	Glaxosmithkline	
UNKNOWN		4 YR		Salmeterol	C	Glaxosmithkline	
2 YR				Combivent	C		
5 MON				Quinine Sulphate	C	Glaxosmithkline	
UNKNOWN	1TAB Per day	5 MON		Montelukast	C		
UNKNOWN	1TAB Per day	4 MON		Co-Codamol	C		
UNKNOWN		1 MON		Barrier Cream	C		
873 DAY							

Date:11/21/03ISR Number: 4238879-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0315354A  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Unknown 6 WK	Hallucination		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged TRANSDERMAL		Paranoia		Corticoid Ointment	C		
		Persecutory Delusion					

Date:11/24/03ISR Number: 4239987-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0440346A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day	Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:11/24/03ISR Number: 4239988-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0440373A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
		Medication Error Pharmaceutical Product Complaint					

Date:11/25/03ISR Number: 4240735-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383113A  
Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest		Digoxin	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged UNKNOWN		Completed Suicide Intentional Misuse		Bupropion Doxazosin	SS SS	Glaxosmithkline	ORAL
		Multiple Drug Overdose Nodal Arrhythmia Urine Output Decreased		Lisinopril Warfarin	SS SS	Glaxosmithkline	

Date:11/25/03ISR Number: 4240736-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383128A  
Age:52 YR Gender:Female I/FU:F

Outcome	PT
Death	Aggression
Hospitalization - Initial or Prolonged	Completed Suicide Convulsion Drug Ineffective Intentional Misuse

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Respiratory Arrest Respiratory Rate Decreased	Report Source	Product	Role	Manufacturer	Route
UNKNOWN				Bupropion Fluvoxamine	PS SS	Glaxosmithkline	ORAL
UNKNOWN				Ethanol	SS		

Date:11/25/03ISR Number: 4240737-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383130A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coma Convulsion		Bupropion Methylphenidate	PS SS	Glaxosmithkline	ORAL
UNKNOWN		Depressed Level Of Consciousness Drug Ineffective Intentional Misuse		Librax Montelukast Synthroid	C C C	Glaxosmithkline	

Date:11/25/03ISR Number: 4240738-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383132A  
Age:67 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Creatinine Increased		Bupropion Sertraline	PS SS	Glaxosmithkline	ORAL
UNKNOWN		Blood Glucose Increased Cardio-Respiratory Arrest Coma Completed Suicide Depressed Level Of Consciousness Dyskinesia General Physical Health Deterioration Intentional Misuse Lethargy Muscle Rigidity					

Pulmonary Congestion  
Tonic Clonic Movements

Date:11/25/03ISR Number: 4240739-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383133A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	900MG Single	Acute Respiratory		Bupropion	PS	Glaxosmithkline	ORAL
Hospitalization - dose		Distress Syndrome					
Initial or Prolonged UNKNOWN	60TAB	Aspiration Single Coma		Topiramate	SS		
dose		Completed Suicide Convulsion		Paracetamol + Propoxyphene	SS		
UNKNOWN	60TAB	Single Depressed Level Of Consciousness Hypotension Intentional Misuse Multiple Drug Overdose					
dose							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/25/03ISR Number: 4240741-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383134A  
Age:19 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Areflexia		Bupropion	PS	Glaxosmithkline	ORAL
UNKNOWN		Cardio-Respiratory Arrest		Valproic Acid	SS		
UNKNOWN		Coma		Olanzapine	SS		
		Completed Suicide					
		Convulsion					
		Depressed Level Of Consciousness					
		Electroencephalogram Abnormal					
		Heart Rate Increased					
		Intentional Misuse					
		Pyrexia					

Date:11/25/03ISR Number: 4240742-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383136A  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Bupropion	PS	Glaxosmithkline	ORAL
UNKNOWN		Cardio-Respiratory Arrest		Doxepin	SS		
		Completed Suicide					
		Convulsion					
		Intentional Misuse					
		Respiratory Arrest					

Date:11/25/03ISR Number: 4240743-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383142A  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest		Bupropion	PS	Glaxosmithkline	ORAL
30TAB Single dose		Coma					



UNKNOWN		Completed Suicide	Aspirin	SS	Glaxosmithkline
UNKNOWN	40TAB	Confusional State Single Intentional Misuse	Olanzapine	SS	
dose		Multiple Drug Overdose Postictal State Pyrexia Tachycardia			

Date:11/25/03ISR Number: 4240744-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383144A  
Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia Supraventricular		Bupropion Warfarin	PS SS	Glaxosmithkline Glaxosmithkline	ORAL
UNKNOWN		Completed Suicide		Carbamazepine	SS		
UNKNOWN		Depressed Level Of Consciousness Intentional Misuse International Normalised Ratio Increased Multiple Drug Overdose Tachycardia		Risperidone Sertraline Marijuana Cocaine	SS SS C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/25/03ISR Number: 4240745-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383149A  
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Accidental Overdose	Bupropion	PS	Glaxosmithkline	ORAL
			Blood Sodium Increased	Risperidone	SS		
UNKNOWN			Cardiac Arrest	Diphenhydramine	SS		
UNKNOWN			Disseminated Intravascular Coagulation Feeling Abnormal Haematocrit Decreased Medication Error Mental Status Changes Pyrexia Staring Tremor				

Date:11/25/03ISR Number: 4240747-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439228A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Medication Error Pharmaceutical Product Complaint	Wellbutrin	PS	Glaxosmithkline	

Date:11/25/03ISR Number: 4240750-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0440827A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Medication Error Pharmaceutical Product Complaint	Wellbutrin	PS	Glaxosmithkline	

Date:11/25/03ISR Number: 4240751-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0440829A  
 Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
		Claustrophobia		Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day							
		Confusional State		Xanax	C		
		Convulsion		Vitamins	C		
		Disorientation					
		Disturbance In Attention					
		Dizziness					
		Feeling Abnormal					
		Flashback					
		Loss Of Consciousness					
		Palpitations					
		Therapeutic Response					
		Unexpected					
		Weight Decreased					

Date:11/25/03ISR Number: 4240752-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0440836A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	
				Wellbutrin	SS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/25/03ISR Number: 4240765-5Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0311195A  
Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	34 DAY	Convulsion		Zyban	PS	Glaxosmithkline	ORAL
		Disorientation					
1MG Per day		Dizziness		Oestradiol Valerate	C		ORAL
10MG Per day		Loss Of Consciousness		Lisinopril	C		ORAL
RESPIRATORY (INHALATION)		Visual Disturbance		Salbutamol	C	Glaxosmithkline	
		Vomiting					

Date:11/25/03ISR Number: 4240768-0Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0314049A  
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	22 DAY	Psoriasis		Zyban	PS	Glaxosmithkline	ORAL

Date:11/25/03ISR Number: 4240771-0Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0314643A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Level Increased		Zyban	PS	Glaxosmithkline	ORAL
		Epilepsy		Librax	C		ORAL
		Sudden Death		Levothyrox	C	Glaxosmithkline	ORAL
				Catapressan	C		
				Agreal	C		ORAL

Date:11/25/03ISR Number: 4240773-4Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0314874A  
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Fall		Zyban	PS	Glaxosmithkline	ORAL
150MG See							
Initial or Prolonged		Tendon Rupture					
dosage text	38	DAY					
				Avandia	C	Glaxosmithkline	ORAL
				Metformine	C		ORAL

Date:11/25/03ISR Number: 4240777-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0315370A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dermatitis Bullous		Zyban	PS	Glaxosmithkline	ORAL
150MG See							
Initial or Prolonged		Rash Morbilliform					
dosage text	17	DAY					
		Rash Vesicular					
		Scar					
		Skin Exfoliation					

Date:11/25/03ISR Number: 4241927-3Report Type:Direct Company Report #CTU 206878  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Erythema Multiforme		Wellbutrin 100 Mg			
Intervention to		Rash Erythematous		Tablets	PS		ORAL
PO BID							
Prevent Permanent		Stevens-Johnson Syndrome					
Impairment/Damage							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/25/03ISR Number: 4242123-6Report Type:Direct  
Age:10 YR Gender:Female I/FU:I

Company Report #CTU 206884

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Constipation		Bupropion Sr	PS		
100 MG Q DAY							
Required		Urinary Retention		Trazodone	C		
Intervention to							
Prevent Permanent							
Impairment/Damage							

Date:11/25/03ISR Number: 4242177-7Report Type:Expedited (15-DaCompany Report #2003-03934  
Age:28 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Literature Health Professional	Verapamil (Verapamil Hydrochloride)			
				Unknown	PS	Watson Laboratories	
				Venlafaxine (Venlafaxine)	SS		
				Bupropion (Bupropion)	SS		

Date:11/25/03ISR Number: 4242309-0Report Type:Expedited (15-DaCompany Report #2003-04021  
Age:47 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardio-Respiratory Arrest	Literature Health Professional	Alprazolam (Watson Laboratories)			
				(Alprazolam) Tablet	PS		
UNKNOWN	UNK, UNK, UNK			Methadone (Methadone)	SS		
UNKNOWN	UNK, UNK, UNK			Bupropion (Bupropion)	SS		
UNKNOWN	UNK, UNK, UNK						

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Acidosis	Study	Oxycodone			
Initial or Prolonged	Depressed Level Of	Health	Hydrochloride			
Other	Consciousness	Professional	(Similar To Nda			
	Hypotension	Other	20-553) (Oxycodone			
	Multiple Drug Overdose		Hydrochloride)	PS		
			Hydrocodone			
			W/Acetaminophen			
			(Paracetamol,			
			Hydrocodone			
			Bitartrate)	SS		
			Benzodiazepine			
			Derivates	SS		
			Cox-2 Inhibitor	SS		
			Gastrointestinal			
			Prep	SS		
			Laxatives	SS		
			Anticonvulsant	SS		
			Muscle Relaxants	SS		
			Bupropion(Amfebutamo			
			ne)	SS		
			Cyproheptadine(Cypro			

Freedom Of Information (FOI) Report

heptadine) SS  
 Antipsychotics SS  
 Proton Pump  
 Inhibitor SS  
 Ssri SS  
 Trazodone(Trazodone) SS

Date:11/25/03ISR Number: 4242470-8Report Type:Expedited (15-DaCompany Report #03P-163-0241040-00  
 Age:28 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Verapamil (Isoptin) (Verapamil) (Verapamil)			
				Venlafaxine	PS		
				Bupropion	SS		

Date:11/25/03ISR Number: 4242499-XReport Type:Expedited (15-DaCompany Report #B0313503A  
 Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Increased Appetite Sleep Walking	Foreign Literature Health Professional	Zyban (Bupropion Hydrochloride)			
					PS		ORAL

150 MG / SEE

DOSAGE TEXT

Date:11/25/03ISR Number: 4242501-5Report Type:Expedited (15-DaCompany Report #B0314910A  
 Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Disability		Angioneurotic Oedema Arthralgia C-Reactive Protein Increased Chest Pain Dizziness Electrocardiogram T Wave	Foreign Literature Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)			
					PS		



Inversion  
Liver Function Test  
Abnormal  
Myalgia  
Neutrophilia  
Oedema Peripheral  
Palpitations  
Pyrexia  
Serum Sickness  
Sinus Tachycardia  
Skin Discolouration  
Urticaria

Date:11/26/03ISR Number: 4243980-XReport Type:Expedited (15-DaCompany Report #2003-06064  
Age:26 YR Gender:Male I/FU:I

Outcome PT  
Other Blood Phosphorus  
Decreased  
Cholelithiasis  
Dyspepsia

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hepatitis					
		Nausea					
		Pancreatitis					
		Vomiting	Health	Viread	PS		ORAL
300 MG QD PO			Professional	Combivir	SS		ORAL
450 MG BID PO				Wellbutrin	SS		
				Ambien	C		
				Effexor	C		
				Asacol	C		
				Clindagel	C		
				Elidel	C		
				Clotrim/Beta	C		
				Protopic	C		
				Flonase	C		

Date:11/26/03ISR Number: 4244193-8Report Type:Expedited (15-DaCompany Report #2003039894

Age:45 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature	Doxepin (Caps)			
Other		Completed Suicide	Health	(Doxepin)	PS		ORAL
ORAL		Multiple Drug Overdose	Professional	Bupropion			
				(Bupropion)	SS		ORAL
				Clorazepate			
				Dipotassium			
				(Clorazepate			
				Dipotassium)	SS		ORAL
ORAL							

Date:11/28/03ISR Number: 4244975-2Report Type:Expedited (15-DaCompany Report #2003-04125

Age:45 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature	Clorazepate(Watson			
			Health	Laboratories)(Cloraz			
			Professional	epate Dipotassium)			

Tablet, 3.75mg	PS	
Doxepin(Watson Laboratories)(Doxepin Hydrochloride)		
Capsule	SS	Watson Laboratories
Bupropion(Long Acting)	SS	

Date:11/28/03ISR Number: 4245222-8Report Type:Expedited (15-DaCompany Report #2003-04088  
 Age:38 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature Health Professional	Alprazolam (Watson Laboratories)(Alprazolam)Tablet	PS	Watson Laboratories	
				Bupropion(Bupropion)	SS		
				Acetaminophen(Paracetamol)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/28/03 ISR Number: 4245390-8 Report Type:Expedited (15-DaCompany Report #USA-2003-0006512  
 Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Consumer	Oxycontin Tablets			
		Hepatomegaly	Health	(Oxycodone			
		Portal Triaditis	Professional	Hydrochloride) Cr			
		Postoperative Infection	Other	Tablet	PS		ORAL
40 MG, BID, ORAL; 20 MG		Pulmonary Congestion					
		Pulmonary Oedema		Celexa (Citalopram			
				Hydrobromide)	SS		
				Carisoprodol			
				(Carisoprodol)	SS		
				Meprobamate			
				(Meprobamate)	SS		
				Bupropion			
				(Amfebutamone)	SS		
				Biaxin			
				(Clarithromycin)			
				Tablet	C		

Date:12/01/03 ISR Number: 4243427-3 Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0289128A  
 Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 40TABS Single		Blood Creatine		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged dose	1 DAY	Phosphokinase Increased					
1PACK Single		Dizziness		Zopiclone	SS		ORAL
dose	1 DAY	Dysarthria					
1PACK Single		Intentional Misuse		Valium	SS		ORAL
dose	1 DAY	Multiple Drug Overdose					
		Oxygen Saturation Decreased					
		Sinus Tachycardia					
		Somnolence					

Vomiting

Date:12/01/03ISR Number: 4243430-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0300655A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	3.7G per day 1 DAY	Agitation		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged	6.25MG per day 1 DAY	Coordination Abnormal Dysarthria Hallucinations, Mixed Intentional Misuse Suicide Attempt Tremor		Xanax	C		ORAL

Date:12/01/03ISR Number: 4243432-7Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0308326A  
Age:65 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG per day 1 DAY	Myocardial Infarction		Zyban	PS	Glaxosmithkline	
1UNIT See dosage text				Tobacco	C		
20MG Twice per day				Isosorbide	C		ORAL
150MG Per day				Aspirin	C	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/01/03ISR Number: 4243455-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0316048A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Clumsiness		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day		Escherichia Infection		Urbanyl	SS		ORAL
5MG Three		Pyelonephritis					
times per day	6	Sensation Of Heaviness					
			DAY				

Date:12/02/03ISR Number: 4243884-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0315354A  
Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anxiety		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	6	Delirium		Corticoid Ointment	SS		
Initial or Prolonged		Hallucination					
TRANSDERMAL		Paranoia					
		Persecutory Delusion					
		Psychotic Disorder					

Date:12/02/03ISR Number: 4243908-2Report Type:Expedited (15-DaCompany Report #DK-GLAXOSMITHKLINE-B0316211A  
Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Atrial Fibrillation		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							
	51		DAY				

Date:12/02/03ISR Number: 4244818-7Report Type:Direct Company Report #CTU 207180  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	200 MG QD	Balance Disorder Dry Mouth		Amantadine 100mg Apothecon	PS	Apothecon	ORAL
Required ORAL		Heart Rate Increased					
Intervention to Prevent Permanent 100MG HS ORAL Impairment/Damage		Orthostatic Hypotension Thinking Abnormal Tremor		Bupropion Sr 100mg Glaxowellcome .. .. Lorazepam Paroxetine Ranitidine Quetiapine Vit E Vit C Ecasa Ibuprofen Loratadine Meclizine Hcl	SS  C C C C C C C C C C	Glaxowellcome	ORAL

Date:12/02/03ISR Number: 4246222-4Report Type:Expedited (15-DaCompany Report #USA-2003-0010936  
Age:57 YR Gender:Female I/FU:I

Outcome  
Death

PT  
Accidental Overdose  
Coronary Artery  
Atherosclerosis  
Drowning  
Drug Toxicity

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Glomerulosclerosis Hepatomegaly Pyelonephritis Chronic	Report Source	Product	Role	Manufacturer	Route
			Health	Oxycodone			
			Professional	Hydrochloride	PS		
			Other	Hydrocodone			
				Bitartrate (Similar To Ind 59,175)			
				(Hydrocodone Bitartrate)	SS		
				Chlorpheniramine (Chlorpheniramine)	SS		
				Citalopram (Citalopram)	SS		
				Diazepam (Diazepam)	SS		
				Lorazepam (Lorazepam)	SS		
				Mirtazapine (Mirtazapine)	SS		
				Bupropion (Amfebutamone)	SS		
				Oxazepam (Oxazepam)	SS		
				Temazepam (Temazepam)	SS		
				Verapamil (Verapamil)	SS		
				Triazolam (Triazolam)	SS		

Date:12/03/03ISR Number: 4244527-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0383141A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Acute Pulmonary Oedema		Bupropion	PS	Glaxosmithkline	ORAL
		Cardio-Respiratory Arrest		Acetaminophen +			
		Completed Suicide		Hydrocodone	SS		
UNKNOWN							
		Intentional Misuse		Propoxyphene	SS		
UNKNOWN							
		Pulmonary Congestion		Doxepin	SS		
				Fluoxetine	SS		
				Zolpidem	SS		
				Ammonia	SS		



Bleach SS  
Nicotine C Glaxosmithkline

Date:12/03/03ISR Number: 4244532-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441016A  
Age:60 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150MG See	Anal Fissure		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged dosage text 6 WK	Constipation					
	Proctalgia		Vitamins Fosamax	C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/03/03ISR Number: 4244544-4Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0315394A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Per day	3	DAY	Anaphylactic Reaction	Zyban	PS	Glaxosmithkline	ORAL
			Asthma				

Date:12/03/03ISR Number: 4244545-6Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0315395A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300MG per day	3	WK	Joint Swelling	Zyban	PS	Glaxosmithkline	ORAL
			Odynophagia				
			Pruritus				
			Pyrexia				
			Rash				
			Serum Sickness				

Date:12/03/03ISR Number: 4244547-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0315632A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
			Feeling Abnormal	Zyban	PS	Glaxosmithkline	
			Hyperhidrosis				
			Insomnia				
			Loss Of Consciousness				

Date:12/03/03ISR Number: 4244548-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0315793A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
150MG Unknown			Asthenia	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged			Chest Pain				
			Dysaesthesia				

Headache  
Hypoaesthesia  
Speech Disorder  
Vertigo

Date:12/03/03ISR Number: 4246221-2Report Type:Direct  
Age:52 YR Gender:Female I/FU:I

Company Report #CTU 207438

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Wellbutrin 200 Mg			
200 MG BID		Blood Pressure Increased		Per Patient	PS		ORAL
ORAL		Heart Rate Increased					
500 MG ONC		Nervousness		Cipro 500 Mg Per			
ORAL				Patient	SS		ORAL

Date:12/03/03ISR Number: 4246369-2Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 207418

Outcome  
Required  
Intervention to  
Prevent Permanent

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG PO QD;		Blister		Bupropion	PS		ORAL
75 MG PO BID		Candidiasis					
		Rash		Luvox	C		
		Rash Papular		Fosamax	C		
				Lasix	C		
				Lipitor	C		
				Asa	C		

Date:12/03/03ISR Number: 4246919-6Report Type:Expedited (15-DaCompany Report #DEWYE424826NOV03  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Study	Trevilor			
Other		Tremor		(Venlafaxine Hydrochloride, Tablet, 0)	PS		ORAL
SEE IMAGE	46 DAY			Zyban (Amfebutamone Hydrochloride, , 0)	SS		ORAL
SEE IMAGE	11 DAY			Zopiclone (Zopiclone)	C		

Date:12/03/03ISR Number: 4247125-1Report Type:Direct Company Report #CTU 207333  
 Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abnormal Behaviour		Wellbutrin	PS		
		Aggression					
		Alcoholism					
		Cardiac Arrest					
		Coma					
		Convulsion					
		Delusion					
		Depressed Level Of					

Consciousness  
 Disorientation  
 Drug Abuser  
 Eye Rolling  
 Fall  
 Feeling Hot  
 Grunting  
 Incoherent  
 Mydriasis  
 Respiration Abnormal

Date:12/03/03ISR Number: 4247187-1Report Type:Direct  
 Age:61 YR Gender:Male I/FU:I

Company Report #CTU 207343

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged	150 MG DAY	Anxiety Blood Pressure Increased Nightmare		Wellbutrin Sr Then Xl 150 Mg Glaxosmithkline	PS	Glaxosmithkline	ORAL
Other ORAL	300 MG DAY			Wellbutrin Xl 300 Mg Glaxosmithkline	SS	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/03/03ISR Number: 4247502-9Report Type:Expedited (15-DaCompany Report #DEWYE424826NOV03

Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Study	Trevilor (Venlafaxine Hydrochloride, Tablet, 0)	PS		ORAL
75 MG 1X PER							
1 DAY ORAL;							
SEE IMAGE	46	DAY					
150 MG 1X PER							
1 DAY ORAL;							
SEE IMAGE	11	DAY		Zyban (Amfebutamone Hydrochloride, , 0)	SS		ORAL
				Zopiclone (Zopiclone)	C		

Date:12/04/03ISR Number: 4245720-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441481A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Intentional Misuse		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN	9000MG	Single					
dose							
		Suicide Attempt		Amphetamines	SS		

Date:12/04/03ISR Number: 4245723-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441720A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Congenital Anomaly		Cleft Lip		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN		Drug Exposure During		Neurontin	SS		

Pregnancy

Date:12/04/03ISR Number: 4245724-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441723A  
 Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs Drug Exposure During Pregnancy		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:12/04/03ISR Number: 4245726-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0298802A  
 Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG Per day 4 DAY		Insomnia		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged 500MG Three times per day		Malaise  Painful Respiration		Dafalgan	SS	Glaxosmithkline	ORAL
100MG Per day		Serotonin Syndrome		Granudoxy	SS	Glaxosmithkline	ORAL
20MG Per day				Seropram	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/04/03ISR Number: 4245728-1Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0312169A  
 Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Zyban	PS	Glaxosmithkline	ORAL

Date:12/04/03ISR Number: 4245732-3Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0316104A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1 WK Initial or Prolonged		Aggression Agitation Bipolar I Disorder Depression Hallucination Mania Mood Swings Suicide Attempt		Zyban	PS	Glaxosmithkline	ORAL

Date:12/04/03ISR Number: 4245733-5Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0316198A  
 Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Per day 11 DAY		Anxiety		Zyban	PS	Glaxosmithkline	ORAL
TRANSDERMAL per day	21MG Twice	Irritability Suicide Attempt		Nicopatch	C	Glaxosmithkline	

Date:12/04/03ISR Number: 4245734-7Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0042190A  
 Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 1TAB Twice		Anxiety		Zyban	PS	Glaxosmithkline	ORAL



per day            17    DAY            Confusional State  
   Depression  
   Dizziness  
   Dry Mouth  
   Dysphonia  
   Fatigue  
   Hyperhidrosis  
   Muscle Spasms  
   Nervousness  
   Rash  
   Stomatitis

Date:12/05/03ISR Number: 4245940-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441718A  
Age:            Gender:Female            I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin	PS	Glaxosmithkline	ORAL
1    YR		Drug Exposure During Pregnancy					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/05/03ISR Number: 4245943-7Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0313519A  
Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Disability		Depressed Level Of Consciousness Grand Mal Convulsion Incontinence Overdose Tongue Biting		Zyban	PS	Glaxosmithkline	ORAL

Date:12/05/03ISR Number: 4245956-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0316469A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Per day		Psoriasis		Zyban	PS	Glaxosmithkline	ORAL

Date:12/08/03ISR Number: 4246798-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0312762A  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG See Hospitalization - dosage text 3 WK Initial or Prolonged		Agitation Anxiety Insomnia Intentional Misuse Suicidal Ideation Suicide Attempt		Zyban	PS	Glaxosmithkline	ORAL

Date:12/08/03ISR Number: 4248380-4Report Type:Direct Company Report #CTU 207655  
Age:79 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Pruritus Rash		Wellbutrin Sr 150 Mg Dsm Pharm/Glaxo			

150 MG 2

TIMES ORAL

Smithkline

PS

Dsm Pharm/Glaxo  
Smithkline

ORAL

Wellbutrin Xl 300 Mg  
Dsm Pharm/Glaxo  
Smithkline

SS

Dsm Pharm/Glaxo  
Smithkline

ORAL

300 MG 1 PER

DAY ORAL

Date:12/09/03ISR Number: 4247950-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0400608A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blister		Flonase	PS	Glaxosmithkline	NASAL
DAY		Burning Sensation		Wellbutrin	SS	Glaxosmithkline	ORAL
8	MON	Erythema Hypersensitivity Local Swelling Pyrexia Swelling Face					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/09/03ISR Number: 4249855-4Report Type:Expedited (15-DaCompany Report #HQWYE843007NOV03

Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion Drug Withdrawal Syndrome Intentional Misuse Loss Of Consciousness Suicide Attempt	Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule Extended Release)	PS		ORAL
OVERDOSE							
AMOUNT,							
UNKNOWN; ORAL							
OVERDOSE				Atenolol (Atenolol, )	SS		
AMOUNT							
UNKNOWN							
OVERDOSE				Bactrim (Sulfamethoxazole/Tr imethoprim)	SS		
AMOUNT							
UNKNOWN							
OVERDOSE				Benzonatate (Benzontate)	SS		
AMOUNT							
UNKNOWN							
OVERDOSE				Flavoxate (Flavoxate)	SS		
AMOUNT							
UNKNOWN							
OVERDOSE				Indocin (Indometacin)	SS		

AMOUNT

UNKNOWN

Lexapro  
(Escitalopram) SS

OVERDOSE

AMOUNT

UNKNOWN

Minocycline  
(Minocycline,  
Unspec) SS

OVERDOSE

AMOUNT

UNKNOWN

Naprosyn (Naproxen) SS

OVERDOSE

AMOUNT

UNKNOWN

Willbutrin  
(Amfebutamone  
Hydrochloride) SS

OVERDOSE

AMOUNT

UNKNOWN

Date:12/11/03ISR Number: 4249634-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0354550A

Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Discomfort
Initial or Prolonged	Aggression
Other	Agitation
	Anger
	Anhedonia
	Anorexia
	Anxiety

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Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
20MG In the morning		Asthenia Confusional State Coordination Abnormal				
		Delusion	Paxil	PS	Glaxosmithkline	ORAL
150MG Twice per day		Depressed Mood				
		Depression	Wellbutrin	SS	Glaxosmithkline	ORAL
		Diarrhoea				
		Disturbance In Attention	Ritalin	C		
		Dizziness	Amoxicillin	C	Glaxosmithkline	
		Drug Ineffective				
		Drug Withdrawal Syndrome				
		Fatigue				
		Gait Disturbance				
		Gingival Infection				
		Hallucinations, Mixed				
		Homicidal Ideation				
		Influenza				
		Insomnia				
		Intentional Self-Injury				
		Irritability				
		Lethargy				
		Malaise				
		Memory Impairment				
		Myalgia				
		Nausea				
		Nightmare				
		Obsessive Thoughts				
		Obsessive-Compulsive Disorder				
		Overdose				
		Paraesthesia				
		Paranoia				
		Psychotic Disorder				
		Pyrexia				
		Sleep Disorder				
		Suicidal Ideation				
		Suicide Attempt				
		Tremor				
		Vertigo				
		Vomiting				

Age:23 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Amnesia	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged	Constipation	Professional	Lithobid	SS	Glaxosmithkline	
1200MG Per						
day	Depressed Mood					
30MG Per day	Fall		Lexapro	C		
150MG At	Fatigue		Nortriptyline	C		
night	Grand Mal Convulsion					
	Loss Of Consciousness					
	Medication Error					
	Tremor					
	Weight Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/11/03ISR Number: 4249662-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0316325A

Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Chest Discomfort		Amoxil	PS	Glaxosmithkline	
UNKNOWN	250MG Per day 1 DAY						
Initial or Prolonged		Dyspnoea		Zyban	SS	Glaxosmithkline	ORAL
1 WK		Erythema					
		Swelling Face					

Date:12/11/03ISR Number: 4249666-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0316749A

Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Aggression		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day 1 DAY		Amnesia					
		Confusional State					
		Disinhibition					

Date:12/11/03ISR Number: 4249673-7Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0317068A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Affect Lability		Zyban	PS	Glaxosmithkline	
		Belligerence		Clonazepam	C		
2MG Three		Homicidal Ideation					
times per day		Intentional Self-Injury					

Date:12/11/03ISR Number: 4251379-5Report Type:Expedited (15-DaCompany Report #S03-FRA-05070-01

Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Diarrhoea	Foreign	Seropram(Citalopram			



Initial or Prolonged 20 MG QD PO	Flushing	Health	Hydrobromide)	PS	ORAL
	Insomnia Malaise	Professional Other	Dafalgan (Paracetamol)	SS	ORAL
500 MG TID PO	Respiratory Disorder Vomiting		Zyban (Bupropion Hydrochloride)	SS	ORAL
1 QD PO			Granudoxy (Doxycycline Hyclate)	SS	ORAL
100 MG QD PO					

Date:12/12/03ISR Number: 4250924-3Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0316714A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Arthralgia Fatigue		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
77 DAY		Headache Pyrexia Skin Disorder		Unspecified Medication	C		

Date:12/12/03ISR Number: 4251791-4Report Type:Expedited (15-DaCompany Report #S03-USA-05003-01  
Age:42 YR Gender:Female I/FU:I

Outcome Other	PT Contusion Convulsion
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Pharmaceutical Product  
per day 1 WK  
Complaint

Date:12/15/03ISR Number: 4251472-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0442750A  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Bupropion	PS	Glaxosmithkline	
300MG per day		Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus					

Date:12/15/03ISR Number: 4251473-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0442800A  
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Medication Error	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
per day			Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/15/03ISR Number: 4252023-3Report Type:Direct  
Age:45 YR Gender:Female I/FU:I

Company Report #CTU 208108

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased		Zyban 150 Mg Bid	PS		
150 MG BID		Condition Aggravated					

Date:12/15/03ISR Number: 4252085-3Report Type:Direct  
Age:16 YR Gender:Male I/FU:I

Company Report #CTU 208011

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Anaphylactic Reaction		Wellbutrin Xl 150 Mg			
		Inflammation		Glaxxo	PS	Glaxxo	ORAL
TWO TABLET		Pharmaceutical Product					
ONCE DAILY		Complaint					
ORAL		Tracheal Disorder					
		Tracheal Oedema					

Date:12/15/03ISR Number: 4252135-4Report Type:Direct  
Age:28 YR Gender:Female I/FU:I

Company Report #CTU 208031

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Oedema Peripheral		Zyban 150 Mg			
150 MG BID		Rash		Glaxosmithkline	PS	Glaxosmithkline	ORAL
ORAL		Urticaria					

Date:12/16/03ISR Number: 4252212-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0443093A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Paxil	PS	Glaxosmithkline	ORAL

UNKNOWN	Bipolar I Disorder	Wellbutrin	SS	Glaxosmithkline	ORAL
	Drug Ineffective	Lithium	SS	Glaxosmithkline	
UNKNOWN	Paranoia	Prozac	SS		
	2 YR				
	Suicidal Ideation				

Date:12/16/03ISR Number: 4252218-9Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0314554A  
 Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypersensitivity		Zyban	PS	Glaxosmithkline	ORAL
		Medication Error		Doxycycline	SS		ORAL
		Oropharyngeal Swelling					

Date:12/16/03ISR Number: 4252223-2Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0316104A  
 Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Aggression	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
Hospitalization -		Agitation	Professional				
per day 19 DAY							
Initial or Prolonged		Depression		Qvar	C	Glaxosmithkline	
RESPIRATORY							
(INHALATION)		Mood Swings					
RESPIRATORY		Suicide Attempt		Ventolin	C	Glaxosmithkline	
(INHALATION)							
				Alcohol	C		



Other 150MG Twice	Neutropenia	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	MON				
UNKNOWN		Adriamycin	SS		
UNKNOWN		Taxol	SS		

Date:12/17/03ISR Number: 4252745-4Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0316627A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 30TAB Unknown Initial or Prolonged		Convulsion		Zyban	PS	Glaxosmithkline	ORAL
		Intentional Misuse Intentional Self-Injury Loss Of Consciousness Somnolence Suicide Attempt					

Date:12/17/03ISR Number: 4252752-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0317217A  
 Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Unknown 41 DAY		Malaise		Zyban	PS	Glaxosmithkline	ORAL
		Myocardial Infarction Sudden Death		Esperial Tranxene	C C		ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/17/03ISR Number: 4253781-4Report Type:Direct  
Age: Gender: I/FU:I

Company Report #CTU 208333

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG PO QD		Accidental Overdose		Bupropion	PS		ORAL
Initial or Prolonged 60 MG PO Q AM		Confusional State		Citalopram	SS		ORAL
		Dysarthria Gait Disturbance Mental Status Changes		Diazepam 5mg Po Qid Prn Anxiety	SS		ORAL

Date:12/17/03ISR Number: 4253869-8Report Type:Direct  
Age:31 YR Gender:Male I/FU:I

Company Report #CTU 208279

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG PO QD		Abdominal Pain Upper		Wellbutrin 150 Mg	PS		ORAL
		Suicide Attempt					

Date:12/17/03ISR Number: 4253882-0Report Type:Direct  
Age:38 YR Gender:Male I/FU:I

Company Report #CTU 208277

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG BID PO		Overdose Suicide Attempt		Wellbutrin Sr 150 Mg	PS		ORAL
				Prevacid Beano	C C		

Date:12/17/03ISR Number: 4254661-0Report Type:Expedited (15-DaCompany Report #B0316252A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alcohol Poisoning Coma	Literature Health	Unspecified Tablet (Bupropion			



ORAL	Completed Suicide	Professional	Hydrochloride)	PS	ORAL
	Pulmonary Congestion		Ethanol (Formulation		
	Pulmonary Oedema		Unknown) (Alcohol)	SS	
			Fluoxetine	C	
			Flurazepam	C	
			Lorazepam	C	
			Ibuprofen	C	
			Salbutamol Sulphate	C	

Date:12/17/03ISR Number: 4271077-1Report Type:Periodic Company Report #KII-2003-0004053  
 Age:52 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Other SEE TEXT,	Depressed Level Of Consciousness Multiple Drug Overdose	Health Professional	Oxycontin Tablets (Oxycodone Hydrochloride)	PS		ORAL
ORAL	Nodal Rhythm					
SEE TEXT,	Somnolence		Wellbutrin (Amfebutamone Hydrochloride)	SS		ORAL
ORAL			Verapamil (Verapamil)	SS		ORAL
SEE TEXT,						
ORAL			Hydrochlorothiazide (Hydrochlorothiazide)			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

SEE TEXT,	)	SS	ORAL
ORAL			
SEE TEXT,	Zoloft (Sertraline Hydrochloride)	SS	ORAL
ORAL			
SEE TEXT,	Neuroton (Citicoline Sodium)	SS	ORAL
ORAL			
ORAL	Tricyclic Antidepressants	SS	ORAL
	Cannabis (Cannabis)	SS	

Date:12/17/03ISR Number: 4272416-8Report Type:Periodic Company Report #USA-2003-0008572  
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose Multiple Drug Overdose	Health Professional Other	Oxycodone Hydrochloride Methadone (Methadone) Bupropion (Amfebutamone) Caffeine (Caffeine) Diazepam (Diazepam0 Nicotine (Nicotine) Oxymorphone (Oxymorphone)	PS SS SS SS SS		

Date:12/18/03ISR Number: 4253919-9Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0084018A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Per day	22 DAY	Dysgeusia Eating Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL

Oral Discomfort  
Weight Decreased

Date:12/18/03ISR Number: 4253921-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441720A  
Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Congenital Anomaly	Cleft Lip		Wellbutrin	PS	Glaxosmithkline	
	Drug Exposure During		Neurontin	SS		
UNKNOWN	Pregnancy		Multivitamins	C		
			Alcohol	C		
			Allegra	C		

Date:12/18/03ISR Number: 4253922-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0442426A  
Age:54 YR Gender:Male I/FU:F

Outcome	PT
Life-Threatening	Abnormal Behaviour
Hospitalization -	Anxiety
Initial or Prolonged	Chest Pain
	Circulatory Collapse
	Disorientation
	Heart Rate Increased
	Myocardial Infarction

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Panic Disorder Stress Ventricular Tachycardia	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
51	DAY			Glucosamine	C		ORAL
				Nexium	C		ORAL

Date:12/18/03ISR Number: 4273966-0Report Type:Periodic Company Report #HQWYE078213AUG03  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release	PS		
Hospitalization - Initial or Prolonged		Condition Aggravated Mania	Consumer				

ORAL/SEE

IMAGE

Wellbutrin (Amfebutamone Hydrochloride,Extend ed Release	SS
Lithobid (Lithium Carbonate)	C

Date:12/19/03ISR Number: 4255081-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0443009A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
		Insomnia Medication Error		Wellbutrin	SS	Glaxosmithkline	ORAL

150MG Per day

Date:12/19/03ISR Number: 4255096-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0317420A  
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day 3 DAY	Completed Suicide		Zyban	PS	Glaxosmithkline	ORAL
		Disinhibition Suicidal Ideation					

Date:12/22/03ISR Number: 4255852-5Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0313883A  
 Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG See dosage text 21 DAY	Amnesia		Zyban	PS	Glaxosmithkline	ORAL
		Crying Insomnia Malaise Petit Mal Epilepsy Visual Disturbance					

Date:12/22/03ISR Number: 4255857-4Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0317324A  
 Age:33 YR Gender:Female I/FU:F

Outcome	PT
Other	Grand Mal Convulsion Loss Of Consciousness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Tongue Biting Urinary Incontinence				
Dose	Duration		Report Source	Product	Role	Manufacturer
10	DAY			Zyban	PS	Glaxosmithkline
				No Concurrent Medication	C	

Date:12/22/03ISR Number: 4257821-8Report Type:Expedited (15-DaCompany Report #B0316387A  
Age:27 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - Initial or Prolonged	Dizziness Grand Mal Convulsion Hallucination, Visual	Foreign Literature Health	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG / UNK  / ORAL	Intentional Misuse  Nausea Sinus Tachycardia Vomiting	Professional				

Date:12/23/03ISR Number: 4257090-9Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0443220A  
Age:29 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Life-Threatening 150MG Twice Hospitalization - per day Initial or Prolonged Other	Eyelid Oedema  Hyperglycaemia  Hypokalaemia Loss Of Control Of Legs Oedema Peripheral Urticaria Generalised	Health  Professional	Zyban	PS	Glaxosmithkline	ORAL

Date:12/23/03ISR Number: 4257114-9Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0317731A  
Age:60 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Sudden Death		Bupropion Hydrochloride	PS	Glaxosmithkline	
	150MG per day	3 DAY			Metoprolol	C		
	100MG per day				Carbasalate Calcium	C		
	100MG per day				Furosemide	C	Glaxosmithkline	
	40MG per day							

Date:12/23/03ISR Number: 4257148-4Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0084018A  
Age: Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability			Burning Sensation	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
	150MG Per day	22 DAY	Dysgeusia	Professional				
			Eating Disorder					
			Weight Decreased					

Date:12/23/03ISR Number: 4257162-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0443236A  
Age:28 YR Gender:Female I/FU:I

Outcome	PT
Other	Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Drugs  
Maternal Drugs Affecting  
Foetus

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300MG per day			Bupropion	PS	Glaxosmithkline	
			Lexapro	C		

Date:12/23/03ISR Number: 4257163-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0443376A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation		Wellbutrin	PS	Glaxosmithkline	
300MG per day		Confusional State Headache Myalgia Serotonin Syndrome		Lexapro	SS		

Date:12/23/03ISR Number: 4257164-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0443378A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation		Wellbutrin	PS	Glaxosmithkline	
300MG per day		Confusional State Headache Myalgia Serotonin Syndrome		Lexapro	SS		

Date:12/23/03ISR Number: 4257166-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0443651A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day	YR	Muscle Twitching					



Pharmaceutical Product  
Complaint

Vitamins  
Calcium  
Advil

C  
C  
C

Glaxosmithkline

Date:12/23/03ISR Number: 4257174-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0443925A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pancreatitis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:12/23/03ISR Number: 4257189-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0315793A  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Abdominal Pain	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG Per day 39 DAY							
Initial or Prolonged		Asthenia Chest Pain Dysaesthesia Headache Hypoaesthesia Malaise Speech Disorder Vertigo	Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/23/03ISR Number: 4257196-4Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0317735A  
 Age:27 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other UNKNOWN		Amnesia 10 DAY Dizziness Feeling Abnormal Hyperventilation Loss Of Consciousness Nausea Pallor Salivary Hypersecretion		Zyban	PS	Glaxosmithkline	

Date:12/23/03ISR Number: 4257212-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0318341A  
 Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Twice per day	67 DAY	Cardiac Arrest Respiratory Arrest Sudden Death		Zyban	PS	Glaxosmithkline	ORAL

Date:12/23/03ISR Number: 4257213-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0318367A  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Per day Disability	3 DAY	Loss Of Consciousness Myocardial Infarction		Zyban	PS	Glaxosmithkline	ORAL

Date:12/23/03ISR Number: 4258205-9Report Type:Direct Company Report #USP 56170  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
TABLET-SUSTAINED RELEASE		Depression Myalgia	Health Professional	Lipitor (Atorvastatin)	PS	Glaxo Smithkline	ORAL
		Sexual Dysfunction		Paroxetine Hydrochloride (Paroxetine Hydrochloride)	SS		ORAL
	20 MG DAILY,			Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		ORAL
	ORAL						
	300 MG (BID),						
	ORAL						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/24/03ISR Number: 4257597-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0444101A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
			Medication Error Pharmaceutical Product Complaint				

Date:12/24/03ISR Number: 4257602-5Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0444142A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
Other			Renal Tubular Necrosis				

Date:12/24/03ISR Number: 4257616-5Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0317878A

Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	
Hospitalization - Initial or Prolonged			Fall Hypoglycaemia Limb Injury	Dothiepin Hydrochloride	C		
75MG per day							

Date:12/24/03ISR Number: 4257623-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0317972A

Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
Life-Threatening 150MG Per day 5 DAY			Blood Glucose Decreased				
Hospitalization - 100MG Twice			Circulatory Collapse	Becotide	C	Glaxosmithkline	
Initial or Prolonged per day			Feeling Abnormal				
				Combivent	C		
				Salbutamol	C	Glaxosmithkline	
				Antiemetic	C		

Date:12/24/03ISR Number: 4257639-6Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0318401A  
Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fatigue		Zyban	PS	Glaxosmithkline	ORAL
300MG per day		Feeling Jittery Sleep Apnoea Syndrome					

Date:12/29/03ISR Number: 4261311-6Report Type:Direct Company Report #CTU 208864  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Simple Partial Seizures		Wellbutrin	PS		

Date:12/30/03ISR Number: 4267021-3Report Type:Periodic Company Report #HQWYE078213AUG03  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Condition Aggravated Mania	Consumer	Effexor Xr (Venlafaxine Hydochloride, Capsule, Extended Release)	PS		ORAL

ORAL; 75 MG

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Freedom Of Information (FOI) Report

1X PER 1 DAY,  
 ORAL; 150 MG  
 1X PER 1 DAY,  
 ORAL; 75 MG

Wellbutrin  
 (Amfebutamone  
 Hydrochloride, ) SS  
 Lithobid (Lithium  
 Carbonate) C  
 Oral Contraceptive  
 Nos (Oral  
 Contraceptive Nos) C

Date:12/31/03ISR Number: 4262581-0Report Type:Expedited (15-DaCompany Report #2003125837  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Burning Sensation Feeling Abnormal	Consumer	Neurontin (Gabapentin)	PS		ORAL
100 MG, ORAL		Myalgia		Geodon (Ziprasidone)	SS		ORAL
ORAL		Nausea		Zoloft (Sertraline)	SS		ORAL
50 MG		Pain					
(DAILY), ORAL				Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		ORAL
ORAL							

Date:01/02/04ISR Number: 4262055-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0490866A  
 Age:2 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 0 DAY		Accidental Exposure		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/02/04    ISR Number: 4264230-4    Report Type:Expedited (15-DaCompany Report #B0317415A  
 Age:40 YR    Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Drug Toxicity Mood Swings Treatment Noncompliance	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Alprazolam (Formulation Unknown) (Alprazolam) Temazepam (Formulation Unknown) (Temazepam)	SS SS		

Date:01/04/04    ISR Number: 4263229-1    Report Type:Direct      Company Report #CTU 209267  
 Age:55 YR    Gender:Male      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG BID	1 DAY	Clavicle Fracture		Bupropion 100mg Tab	PS		
Initial or Prolonged		Convulsion Fall					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/05/04ISR Number: 4263257-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0490813A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aorta Hypoplasia Atrial Septal Defect Drug Exposure During Pregnancy		Bupropion	PS	Glaxosmithkline	

Date:01/05/04ISR Number: 4263259-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0490904A

Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Avandia	PS	Glaxosmithkline	ORAL
4MG Twice per day		Hangover					
150MG Twice per day				Wellbutrin	SS	Glaxosmithkline	ORAL
				Starlix	C		
				Plavix	C		
				Synthroid	C	Glaxosmithkline	
				Aciphex	C		

Date:01/05/04ISR Number: 4263812-3Report Type:Direct Company Report #CTU 209312

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Paxil - Gsk 60 Mg	PS	Gsk	ORAL
60 MG ORAL @ NIGHT; A TOTAL OF 15 YRS ON SSRI 300 MG, SSRI		Anger Drug Withdrawal Syndrome Emotional Distress Fear					
				Wellbutrin	SS		



Feeling Abnormal  
Malaise

Date:01/05/04ISR Number: 4263832-9Report Type:Direct  
Age:25 YR Gender: I/FU:I

Company Report #CTU 209308

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required		Concussion Fall Grand Mal Convulsion		Wellbutrin Xr 150mg Began On 12/05/03, Increased To 300mg On 12/11	PS		
Intervention to 150 MG BEGAN Prevent Permanent 12/5/03 Impairment/Damage INCREASED TO  300 MG 12/11		Skull Fracture		Clomipramine	C		

Date:01/06/04ISR Number: 4264427-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0309783A  
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Unknown 3 WK		Diabetes Mellitus Headache Thirst Visual Acuity Reduced		Zyban	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/06/04ISR Number: 4264430-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0316252A  
 Age:38 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Alcohol Poisoning	Consumer	Bupropion	PS	Glaxosmithkline	ORAL
UNKNOWN			Coma		Ethanol	SS		
UNKNOWN			Completed Suicide		Fluoxetine	C		
UNKNOWN			Pulmonary Congestion		Flurazepam	C		
UNKNOWN			Pulmonary Oedema		Lorazepam	C		
UNKNOWN					Ibuprofen	C	Glaxosmithkline	
UNKNOWN					Albuterol	C	Glaxosmithkline	
RESPIRATORY								
(INHALATION)								

Date:01/06/04ISR Number: 4264431-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0316261A  
 Age:37 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL
			Intentional Misuse		Carbamazepine	C		
					Thioridazine	C		

Date:01/06/04ISR Number: 4264432-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0316263A  
 Age:18 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN			Intentional Misuse		Diphenhydramine	SS		

Date:01/06/04ISR Number: 4264433-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0316265A  
 Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN		Intentional Misuse		Benzodiazepines	C		
UNKNOWN				Alprazolam	C		
UNKNOWN				Maprotiline	C		

Date:01/06/04ISR Number: 4264435-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0317415A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Bupropion	PS	Glaxosmithkline	ORAL
UNKNOWN		Drug Toxicity		Alprazolam	SS		
UNKNOWN		Feeling Abnormal		Temazepam	SS		
UNKNOWN		Intentional Misuse					
UNKNOWN		Mood Swings					
UNKNOWN		Treatment Noncompliance					

Date:01/06/04ISR Number: 4264444-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0318606A  
Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Eczema		Zyban	PS	Glaxosmithkline	ORAL
1 MON				Salbutamol	C	Glaxosmithkline	
UNKNOWN				Fluticasone	C	Glaxosmithkline	
UNKNOWN				Betamethasone	C	Glaxosmithkline	
UNKNOWN				Oxytetracycline	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/06/04ISR Number: 4264449-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0318887A  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 12 DAY Initial or Prolonged		Diplopia Headache Iiird Nerve Paralysis Strabismus		Zyban	PS	Glaxosmithkline	ORAL

Date:01/06/04ISR Number: 4264450-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0318901A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 1 WK Congenital Anomaly		Congenital Anomaly Drug Exposure During Pregnancy Talipes		Zyban	PS	Glaxosmithkline	

Date:01/06/04ISR Number: 4264486-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0491332A  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Amnesia Bipolar Disorder Convulsion		Wellbutrin Valium Antidepressant (Unspecified) Narcotics (Unspecified)	PS C C C	Glaxosmithkline	ORAL

Date:01/06/04ISR Number: 4265056-8Report Type:Direct Company Report #CTU 209440  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 1 TABLET				Wellbutrin Sr 150 Mg Tab. Sa Glx	PS		
Initial or Prolonged TWICE A DAY							
Required (A WEEK OR Intervention to TWO BEFORE Prevent Permanent PERIOD) Impairment/Damage							

Date:01/07/04ISR Number: 4265664-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0490959A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	30 DAY	Condition Aggravated					
Other 300MG See dosage text	7 WK	Convulsion		Wellbutrin	SS	Glaxosmithkline	ORAL
		Medication Error					
				Lipitor	C		
				Synthroid	C	Glaxosmithkline	
				Unknown Medication	C		
				Vitamins	C		
				Aricept	C		

3 MON

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/07/04ISR Number: 4265673-5Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0314545A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Zyban	PS	Glaxosmithkline	
Other		Anxiety		Oxycodone	C		
15 DAY		Disturbance In Attention					
180MG per day		Dry Mouth					
		Dyskinesia					
		Dysstasia					
		Hyperhidrosis					
		Lack Of Spontaneous					
		Speech					
		Muscle Twitching					
		Nausea					
		Pallor					
		Tremor					

Date:01/09/04ISR Number: 4267508-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0414556A

Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
				Birth Control	C		ORAL

Date:01/09/04ISR Number: 4267509-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0415800A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Captopril	C	Glaxosmithkline	
per day							

Date:01/09/04ISR Number: 4267510-1Report Type:Periodic  
Age:64 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0415804A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Creatine Abnormal	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Three		Blood Urea					
times per day 1	YR	Hypokalaemia		Extra Strength			
500MG Twice		Renal Injury		Tylenol	SS	Glaxosmithkline	ORAL
per day	YR						
650MG Twice				Arthritis/Tylenol	SS	Glaxosmithkline	ORAL
per day	YR						

Date:01/09/04ISR Number: 4267511-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0416376A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
6 YR							

Date:01/09/04ISR Number: 4267512-5Report Type:Periodic  
Age:38 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0417155A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mania		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/09/04ISR Number: 4267513-7Report Type:Periodic  
 Age:54 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0417661A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dry Mouth		Wellbutrin	PS	Glaxosmithkline	ORAL
		Headache		Zestril	C		
		Memory Impairment		Elavil	C	Glaxosmithkline	
		Somnolence					

Date:01/09/04ISR Number: 4267514-9Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418009A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN		Drug Interaction		Demerol	SS		
UNKNOWN				Versed	SS		

Date:01/09/04ISR Number: 4267515-0Report Type:Periodic  
 Age:66 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0418060A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Three times per day		Insomnia		Bupropion	PS	Glaxosmithkline	ORAL
				Mirapex	C		
				Evista	C		

Date:01/09/04ISR Number: 4267516-2Report Type:Periodic  
 Age:10 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418077A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:01/09/04ISR Number: 4267517-4Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418604A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Behaviour		Wellbutrin	PS	Glaxosmithkline	ORAL
1	DAY						

Date:01/09/04ISR Number: 4267518-6Report Type:Periodic  
Age:27 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418606A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression Stress Swelling Face Tension Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267519-8Report Type:Periodic  
Age:31 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418608A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Smoker		Wellbutrin Xanax	PS C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/09/04ISR Number: 4267520-4Report Type:Periodic  
Age:82 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0418955A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	75MG Twice	Crying		Wellbutrin	PS	Glaxosmithkline	ORAL
	per day	Tinnitus		Tamoxifen	C		
				Fosamax	C		
				Lasix	C	Glaxosmithkline	
				Potassium	C		
				Zocor	C		
				Aricept	C		
				Klonopin	C		

Date:01/09/04ISR Number: 4267521-6Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0419167A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Unknown	Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
	YR	Sleep Disorder					

Date:01/09/04ISR Number: 4267522-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0419171A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hepatitis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267523-XReport Type:Periodic  
Age:79 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0419180A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	75MG Per day	Abdominal Distension		Wellbutrin	PS	Glaxosmithkline	ORAL

2	WK	Dizziness	Wellbutrin	SS	Glaxosmithkline	ORAL
		Nausea	Atenolol	C		
		Vision Blurred	Xanax	C		

Date:01/09/04ISR Number: 4267524-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0420194A  
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150TAB Four							
times per day 2 WK							

Date:01/09/04ISR Number: 4267525-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0420538A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Asthenia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Developmental		Zyprexa	C		
		Coordination Disorder		Diazide	C		
				Vasotec	C		
				Insulin	C		
				Digitex	C	Glaxosmithkline	
				Aggrenox	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/09/04ISR Number: 4267526-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420574A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness Vertigo		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267527-7Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0420764A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	3 MON	Tinnitus		Bupropion	PS	Glaxosmithkline	ORAL
UNKNOWN	81MG Per day			Aspirin	C	Glaxosmithkline	
				Concurrent Medications	C		

Date:01/09/04ISR Number: 4267528-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420766A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267529-0Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420767A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267530-7Report Type:Periodic  
Age:51 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0420811A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 75MG Per day	2 WK	Coma		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged Other		Confusional State Convulsion Dyspnoea Somnolence		Synthroid Detrol Methylprednisolone	C C C	Glaxosmithkline	

Date:01/09/04ISR Number: 4267532-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0421215A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	8 WK	Tinnitus Weight Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
				Birth Control Pills Zoloft Centrum Calcium	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/09/04ISR Number: 4267533-2Report Type:Periodic  
Age:43 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0421266A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:01/09/04ISR Number: 4267534-4Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422162A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nephrolithiasis		Wellbutrin	PS	Glaxosmithkline	ORAL
10MG Unknown	MON			Lexapro	SS		ORAL
				Buspar	C		

Date:01/09/04ISR Number: 4267535-6Report Type:Periodic  
Age:33 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0422163A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Screen False					
per day		Positive		Concerta	SS		ORAL
54MG Per day							

Date:01/09/04ISR Number: 4267536-8Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422192A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sexual Dysfunction		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	14 MON						

Norvasc	C	
Hydrochlorothiazide	C	
Multi-Vitamin	C	
Tylenol	C	Glaxosmithkline

Date:01/09/04ISR Number: 4267537-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0422194A  
 Age:46 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100TAB Per		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
day	27	DAY					
		Headache					
		Thinking Abnormal		Zoloft	C		
				Risperdal	C		

Date:01/09/04ISR Number: 4267538-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0422214A  
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2	MON					
		Drug Withdrawal Syndrome					
		Insomnia		Estradiol	C		
				Claritin D	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/09/04ISR Number: 4267539-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422295A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
				Aspirin	C	Glaxosmithkline	
				Coumadin	C	Glaxosmithkline	
				Proscar	C		
5MG Per day				Metoprolol Tartrate	C		
50MG Twice							
per day				Terazosin	C		
5MG Per day				Zocor	C		
40MG Per day				Xanax	C		

Date:01/09/04ISR Number: 4267540-XReport Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422316A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Herpes Zoster					
150MG Twice				Prozac	C		
per day	5	MON					

Date:01/09/04ISR Number: 4267541-1Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422329A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Panic Attack		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Xanax	C		
per day							



Date:01/09/04ISR Number: 4267542-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0422614A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	7	DAY					

Date:01/09/04ISR Number: 4267543-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0422645A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sluggishness		Wellbutrin	PS	Glaxosmithkline	ORAL
50MG As		Somnolence					
required							

Date:01/09/04ISR Number: 4267544-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0422734A  
Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral		Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Per day							

Date:01/09/04ISR Number: 4267545-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0422844A  
Age:11 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

UNKNOWN 20MG Per day Lexapro SS

Date:01/09/04ISR Number: 4267546-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0422922A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea Vomiting		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267547-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0423052A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort Chest Pain		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267548-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0423388A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice per day	1 MON						

Date:01/09/04ISR Number: 4267549-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0423434A  
Age:15 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267550-2Report Type:Periodic  
Age:51 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0423606A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Three						
times per day							
				B12	C	Glaxosmithkline	
				Ferrous Sulfate	C	Glaxosmithkline	
				Cipro	C	Glaxosmithkline	

Date:01/09/04ISR Number: 4267551-4Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0423700A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267552-6Report Type:Periodic  
Age:60 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0423852A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG	Unknown	Pharmaceutical Product		Wellbutrin	PS	Glaxosmithkline	ORAL
		Complaint					
		Retching					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/09/04ISR Number: 4267570-8Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424133A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Visual Acuity Reduced		Wellbutrin	PS	Glaxosmithkline	

Date:01/09/04ISR Number: 4267571-XReport Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424151A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267572-1Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424329A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Skin Discolouration		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	8	WK		Seroquel	C		

Date:01/09/04ISR Number: 4267573-3Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424379A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Crying		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2	WK					
		Insomnia					
		Nightmare					

Date:01/09/04ISR Number: 4267574-5Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0424554A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267575-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0424689A  
 Age: YR Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Unevaluable Event		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267576-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0424713A  
 Age:48 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice per day	18 MON	Gingivitis		Wellbutrin	PS	Glaxosmithkline	ORAL
		Nausea		Lamictal	SS	Glaxosmithkline	ORAL
25MG Per day				Effexor	C	Glaxosmithkline	
				Elavil	C		
				Thyroid	C		
				Vivactil	C		



Hyperhidrosis

Prozac  
Accupril

C  
C

Date:01/09/04ISR Number: 4267582-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425719A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error		Wellbutrin	PS	Glaxosmithkline	NASAL

Date:01/09/04ISR Number: 4267583-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425735A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 YR		Alopecia Weight Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267584-8Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425969A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	1 DAY	Dysuria		Wellbutrin	PS	Glaxosmithkline	ORAL

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Freedom Of Information (FOI) Report

Date:01/09/04ISR Number: 4267585-XReport Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426358A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Abnormal Behaviour		Phenobarbital	C		
				Propoxyphene	C		
				Vicodin	C		
				Potassium Bromide	C		
				Celexa	C		

Date:01/09/04ISR Number: 4267586-1Report Type:Periodic  
Age:59 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426582A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropion	PS	Glaxosmithkline	ORAL
Disability		Balance Disorder		Lexapro	C		
75MG Per day	2 WK	Visual Acuity Reduced		Maxzide	C	Glaxosmithkline	
				Prevacid	C		

Date:01/09/04ISR Number: 4267587-3Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426898A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
		Weight Decreased		Topamax	C		

Date:01/09/04ISR Number: 4267588-5Report Type:Periodic  
Age:25 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0426931A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Abdominal Pain Upper					
per day	8 DAY	Headache					
		Insomnia					



Nausea

Date:01/09/04ISR Number: 4267589-7Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426932A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Lexapro	PS C	Glaxosmithkline	

Date:01/09/04ISR Number: 4267590-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0426943A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Memory Impairment		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267591-5Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0427072A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amenorrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/09/04ISR Number: 4267592-7Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0427339A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Mania		Wellbutrin	PS	Glaxosmithkline	

Date:01/09/04ISR Number: 4267593-9Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427529A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Photosensitivity Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267594-0Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0427667A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Wellbutrin Alcohol	PS SS	Glaxosmithkline	

Date:01/09/04ISR Number: 4267595-2Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427689A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	

Date:01/09/04ISR Number: 4267596-4Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427745A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Malaise Stomach Discomfort		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267597-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0427855A  
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral	Health Professional	Wellbutrin	PS	Glaxosmithkline	ORAL
75MG Three							
times per day	2	MON		Norvasc	C		

Date:01/09/04ISR Number: 4267598-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0427867A  
Age:15 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	2	YR					

Date:01/09/04ISR Number: 4267599-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0428053A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	

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Freedom Of Information (FOI) Report

Date:01/09/04ISR Number: 4267600-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428128A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
60 DAY		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
		Neck Mass					
		Pharyngolaryngeal Pain					
		Subcutaneous Nodule					
		Thyroid Neoplasm					

Date:01/09/04ISR Number: 4267601-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428220A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
				Paxil	SS	Glaxosmithkline	
				Antidepressants	SS		
UNKNOWN							

Date:01/09/04ISR Number: 4267602-7Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428690A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
14 DAY		Dyspnoea Exertional		Wellbutrin	PS	Glaxosmithkline	ORAL
5 DAY				Lotrel	C		

Date:01/09/04ISR Number: 4267603-9Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428697A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 YR		Hypoglycaemia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Insulin	C		

Date:01/09/04ISR Number: 4267604-0Report Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428698A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hyperglycaemia		Wellbutrin	PS	Glaxosmithkline	ORAL
500MG Unknown	2 YR	Hypoglycaemia		Insulin	C		

Date:01/09/04ISR Number: 4267605-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428732A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin	PS	Glaxosmithkline	
1	DAY						

Date:01/09/04ISR Number: 4267606-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429136A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
UNKNOWN		Headache		Wellbutrin	PS	Glaxosmithkline	
		Local Swelling					
		Musculoskeletal Stiffness					
		Rash Pruritic					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/09/04ISR Number: 4267607-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429237A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
		Therapeutic Response Unexpected					

Date:01/09/04ISR Number: 4267608-8Report Type:Periodic  
 Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0429519A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion		Wellbutrin	PS	Glaxosmithkline	

Date:01/09/04ISR Number: 4267609-XReport Type:Periodic  
 Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430010A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267610-6Report Type:Periodic  
 Age:54 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430084A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
				Prozac	C		
				Zoloft	C		
				Tca	C		

Date:01/09/04ISR Number: 4267611-8Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430470A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Per day	9	DAY	Headache	Wellbutrin	PS	Glaxosmithkline	ORAL
			Palpitations	Tranxene	C		
			Somnolence	Glucovance	C		
				Benicar	C		
				Baby Aspirin	C	Glaxosmithkline	

Date:01/09/04ISR Number: 4267612-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430480A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Apathy		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:01/09/04ISR Number: 4267613-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430640A  
 Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Therapeutic Response Unexpected		Wellbutrin	PS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/09/04ISR Number: 4267614-3Report Type:Periodic  
Age:47 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0430648A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Weight Increased					
per day							

Date:01/09/04ISR Number: 4267619-2Report Type:Periodic  
Age:85 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430717A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Dilantin	C		

Date:01/09/04ISR Number: 4267620-9Report Type:Periodic  
Age:26 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430719A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Rash					
per day	2 WK	Urticaria					

Date:01/09/04ISR Number: 4267621-0Report Type:Periodic  
Age:46 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0430757A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Unknown	146 DAY						

Date:01/09/04ISR Number: 4267622-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430762A



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Fluctuation		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN							

Date:01/09/04ISR Number: 4267623-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430933A  
Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
37.5MG Twice		Pyrexia					
per day				Paxil	C	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267624-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430960A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Insulin	C		

Date:01/09/04ISR Number: 4267625-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431193A  
Age: Gender:Female I/FU:F

Outcome	PT
	Dizziness
	Drug Ineffective

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Nervousness

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice			Wellbutrin	PS	Glaxosmithkline	
per day	2 WK		Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day			Wellbutrin	SS	Glaxosmithkline	ORAL
			Klonopin	C		
			Lipitor	C		
			Insulin	C		

Date:01/09/04ISR Number: 4267626-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431438A  
 Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267627-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431608A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoaesthesia Muscle Spasms Myalgia Paraesthesia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267628-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431807A  
 Age:78 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Influenza Like Illness		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	10 YR	Nervousness		Prevacid	C		
		Pollakiuria		Synthroid	C	Glaxosmithkline	

Tinnitus  
Tremor  
Weight Decreased

Premarin C  
Cozar C

Date:01/09/04ISR Number: 4267629-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431815A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased		Wellbutrin	PS	Glaxosmithkline	

Date:01/09/04ISR Number: 4267631-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431945A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
2 WK				Lexapro	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/09/04ISR Number: 4267632-5Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431951A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1	MON	Depression		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267633-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431976A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG Twice		Fatigue	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	9 MON	Loss Of Consciousness					
Other		Nausea		Cold Medicine	SS		

Date:01/09/04ISR Number: 4267634-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431982A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Chest Pain Increased Appetite Nicotine Dependence		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267635-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432592A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1	WK	Rash		Wellbutrin	PS	Glaxosmithkline	
1	DAY	Urticaria		Zyban	SS	Glaxosmithkline	

Date:01/09/04ISR Number: 4267636-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432897A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flushing		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	5 YR						

Date:01/09/04ISR Number: 4267637-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432981A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Skin Discolouration		Wellbutrin	PS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267638-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0433001A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Heart Rate Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Unknown				Concerta	SS		ORAL
18MG Per day							

Date:01/09/04ISR Number: 4267639-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0433328A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/09/04ISR Number: 4267640-4Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0433346A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	MON	Medication Error					
				Citalopram	C		
				Methylphenidate	C		

Date:01/09/04ISR Number: 4267641-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0438902A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
75MG Three		Eye Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
times per day		Lacrimation Increased					

Date:01/09/04ISR Number: 4267642-8Report Type:Periodic  
Age:66 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439019A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Per day	1 YR	Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
				Ambien	C		
				Lexapro	C		

Date:01/09/04ISR Number: 4267643-XReport Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439050A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Alopecia		Wellbutrin	PS	Glaxosmithkline	
				Prednisone	C		

Date:01/09/04ISR Number: 4267644-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439417A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia Tinnitus Weight Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267645-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439461A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Feeling Jittery Heart Rate Increased		Wellbutrin	PS	Glaxosmithkline	

Date:01/09/04ISR Number: 4267646-5Report Type:Periodic  
Age:30 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439498A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Retinal Vein Occlusion		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/09/04ISR Number: 4267647-7Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439637A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Liver Function Test Abnormal		Wellbutrin	PS	Glaxosmithkline	

Date:01/09/04ISR Number: 4267649-0Report Type:Periodic  
 Age: YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0439815A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dysphonia Hypertrichosis		Wellbutrin	PS	Glaxosmithkline	

Date:01/09/04ISR Number: 4267650-7Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440212A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Screen Positive		Wellbutrin	PS	Glaxosmithkline	

Date:01/09/04ISR Number: 4267651-9Report Type:Periodic  
 Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440377A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
6 WK		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267652-0Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440387A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Chest Discomfort		Wellbutrin	PS	Glaxosmithkline	



Date:01/09/04ISR Number: 4267653-2Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440585A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Convulsion		Wellbutrin	PS	Glaxosmithkline	

Date:01/09/04ISR Number: 4267654-4Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440606A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Alopecia Menstruation Irregular		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267655-6Report Type:Periodic  
Age:82 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440607A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 WK	Amnesia Tinnitus	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
				Luvox	C		
				Risperdal	C		
				Lithium	C	Glaxosmithkline	
				Zocor	C		
40MG Per day							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

50MG Twice  
 per day  
 20MG Per day

Metoprolol C  
 Prilosec C

Date:01/09/04ISR Number: 4267656-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0440715A  
 Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Interaction		Paxil	SS	Glaxosmithkline	ORAL
		Fatigue					
		Weight Decreased					

Date:01/09/04ISR Number: 4267657-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0440975A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tachycardia		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN							

Date:01/09/04ISR Number: 4267658-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441133A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arrhythmia		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN							

Date:01/09/04ISR Number: 4267659-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441521A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Malaise	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267660-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441541A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mood Swings		Wellbutrin	PS	Glaxosmithkline	ORAL
				Lamictal	C	Glaxosmithkline	ORAL

Date:01/09/04ISR Number: 4267814-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0149544A  
Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abasia Amnesia Brain Damage Concussion Convulsion Difficulty In Walking Fall Head Injury Impaired Work Ability Loss Of Consciousness Lumbar Vertebral Fracture Memory Impairment		Wellbutrin Nicotrol Inhaler	PS SS	Glaxosmithkline Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/09/04ISR Number: 4267816-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0432608A  
 Age:81 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Confusional State	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
1 MON		Medication Error	Professional	Bupropion	SS	Glaxosmithkline	
		Nightmare		Heart Medication	C		
		Palpitations		Antihypertensives	C		
		Treatment Noncompliance		Lopressor	C		
		Urinary Retention					

Date:01/09/04ISR Number: 4267817-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0442800A  
 Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Medication Error	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
per day		No Adverse Effect	Professional				

Date:01/09/04ISR Number: 4267820-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0491694A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	
		Pharmaceutical Product					
		Complaint					

Date:01/12/04ISR Number: 4268588-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0431972A  
 Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							

per day 10 YR Arthritis Professional  
 Psoriasis Synthroid C Glaxosmithkline  
 Thiazide C  
 Fosamax C  
 Vasotec C

Date:01/12/04ISR Number: 4268590-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0491314A

Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK	Insomnia		Compazine	C	Glaxosmithkline	
		Medication Error					
		Visual Acuity Reduced					

Date:01/12/04ISR Number: 4268592-3Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0317735A

Age:27 YR Gender:Male I/FU:F

Outcome	PT
Other	Amnesia
	Dizziness
	Feeling Abnormal
	Hyperventilation
	Loss Of Consciousness
	Nausea
	Pallor

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Freedom Of Information (FOI) Report

Salivary Hypersecretion

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
UNKNOWN	10 DAY	Consumer	Zyban	PS	Glaxosmithkline	

Date:01/12/04ISR Number: 4268599-6Report Type:Expedited (15-DaCompany Report #DK-GLAXOSMITHKLINE-B0318928A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG per day		Depression Irritability Respiratory Distress		Zyban	PS	Glaxosmithkline	ORAL

Date:01/12/04ISR Number: 4268600-XReport Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0319066A  
 Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 9 WK		Depression Drug Withdrawal Syndrome		Zyban Acetylsalicylic Acid	PS C	Glaxosmithkline Glaxosmithkline	

Date:01/13/04ISR Number: 4269183-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0491932A  
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day 5 DAY		Blood Pressure Increased Dizziness Paraesthesia Syncope		Wellbutrin Unknown Supplement Estrace Calcium Progesterone Cream	PS SS C C C	Glaxosmithkline	ORAL

Date:01/13/04ISR Number: 4269184-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0492275A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
		Medication Error					

Date:01/13/04ISR Number: 4269192-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0316388A  
Age:21 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	
Other		Abnormal Behaviour					
UNKNOWN		Aggression		Sedatives			
		Confusional State		(Unspecified)	C		
UNKNOWN		Convulsion					
		Fall					
		Hallucination, Visual					
		Intentional Misuse					
		Nervous System Disorder					
		White Blood Cell Count					
		Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/13/04ISR Number: 4269206-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0318901A  
Age:6 MON Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	1 WK	Congenital Anomaly		Zyban	PS	Glaxosmithkline	
Congenital Anomaly		Drug Exposure During Pregnancy Talipes					

Date:01/13/04ISR Number: 4269217-3Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0319105A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Malaise Pain Pyrexia Skin Injury		Bupropion Hydrochloride	PS	Glaxosmithkline	

Date:01/13/04ISR Number: 4270567-5Report Type:Direct Company Report #CTU 209952  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	200 MG BID	Dysgeusia Pharmaceutical Product		Wellbutrin Sr 200 Mg Glaxosmithkline	PS	Glaxosmithkline	ORAL
ORAL		Complaint					

Date:01/13/04ISR Number: 4277275-5Report Type:Expedited (15-DaCompany Report #B0318456A  
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Agitation Choreoathetosis Convulsion	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion			



ORAL  
Electrocardiogram Qrs  
Hydrochloride)  
PS  
ORAL  
Complex Prolonged  
Electrocardiogram Qt  
Corrected Interval  
Prolonged  
Overdose  
Tachycardia  
Urinary Incontinence

Date:01/14/04ISR Number: 4270014-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0389829A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Psychotic Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270015-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0052984A  
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
		Pharmaceutical Product					
		Complaint		Wellbutrin	SS	Glaxosmithkline	ORAL
		Therapeutic Response					
		Decreased		Nortriptyline	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

5MG Per day Vasotec C  
 .1MG Per day Synthroid C Glaxosmithkline

Date:01/14/04ISR Number: 4270016-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0070258A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Female Orgasmic Disorder		Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day	7 WK	Headache		Estrogen	C		
		Libido Decreased					
		Sexual Dysfunction					

Date:01/14/04ISR Number: 4270017-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0379391A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression		Bupropion	PS	Glaxosmithkline	
UNKNOWN		Drug Ineffective					
		Lethargy					
		Weight Increased					

Date:01/14/04ISR Number: 4270018-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0385287A  
 Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Diaphragmalgia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Dystonia					
per day		Formication		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Twice		Hypoaesthesia					
per day	8 YR						

12.5MG Muscle Spasms Zoloft C  
 Alternate Tension  
 days Weight Increased  
 Tenormin C  
 Naprosyn C

Date:01/14/04ISR Number: 4270019-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0388023A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Laceration Pharmaceutical Product Complaint		Wellbutrin	PS	Glaxosmithkline	

Date:01/14/04ISR Number: 4270020-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0388176A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
1 MON		Nausea					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4270021-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0388627A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
2	WK						

Date:01/14/04ISR Number: 4270022-2Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0388679A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hair Colour Changes		Wellbutrin	PS	Glaxosmithkline	ORAL
75MG See							
dosage text	5	YR		Levoxyl	C	Glaxosmithkline	
				Hormones	C		
				Tegretol	C		

Date:01/14/04ISR Number: 4270023-4Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0388713A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
75MG In the							
morning				Temazepam	C		
				Fosamax	C		
				Estradiol	C		
				Prometrium	C		
				Lamictal	C	Glaxosmithkline	
				Celebrex	C		
				Calcium	C		
				Magnesium Oxide	C		
				Boron	C		
				Biotin	C		
				Vitamin E	C		
				Vitamin C	C	Glaxosmithkline	
				Gingko	C		

Multivitamin C  
Glucosamine C  
Chondroitin C  
Vitamin B50 C  
Co Q10 C  
L-Glutamine C

Date:01/14/04ISR Number: 4270024-6Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0389008A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Acne Agitation Amnesia Confusional State Coordination Abnormal Insomnia Nightmare Paranoia		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4270025-8Report Type:Periodic  
Age:11 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0389097A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	24 DAY	Joint Swelling Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270026-XReport Type:Periodic  
Age:19 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0389202A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day	1 MON	Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
				Prozac	SS		ORAL

Date:01/14/04ISR Number: 4270027-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0389499A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG per day	2 MON	Drug Ineffective Rash Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270028-3Report Type:Periodic  
Age:81 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0389788A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day		Blister		Wellbutrin	PS	Glaxosmithkline	ORAL
60MG per day				Paxil	C	Glaxosmithkline	
				Nexium	C		

YR

Date:01/14/04ISR Number: 4270031-3Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0389826A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Unknown	YR	Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270032-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390024A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270033-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390383A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4270034-9Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0390887A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Palpitations Panic Attack		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270035-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391255A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Ocular Hyperaemia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270036-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391316A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Genital Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270037-4Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391458A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN	2 YR	Deafness Hyperhidrosis Tinnitus		Wellbutrin	PS	Glaxosmithkline	

Date:01/14/04ISR Number: 4270038-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391512A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fibromyalgia		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:01/14/04ISR Number: 4270039-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0391522A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Energy Increased Nervous System Disorder		Wellbutrin	PS	Glaxosmithkline	

Date:01/14/04ISR Number: 4270040-4Report Type:Periodic  
Age:25 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0391658A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Disturbance In Attention Memory Impairment Personality Change	Health Professional	Wellbutrin Alcohol	PS SS	Glaxosmithkline	ORAL ORAL

Date:01/14/04ISR Number: 4270041-6Report Type:Periodic  
Age:45 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0391719A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Disturbance In Attention Memory Impairment					

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Freedom Of Information (FOI) Report

Personality Change

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Health Professional	Wellbutrin Alcohol	PS SS	Glaxosmithkline	ORAL ORAL

Date:01/14/04ISR Number: 4270042-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0391879A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
Other		Convulsion					
UNKNOWN							

Date:01/14/04ISR Number: 4270043-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0392136A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Dyspepsia Nausea					

Date:01/14/04ISR Number: 4270044-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0392155A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Unknown	20 DAY	Rash Generalised Urticaria					

Date:01/14/04ISR Number: 4270045-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0392331A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Hallucination					

.05U Per day      YR      Headache      Synthroid      C      Glaxosmithkline      ORAL  
Hot Flush

Date:01/14/04ISR Number: 4270046-5Report Type:Periodic      Company Report #US-GLAXOSMITHKLINE-A0392337A  
Age:      Gender:      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Bupropion	PS	Glaxosmithkline	
UNKNOWN							

Date:01/14/04ISR Number: 4270047-7Report Type:Periodic      Company Report #US-GLAXOSMITHKLINE-A0392415A  
Age:      Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Twice		Suicidal Ideation		Wellbutrin	PS	Glaxosmithkline	ORAL
per day							

Date:01/14/04ISR Number: 4270048-9Report Type:Periodic      Company Report #US-GLAXOSMITHKLINE-A0392515A  
Age:49 YR      Gender:Male      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4270049-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0392602A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dyspnoea Nonspecific Reaction Pharyngolaryngeal Pain		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270050-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393175A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation Anxiety Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270051-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393613A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Choking Sensation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270052-0Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393998A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Bupropion	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270053-2Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0393999A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270054-4Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394011A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
2 DAY		Formication		Zoloft	C		
		Panic Attack		Xanax	C		
				Nicotine Patch	C	Glaxosmithkline	

Date:01/14/04ISR Number: 4270055-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394058A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - 300MG per day				Topamax	C		
Initial or Prolonged				Depakote	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4270056-8Report Type:Periodic  
 Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394267A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	75MG Per day	Loss Of Libido		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270057-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394293A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Hypersensitivity Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270058-1Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394534A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphagia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270059-3Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394742A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270060-XReport Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0394873A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other UNKNOWN		Convulsion		Wellbutrin	PS	Glaxosmithkline	

Date:01/14/04ISR Number: 4270061-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395033A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:01/14/04ISR Number: 4270062-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395175A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270063-5Report Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395178A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
75MG Twice							
per day							

Tylenol #4	C	
Motrin	C	Glaxosmithkline
Viread	C	
Ziagen	C	Glaxosmithkline
Kaletra	C	

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Prevacid	C	
Valium	C	
Zoloft	C	
Atenolol	C	
Imodium	C	
Aspirin	C	Glaxosmithkline

Date:01/14/04ISR Number: 4270064-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395406A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270065-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395449A  
 Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoaesthesia		Bupropion	PS	Glaxosmithkline	ORAL
100MG In the							
morning	3 MON			Luvox	C		

Date:01/14/04ISR Number: 4270074-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395471A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270075-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0395672A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:01/14/04ISR Number: 4270076-3Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0395674A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
5	MON	Crying		Paxil	SS	Glaxosmithkline	ORAL
		Headache		Ibuprofen	C	Glaxosmithkline	
300MG	Unknown	Immobile					
		Mood Swings					
		Oedema Peripheral					
		Pain In Extremity					
		Restless Legs Syndrome					

Date:01/14/04ISR Number: 4270077-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395680A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Wellbutrin	PS	Glaxosmithkline	ORAL
		Therapeutic Response					
		Unexpected					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4270078-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0395719A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Constipation Menstrual Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270079-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0395864A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
YR		Drug Ineffective		Vicodin	C		
				Xanax	C		
				Temazepam	C		

Date:01/14/04ISR Number: 4270080-5Report Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396049A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	2 WK	Constipation Urticaria		Bupropion	PS	Glaxosmithkline	ORAL
				Methocarbamol	C		

Date:01/14/04ISR Number: 4270081-7Report Type:Periodic  
Age:75 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0396277A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	DAY	Dyspepsia Nausea Somnolence		Wellbutrin	PS	Glaxosmithkline	ORAL
				Tylenol #3 W Codeine Glucosamine	SS C		ORAL

Date:01/14/04ISR Number: 4270082-9Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396295A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urinary Retention		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	13	DAY					
2MG Per day	13	DAY		Gabitril	SS		ORAL
				Ambien	C		

Date:01/14/04ISR Number: 4270083-0Report Type:Periodic  
Age:45 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0396494A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Interaction					
per day		Therapeutic Response		Provigil	SS		ORAL
200MG Per day		Decreased					

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Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4270084-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0396662A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depressed Mood		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270085-4Report Type:Periodic  
Age:37 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0396774A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depressed Level Of Consciousness		Wellbutrin	PS	Glaxosmithkline	ORAL
1MG Per day	1 DAY	Drug Ineffective Performance Status Decreased		Lorazepam	SS		ORAL

Date:01/14/04ISR Number: 4270086-6Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0397207A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270087-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0397330A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Discomfort Depressed Mood Headache Pharmaceutical Product Complaint Somnolence		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270088-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0397336A  
Age:14 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270089-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0397337A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Anxiety Irritability		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270090-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0397351A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
10 MON		Weight Decreased		Zoloft	C		

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Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4270091-XReport Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0398324A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Asthma		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270092-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398335A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oligomenorrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270093-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398841A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270094-5Report Type:Periodic  
 Age:35 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0398930A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
10 DAY		Muscle Spasms		Propecia Strattera	C C		

Date:01/14/04ISR Number: 4270096-9Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0399076A

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Convulsion		Wellbutrin Theophylline	PS C	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270097-0Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0399087A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropion	PS	Glaxosmithkline	ORAL
Other		Convulsion		Elavil	C	Glaxosmithkline	

Date:01/14/04ISR Number: 4270099-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0399089A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue Somnolence		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270100-8Report Type:Periodic  
Age:35 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0399225A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bilirubin Urine		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Protein Urine Present					
per day	1	MON		Synthroid	C	Glaxosmithkline	
				Effexor	C		

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Date:01/14/04ISR Number: 4270101-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399449A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Liver Function Test Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270102-1Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0399583A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypotrichosis		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Three times per day 10 MON							

Date:01/14/04ISR Number: 4270103-3Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399589A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other 75MG Per day WK				Effexor	C		

Date:01/14/04ISR Number: 4270104-5Report Type:Periodic  
Age:53 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399601A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort		Wellbutrin	PS	Glaxosmithkline	ORAL
75MG Per day		Drug Ineffective		Remeron	SS		ORAL
45MG Per day 2 YR		Drug Withdrawal Syndrome Influenza Like Illness Malaise Nausea Palpitations					



Restlessness  
Tremor

Date:01/14/04ISR Number: 4270105-7Report Type:Periodic  
Age:72 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399855A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270106-9Report Type:Periodic  
Age:73 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399856A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
				Celexa	C		
				Effexor	C		
				Aspirin	C	Glaxosmithkline	
				Toprol Xl	C		
				Hytrin	C		

UNKNOWN

UNKNOWN

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Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4270107-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0400179A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Emotional Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270108-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0401146A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Screen False Positive		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270109-4Report Type:Periodic  
Age:16 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0401228A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day 1 MON	Drug Ineffective Irritability		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270110-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0401256A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event Drug Ineffective Therapeutic Response Unexpected		Wellbutrin Lexapro	PS C	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270111-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0401802A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction		Wellbutrin	PS	Glaxosmithkline	ORAL
		Tremor		Lexapro	SS		

UNKNOWN

Date:01/14/04ISR Number: 4270112-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0402025A  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Irritability					
per day	10 DAY	Oedema Peripheral		Naltrexone	C		
		Pruritus					
		Psychomotor Hyperactivity					
		Vision Blurred					

Date:01/14/04ISR Number: 4270113-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0402323A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:01/14/04ISR Number: 4270119-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0404218A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Skin Odour Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270120-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0404318A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
				Fat Burner	C		

Date:01/14/04ISR Number: 4270121-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0404373A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day							
UNKNOWN	300MG Twice			Paxil	C	Glaxosmithkline	
per day							
UNKNOWN	3VA per day			Klonopin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

UNKNOWN	10MG At night		Ambien		C	
Date:01/14/04ISR Number: 4270122-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0404377A						
Age:	Gender:Male	I/FU:F				
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Premature Ejaculation		Wellbutrin Prozac	PS C	Glaxosmithkline
						ORAL
Date:01/14/04ISR Number: 4270123-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0404775A						
Age:	Gender:Female	I/FU:I				
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Agitation Depressed Mood Dizziness Malaise Self-Injurious Ideation		Wellbutrin	PS	Glaxosmithkline
						ORAL
Date:01/14/04ISR Number: 4270124-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0404781A						
Age:	Gender:Female	I/FU:I				
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Weight Decreased		Wellbutrin	PS	Glaxosmithkline
						ORAL
Date:01/14/04ISR Number: 4270125-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0404991A						
Age:42 YR	Gender:Female	I/FU:I				
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Bruxism Tremor		Wellbutrin	PS	Glaxosmithkline
						ORAL

Date:01/14/04ISR Number: 4270126-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0405330A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270127-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0405387A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Deafness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270128-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0405397A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dental Caries		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4270129-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0405426A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dyspepsia Gastrooesophageal Reflux Disease		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270130-6Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0405756A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1TAB Twice per day		Aggression Anger Contusion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270136-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0405817A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 WK		Abnormal Dreams Psychomotor Hyperactivity Restlessness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270137-9Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0405902A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	3 WK	Rash Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:01/14/04ISR Number: 4270138-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406446A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	
3 YR				Geodon	C		

Date:01/14/04ISR Number: 4270139-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0406923A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
4 YR							

Date:01/14/04ISR Number: 4270140-9Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0407133A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Feeling Jittery Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4270141-0Report Type:Periodic  
 Age:13 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0407396A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
75MG Per day	1 WK	Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Hypersensitivity					
		Serum Sickness					
		Urticaria					

Date:01/14/04ISR Number: 4270142-2Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0408075A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Bupropion Alcohol	PS SS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270143-4Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408212A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Priapism		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270144-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408689A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other	100MG Twice per day	Convulsion		Bupropion	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270145-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408890A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG At night	6 WK	Crying Depression Dizziness Drug Ineffective Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270146-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0409525A  
 Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270147-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0409539A  
 Age:50 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	MON	Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4270148-3Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0409637A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75MG Twice per day	1 YR	Confusional State Disorientation Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270149-5Report Type:Periodic  
Age:15 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0409942A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin Dexedrine	PS C	Glaxosmithkline Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270150-1Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0410330A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG per day		Anger Irritability Nervousness		Wellbutrin Atarax	PS C	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270152-5Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0411801A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
25MG Per day	3 DAY	Somnolence		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN	20MG Per day	10 MON		Lexapro	C		

Date:01/14/04ISR Number: 4270153-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0411924A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Bupropion	PS	Glaxosmithkline	ORAL
100MG Twice							
per day	19	DAY					

Date:01/14/04ISR Number: 4270162-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0412060A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mania		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270163-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0412415A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyskinesia		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day	1	YR		Prozac	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4270164-1Report Type:Periodic  
Age:59 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0412977A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	75MG Per day	Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270165-3Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0413259A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	7 DAY	Hallucination, Visual		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270166-5Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0413261A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination, Visual		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270167-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0413293A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	200MG Unknown	Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
	WK			Zocor	C		

Date:01/14/04ISR Number: 4270168-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413314A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270169-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413327A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270170-7Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0413934A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Twice		Polymenorrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	16 DAY	Rash					

Date:01/14/04ISR Number: 4270171-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0414002A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Screen Positive		Bupropion	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4270172-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0414062A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Wellbutrin Effexor	PS C	Glaxosmithkline	ORAL ORAL

Date:01/14/04ISR Number: 4270173-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0414188A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Liver Function Test Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270174-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0414234A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
2 MON		Muscle Rigidity Muscle Tightness					

Date:01/14/04ISR Number: 4270175-6Report Type:Periodic  
Age:21 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0414335A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
3 WK		Rash		Avandia	SS	Glaxosmithkline	
UNKNOWN		Rash Papular		Lexapro Trazodone Orthocyclen	C C C		



Date:01/14/04ISR Number: 4270528-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0317875A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dizziness Nausea Swelling Face Syncope Urticaria		Zyban	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270530-4Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0319062A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG per day	8 DAY	Psoriasis		Zyban	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270533-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0319125A  
Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 7 DAY		Irritability		Zyban	PS	Glaxosmithkline	ORAL
		Personality Change		Seretide	C	Glaxosmithkline	
				Salbutamol	C	Glaxosmithkline	
				Prednisolone	C	Glaxosmithkline	ORAL

40MG per day 7 DAY

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4270534-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0319131A  
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 21 DAY		Malaise  Psoriasis Rash Pruritic Skin Exfoliation		Zyban	PS	Glaxosmithkline	ORAL

Date:01/14/04ISR Number: 4270535-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0319266A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Abuser		Zyban Ritalin	PS SS	Glaxosmithkline	

Date:01/15/04ISR Number: 4271781-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0492250A  
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150TAB Twice Initial or Prolonged per day	5 DAY	Amnesia  Convulsion  Hypoaesthesia Oral Salivary Hypersecretion Tongue Biting	Consumer	Wellbutrin   Nicotine Transdermal Patch Protonix	PS   C C	Glaxosmithkline   Glaxosmithkline	ORAL

Date:01/15/04ISR Number: 4271802-XReport Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0319104A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Disability UNKNOWN	4 DAY	Cerebral Infarction		Bupropion Hydrochloride  Ethinylestradiol +	PS	Glaxosmithkline	

Date:01/15/04ISR Number: 4271803-1Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0319175A  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Vaginal Haemorrhage		Bupropion Hydrochloride No Concurrent Medication	PS  C	Glaxosmithkline	

Date:01/16/04ISR Number: 4273114-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0490813A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other	300MG per day	Abortion Induced	Health	Bupropion	PS	Glaxosmithkline	
		Aorta Hypoplasia Atrial Septal Defect Complications Of Maternal Exposure To Therapeutic Drugs Drug Exposure During Pregnancy	Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/16/04ISR Number: 4273118-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0492588A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Accident		Wellbutrin	PS	Glaxosmithkline	

Date:01/16/04ISR Number: 4273119-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0492602A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bite		Wellbutrin	PS	Glaxosmithkline	ORAL
100TAB Twice		Confusional State					
per day	10 WK	Cyanosis		Prozac	C		
80MG per day		Grand Mal Convulsion					
		Respiratory Arrest					

Date:01/16/04ISR Number: 4273120-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0492882A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	
		Pharmaceutical Product					
		Complaint					

Date:01/16/04ISR Number: 4273121-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0492888A  
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Three		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
times per day	2 WK	Medication Error					
		Nausea		Glucophage	C		
		Overdose		Spirolactone	C		

Zelnorm C  
Axid C  
Alcon C

Date:01/16/04ISR Number: 4273122-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0492909A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:01/16/04ISR Number: 4273123-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0493122A  
Age:25 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Unknown		Concussion		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged 150MG At Other night		Fall		Clomipramine	C		
		Grand Mal Convulsion					
		Medication Error					
		Skull Fracture					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/16/04ISR Number: 4273134-2Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0317539A

Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Agraphia		Bupropion			
		Aphagia		Hydrochloride	PS	Glaxosmithkline	ORAL
		Aphasia		Warfarin	C	Glaxosmithkline	ORAL
5MG Per day							
		Transient Ischaemic		Perindopril Erbumine	C		ORAL
6MG per day							
		Attack		Amiloride /			
				Hydrochlorothiazide	C		ORAL
1U per day							
				Rofecoxib	C		ORAL
25MG per day							
				Conjugated Oestrogen	C		
				Arthroguard	C		
				Digoxin	C	Glaxosmithkline	ORAL
.25MG per day							

Date:01/19/04ISR Number: 4273494-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0314463A

Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Blood Pressure Decreased	Health	Zyban	PS	Glaxosmithkline	ORAL
13 DAY							
Initial or Prolonged		Circulatory Collapse	Professional	No Concurrent			
		Dizziness		Medications	C		
		Pharyngolaryngeal Pain					
		Pruritus					
		Rash Erythematous					
		Rash Generalised					
		Type I Hypersensitivity					

Date:01/20/04ISR Number: 4274086-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0493247A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	30 DAY						

Facial Bones Fracture

Date:01/20/04ISR Number: 4274095-2Report Type:Expedited (15-DaCompany Report #TR-GLAXOSMITHKLINE-B0314678A  
 Age:30 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 27 DAY Initial or Prolonged	Genital Lesion Headache Pruritus		Zyban	PS	Glaxosmithkline	ORAL

Date:01/20/04ISR Number: 4274102-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0318367A  
 Age:55 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150MG Per day 3 DAY Hospitalization - Initial or Prolonged Disability	Coronary Artery Stenosis Loss Of Consciousness Myocardial Infarction	Consumer	Zyban	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/20/04ISR Number: 4274104-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0319137A  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Condition Aggravated Overdose Suicidal Ideation		Zyban	PS	Glaxosmithkline	ORAL

Date:01/20/04ISR Number: 4274105-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0319365A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG See Hospitalization - dosage text 17 DAY Initial or Prolonged		Angioneurotic Oedema Paraesthesia Oral Pruritus Generalised		Zyban	PS	Glaxosmithkline	ORAL

Date:01/20/04ISR Number: 4274109-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0319493A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG See dosage text 8 WK		Anorexia Balance Disorder Depression Lymphadenopathy Weight Decreased		Zyban	PS	Glaxosmithkline	ORAL

Date:01/20/04ISR Number: 4274113-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0319635A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG See		Psoriasis		Zyban	PS	Glaxosmithkline	ORAL



dosage text

Date:01/21/04ISR Number: 4275504-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441481A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Screen Positive	Health	Wellbutrin	PS	Glaxosmithkline	
UNKNOWN	9000MG	Single					
		Suicide Attempt	Professional				

dose

Date:01/21/04ISR Number: 4275523-9Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0319066A

Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Depression		Zyban	PS	Glaxosmithkline	
9 WK		Drug Withdrawal Syndrome		Acetylsalicylic Acid	C	Glaxosmithkline	ORAL
		Headache					
		Nausea					

Date:01/21/04ISR Number: 4275680-4Report Type:Direct Company Report #USP 56267

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin Xl	PS	Glaxo Smith Kline	
TABLET				Wellbutrin Sr	SS	Glaxo Smith Kline	
TABLET							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/21/04ISR Number: 4276114-6Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427536A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Valtrex	PS	Glaxosmithkline	ORAL
500MG Per day	4 WK	Depression		Wellbutrin	SS	Glaxosmithkline	
UNKNOWN		Drug Ineffective Obsessive-Compulsive Disorder		Serzone	C		

Date:01/21/04ISR Number: 4277332-3Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 210552

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 500MG ONCE		Chorea		Levaquin 500mg	PS		ORAL
Initial or Prolonged DAILY ORAL		Dyskinesia		Welbutrin Sr 150mg	SS		ORAL
150MG TWICE				..	C		
DAILY ORAL				Nyquil	C		
				Diovan Hct	C		

Date:01/21/04ISR Number: 4277616-9Report Type:Expedited (15-DaCompany Report #2004001589  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other (DAILY), ORAL		Anorexia	Health	Zoloft (Sertraline)	PS		ORAL
ORAL		Blood Potassium Decreased Convulsion Diarrhoea Drug Hypersensitivity	Professional	Terbinafine Hydrochloride (Terbinafine Hydrochloride)	SS		ORAL
		Feeling Abnormal		Bupropion			

	Tremor		Hydrochloride (Bupropion Hydrochloride)	SS		ORAL
ORAL			Paroxetine Hydrochloride (Paroxetine Hydrochloride)	SS		ORAL
ORAL						

Date:01/22/04ISR Number: 4277402-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0438799A  
Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
3 WK		Fall	Professional	Mircette	C		
		Grand Mal Convulsion					
		Skin Laceration					

Date:01/22/04ISR Number: 4277407-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0493591A  
Age:61 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Aphasia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK	Convulsion		Trazodone	C		
Initial or Prolonged		Paraesthesia		Plavix	C		
		Paralysis					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/22/04ISR Number: 4277409-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0493935A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:01/22/04ISR Number: 4277412-2Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0494028A

Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diplopia		Zyban	PS	Glaxosmithkline	ORAL
Other		Vision Blurred					
150MG See		Visual Acuity Reduced		Birth Control	C		ORAL
dosage text				Herbal Multivitamin	C		

Date:01/22/04ISR Number: 4277418-3Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0317324A

Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Health	Zyban	PS	Glaxosmithkline	ORAL
Other		Grand Mal Convulsion	Professional	No Concurrent			
11 DAY		Loss Of Consciousness		Medication	C		
		Tongue Biting					
		Urinary Incontinence					

Date:01/22/04ISR Number: 4277420-1Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0319435A

Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization -		Fall					
150MG per day 1 DAY		Grand Mal Convulsion					
Initial or Prolonged		Musculoskeletal Stiffness					

Paralysis

Date:01/22/04ISR Number: 4277424-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0320051A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	5 DAY						

Date:01/22/04ISR Number: 4278579-2Report Type:Expedited (15-DaCompany Report #US012622  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Provigil Wellbutrin	PS SS		

Date:01/22/04ISR Number: 4278906-6Report Type:Direct Company Report #CTU 210620  
Age:45 YR Gender:Male I/FU:I

Outcome	PT
Life-Threatening	Anxiety Depression Pharmaceutical Product Complaint

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Suicidal Ideation

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
ONE QAM ORAL			Wellbutrin-Xl 300 Mg Glaxosmithkline	PS	Glaxosmithkline	ORAL

Date:01/22/04ISR Number: 4278918-2Report Type:Expedited (15-DaCompany Report #B0318456A  
Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL		Complex Prolonged Overdose Tachycardia Urinary Incontinence					

Date:01/23/04ISR Number: 4277908-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0494266A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 3 YR			Angioneurotic Oedema	Wellbutrin	PS	Glaxosmithkline	
300MG Per day			Condition Aggravated	Wellbutrin	SS	Glaxosmithkline	ORAL
		Gastrooesophageal Reflux Disease Retinal Detachment		Buspar Seizure Medication	C C		

Date:01/23/04ISR Number: 4277909-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0494501A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Medication Error

Wellbutrin

PS

Glaxosmithkline

Date:01/23/04ISR Number: 4277920-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0319950A

Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	23G Unknown	Convulsion	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged	UNKNOWN	Overdose	Professional	Ethanol	SS		
		Paralysis					

Date:01/23/04ISR Number: 4277922-8Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0320073A

Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	22.5G per day	Grand Mal Convulsion		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization -	Initial or Prolonged	Obstructive Airways Disorder					
		Overdose					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/27/04ISR Number: 4279455-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0412091A  
 Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	10 MON	Chorioamnionitis Dizziness Drug Exposure During Pregnancy Fatigue Stillbirth	Health Professional	Wellbutrin Prenatal Vitamins Folic Acid Alcohol Juice Plus	PS C C C C	Glaxosmithkline	ORAL  ORAL ORAL

Date:01/27/04ISR Number: 4279456-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0412091B  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Twice Congenital Anomaly per day		Abdominal Wall Anomaly Arteriopathic Disease Atrial Septal Defect Chorioamnionitis Cleft Palate Drug Exposure During Pregnancy Ear Malformation Intra-Uterine Death Jaw Fracture Kyphosis Liver Disorder Lymphangiectasia Macrogathia Nose Deformity Pectus Excavatum Placental Disorder Spleen Malformation Thyroid Disorder Traumatic Delivery	Health Professional	Wellbutrin Alcohol Folic Acid Juice Plus Prenatal Vitamins	PS C C C C	Glaxosmithkline	



Date:01/27/04ISR Number: 4279467-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0494894A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Electrocardiogram Qt Prolonged		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/27/04ISR Number: 4279476-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0318901A

Age:6 MON Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Congenital Anomaly	Health	Zyban	PS	Glaxosmithkline	
150MG Twice							
Congenital Anomaly		Drug Exposure During	Professional				
per day	1 WK	Pregnancy					
		Talipes					

Date:01/27/04ISR Number: 4279483-6Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0320169A

Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Haematemesis		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	27 DAY						
Initial or Prolonged							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/27/04ISR Number: 4279490-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0320817A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Zyban	PS	Glaxosmithkline	ORAL
Other		Drug Interaction	Professional				
150MG See dosage text	1	MON					
		Hypoaesthesia					
100MG per day	5	DAY		Corticoid	SS		ORAL
		Muscular Weakness					
		Paraesthesia					

Date:01/27/04ISR Number: 4280802-5Report Type:Expedited (15-DaCompany Report #163-20785-03120498(1)  
 Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Thalomid			
Required Intervention to Prevent Permanent Impairment/Damage		Abdominal Distension Condition Aggravated Follicular Thyroid Cancer	Health Professional	(Thalidomide) (100 Milligram, Capsules)	PS		ORAL
400-800 MG, QD, ORAL;		Oedema Peripheral					
500 MG, DAILY, ORAL;		Oropharyngeal Swelling					
400 MG,		Rash Pruritic					
		Thyroglobulin Increased					
		White Blood Cell Count Decreased		Wellbutrin (Bupropion Hydrochloride)	SS		
				Cozaar (Losartan Potassium)	SS		
				Alprazolam (Alprazolam)	C		
				Nifedipine (Nifedipine)	C		
				Synthroid (Levothyroxine Sodium)	C		
				Calcitrol (Calcitrol)	C		
				Pepcid (Famotidine)	C		

Slo-Mag (Magnesium  
Chloride Anhydrous) C  
K-Dur (Potassium  
Chloride) C  
Fluvoxamine  
(Fluvoxamine) C  
Avalide C

Date:01/28/04ISR Number: 4280190-4Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0320004A  
Age:33 YR Gender:Female I/FU:F

Outcome PT  
Other Anxiety  
Aptyalism  
Asthenia  
Cytolytic Hepatitis  
Dermatitis Exfoliative  
Hypersensitivity  
Hypotension  
Inflammation  
Insomnia  
Leukopenia  
Lymphadenopathy  
Nausea  
Painful Respiration

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG	Unknown	22 DAY		Zyban	PS	Glaxosmithkline	ORAL

Date:01/28/04ISR Number: 4280202-8Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0320703A  
 Age:47 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG per day		Convulsion	Foreign	Bupropion	PS	Glaxosmithkline	
Initial or Prolonged	400MG per day		Muscle Twitching		Tramadol	C		
			Salivary Hypersecretion		Salbutamol / Ipratropium Salmeterol Xinafoate + Fluticasone Propionate	C	Glaxosmithkline	
RESPIRATORY								
(INHALATION)								
	30MG per day				Codeine	C		ORAL
	20MG per day				Morphine Sulphate	C	Glaxosmithkline	ORAL

Date:01/28/04ISR Number: 4280209-0Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0042891A  
 Age:34 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day	4 DAY	Chest Pain		Zyban	PS	Glaxosmithkline	ORAL
Other	UNKNOWN		Circulatory Collapse		Fluoxetine	C		
			3 MON					
			Dyspnoea					
			Tachycardia					

Date:01/28/04ISR Number: 4281982-8Report Type:Expedited (15-DaCompany Report #SUS1-2003-00554

Age:51 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional	Carbatrol (Carbamazepine) Capsule Xr Acetaminophen/Propox yphene() Bupropion (Bupropion)	PS  SS  SS		

Date:01/29/04ISR Number: 4280888-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0431158A

Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 300MG Per day 1 WK Hospitalization - Initial or Prolonged		Amnesia  Grand Mal Convulsion Joint Dislocation Tongue Biting	Health Professional	Wellbutrin  None	PS  C	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/29/04ISR Number: 4280889-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439046A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	300MG Per day	2 DAY	Blindness	Health	PS	Glaxosmithkline	ORAL
Other			Optic Ischaemic Neuropathy	Professional	C		
			Visual Field Defect		C		
				Norvasc	C		
				Xanax	C		
				Zocor	C		
				Remeron	C		

Date:01/29/04ISR Number: 4280892-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0494241A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day		Medication Error	Wellbutrin	PS	Glaxosmithkline	ORAL
			Pharmaceutical Product Complaint				

Date:01/29/04ISR Number: 4280914-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0319950A

Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	23G Unknown		Alcohol Interaction	Consumer	PS	Glaxosmithkline	ORAL
Initial or Prolonged	UNKNOWN		Overdose	Ethanol	SS		
			Status Epilepticus				

Date:01/30/04ISR Number: 4282065-3Report Type:Periodic Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-12130480

Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Alopecia	Trazodone Hcl	PS	Apothecon	
				Wellbutrin	SS		ORAL

Date:01/30/04ISR Number: 4282078-1Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-12199782

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Supraventricular		Trazodone Hcl Tabs	PS	Apothecon	ORAL
Initial or Prolonged	Tachycardia		Wellbutrin Sr	SS		ORAL
VARIABLE						

DOSE-200-300m

g/day

Date:01/30/04ISR Number: 4282181-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0139855A  
Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Brain Damage		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
Initial or Prolonged	Cardiac Arrest		Buspar	C		
Disability	Drowning		Sedative(Unknown)	C		
Other	Dysarthria					
	Grand Mal Convulsion					
	Memory Impairment					
	Motor Dysfunction					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/30/04ISR Number: 4282202-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0495331A  
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Muscle Contractions					
per day		Involuntary		Paxil	C	Glaxosmithkline	
		Partial Seizures		Ativan	C		
		Tremor		Zyrtec	C	Glaxosmithkline	

Date:01/30/04ISR Number: 4282220-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0320766A  
 Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris		Bupropion			
13 DAY				Hydrochloride	PS	Glaxosmithkline	ORAL
UNKNOWN				Ramipril	C		
UNKNOWN				Aspirin	C	Glaxosmithkline	
UNKNOWN				Diltiazem			
UNKNOWN				Hydrochloride	C	Glaxosmithkline	
UNKNOWN				Atorvastatin	C		
UNKNOWN				Elantan La	C		
UNKNOWN	50MG Per day						

Date:01/30/04ISR Number: 4282864-8Report Type:Periodic Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-12333696  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Liver Function Test		Serzone Tabs	PS	Bristol-Myers Squibb	
		Abnormal				Company	ORAL
prescribed				Effexor	SS		



for "one  
week"  
prescribed  
for the  
"second week"

Wellbutrin SS

Date:01/30/04ISR Number: 4283380-XReport Type:Direct  
Age: Gender: I/FU:I

Company Report #USP 56276

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error		Wellbutrin Xl	PS	Glaxosmithkline	
				Wellbutrin Sr	SS	Glaxosmithkline	

TABLET

Date:01/30/04ISR Number: 4284481-2Report Type:Direct  
Age:46 YR Gender:Female I/FU:I

Company Report #CTU 211209

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Asthenia		Wellbutrin Sr 200 Mg	PS		ORAL
		Chills		Omeprazole	C		
		Dizziness		Hctz	C		
		Muscle Spasms					
		Tremor					

200 MG PO BID

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/02/04ISR Number: 4283565-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441482A  
 Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 100MG Twice per day	5 WK	Arthritis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/02/04ISR Number: 4283567-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0495325A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:02/02/04ISR Number: 4283572-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0495942A  
 Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/02/04ISR Number: 4283586-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0320504A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2TABS Per day	7 DAY	Blister		Zyban	PS	Glaxosmithkline	ORAL
Burning Sensation Eye Swelling Local Swelling Oedema Peripheral Psoriasis Rash Rash Macular Swelling Face Thrombosis Tremor							

Date:02/02/04ISR Number: 4283594-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0320609A

Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Astrocytoma		Zyban	PS	Glaxosmithkline	ORAL
28 DAY				Fluoxetine Hydrochloride	C		
UNKNOWN							

Date:02/02/04ISR Number: 4283595-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0320767A

Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Angioneurotic Oedema Drug Interaction	Consumer	Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
18 DAY				Co-Codamol	SS		
UNKNOWN							
UNKNOWN							
UNKNOWN							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/02/04ISR Number: 4283596-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0320770A  
 Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice		Psoriasis		Zyban	PS	Glaxosmithkline	ORAL
per day	1	MON					
UNKNOWN				Dovonex	C		
UNKNOWN				Bendrofluazide	C	Glaxosmithkline	
UNKNOWN				Canesten	C	Glaxosmithkline	
UNKNOWN				Beclomethasone Dipropionate	C	Glaxosmithkline	
UNKNOWN	1PUFF Twice						
per day							
UNKNOWN				Aqueous Cream Bp	C		

Date:02/02/04ISR Number: 4283599-8Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0043010A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Five		Alcohol Use	Health	Zyban	PS	Glaxosmithkline	ORAL
times per day		Suicide Attempt	Professional				
				Alcohol	SS		ORAL

Date:02/02/04ISR Number: 4285172-4Report Type:Expedited (15-DaCompany Report #2004196093US  
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged BUCCAL	BUCCAL	Abasia Amnesia	Consumer	Nicotrol (Nicotine) Inhaler	PS		
		Brain Damage		Wellbutrin			

ORAL  
 Convulsion (Amfebutamone Hydrochloride) SS ORAL  
 Fall  
 Head Injury  
 Loss Of Consciousness  
 Lumbar Vertebral Fracture

Date:02/03/04ISR Number: 4284134-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0495505A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	
				Wellbutrin	SS	Glaxosmithkline	

Date:02/03/04ISR Number: 4284135-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0495535A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 MON	Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	

Date:02/03/04ISR Number: 4284157-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0320606A

Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Sudden Death		Zyban	PS	Glaxosmithkline	ORAL
28 DAY				Aspirin	C	Glaxosmithkline	
Hospitalization - UNKNOWN				Glyceryl Trinitrate	C	Glaxosmithkline	
Initial or Prolonged UNKNOWN							

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Hospitalization - Renal Failure Acute Wellbutrin PS Glaxosmithkline ORAL  
 150MG Twice  
 Initial or Prolonged  
 per day

Lexapro C  
 Avapro C  
 Coumadin C Glaxosmithkline  
 Amiodarone C  
 Lipitor C  
 Saw Palmetto C  
 Multivitamin C  
 Tricor C  
 Viagra C

Date:02/04/04ISR Number: 4285106-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0495985A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Drug Exposure During Pregnancy Dyspepsia Headache Loss Of Consciousness Neck Pain					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/04/04ISR Number: 4285108-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0496012A  
Age:1 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other			Accidental Overdose Agitation Grand Mal Convulsion Metabolic Acidosis	Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:02/04/04ISR Number: 4285113-XReport Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0319175A  
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Vaginal Haemorrhage	Bupropion Hydrochloride Hormone Replacement Therapy	PS C	Glaxosmithkline	

YR

Date:02/04/04ISR Number: 4285127-XReport Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0042959A  
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day			Abdominal Pain Upper Anaemia Fat Intolerance Gamma-Glutamyltransferase Increased Lipase Increased	Zyban	PS	Glaxosmithkline	ORAL

Date:02/05/04ISR Number: 4285907-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439007A  
Age:78 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Hospitalization -	Balance Disorder	Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Twice					
Initial or Prolonged	Chills				
per day	1 MON				
Other	Depression	Flomax	C		
	Erectile Dysfunction	Cozaar	C		
	Insomnia	Indapamide	C		
	Myalgia	Duragesic	C		
	Myoclonus				
	Tremor				
	Vestibular Disorder				

Date:02/05/04ISR Number: 4285915-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0307572A  
Age:70 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dysarthria	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK	Dysgraphia	Professional	Chondrosulf	C		
		Metastases To Central Nervous System					
		Speech Disorder					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/05/04ISR Number: 4285916-1Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0318010A  
 Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1TAB Twice		Aggression		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	5 DAY	Agitation					
Other		Blood Pressure Increased Depression Dizziness Feeling Abnormal Headache					

Date:02/05/04ISR Number: 4285926-4Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0321039A  
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Per day	11 DAY	Chromatopsia		Zyntabac	PS	Glaxosmithkline	ORAL
		Constipation Dysgeusia Neuralgia Thirst Vertigo Visual Disturbance					

Date:02/05/04ISR Number: 4285930-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0321213A  
 Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See		Anxiety		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged dosage text	2 DAY	Dry Mouth Headache Intentional Misuse Overdose Palpitations Vertigo					

Date:02/05/04ISR Number: 4286479-7Report Type:Direct  
Age:49 YR Gender:Male I/FU:I

Company Report #CTU 211616

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Vicodin 10 Mg	PS		
Other		Pharmaceutical Product Complaint		Wellbutrin	SS		
				Ativan 2mg	SS		

Date:02/06/04ISR Number: 4286730-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0495765A  
Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Death		Death		Klonopin	C		
5 DAY				Unknown Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/06/04ISR Number: 4286731-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0495776A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	1 DAY			Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Medication Error					

Date:02/06/04ISR Number: 4286740-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0319984A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	
Other		Abdominal Pain Upper Anorexia Balance Disorder Constipation Deafness Disturbance In Attention Dysgeusia Ear Discomfort Ear Disorder Feeling Abnormal Nausea Sleep Disorder Tinnitus					

Date:02/06/04ISR Number: 4286745-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0321121A  
 Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	13 DAY			Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
Other		Angina Pectoris					
1TAB Twice per day				Ramipril	C		
UNKNOWN				Aspirin	C	Glaxosmithkline	
UNKNOWN				Diltiazem			

UNKNOWN Hydrochloride C Glaxosmithkline  
 UNKNOWN Atorvastatin C  
 UNKNOWN Elantan La C

Date:02/06/04ISR Number: 4286746-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0321125A  
 Age:25 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination, Visual		Zyban	PS	Glaxosmithkline	ORAL
150MG See							
dosage text	5	Muscle Spasms					
	DAY			Nicotinell	C	Glaxosmithkline	
TRANSDERMAL							

Date:02/06/04ISR Number: 4286747-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0321126A  
 Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Neutropenia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	69						
	DAY			Co-Proxamol	C		
UNKNOWN				Ciprofloxacin	C		
UNKNOWN				Amoxicillin	C	Glaxosmithkline	
UNKNOWN							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/06/04ISR Number: 4286748-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0321129A

Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Conjunctival Haemorrhage		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							
UNKNOWN				Amoxicillin	C	Glaxosmithkline	

Date:02/06/04ISR Number: 4290056-1Report Type:Direct Company Report #CTU 211692

Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Abnormal Behaviour		Wellbutrin 100 Mg Sr			
100 MG SR DAY		Completed Suicide		Glaxowellcome	PS	Glaxowellcome	ORAL
ORAL		Confusional State					
		Depression		Remeron	C		
		Impaired Driving Ability					
		Irritability					
		Medication Error					
		Social Avoidant Behaviour					

Date:02/06/04ISR Number: 4290313-9Report Type:Direct Company Report #CTU 211702

Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypersensitivity		Wellbutrin Sr 150 Mg	PS		ORAL
ONE DAY ORAL							

Date:02/09/04ISR Number: 4287490-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0416017A

Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

12.5MG Per day      6      DAY      Anxiety  
 Tension  
 Paxil Cr      PS      Glaxosmithkline  
 Wellbutrin Sr      SS      Glaxosmithkline

Date:02/09/04ISR Number: 4287664-0Report Type:Periodic      Company Report #US-GLAXOSMITHKLINE-A0423063A  
 Age:47 YR      Gender:Male      I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Paxil Cr	PS	Glaxosmithkline	
		Nausea		Wellbutrin	SS	Glaxosmithkline	
		Sexual Dysfunction					

Date:02/09/04ISR Number: 4287876-6Report Type:Periodic      Company Report #US-GLAXOSMITHKLINE-A0430895A  
 Age:40 YR      Gender:Male      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Paxil	PS	Glaxosmithkline	ORAL
12.5MG Per day	6	DAY					
		Diarrhoea					
		Dyskinesia		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Per day	4	WK					
		Euphoric Mood		Ambien	C		
		Libido Decreased					
		Panic Attack					
		Weight Decreased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/09/04ISR Number: 4287900-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431593A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Paxil	PS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	ORAL
				Paxil	C	Glaxosmithkline	ORAL

Date:02/09/04ISR Number: 4288013-4Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439982A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Withdrawal Syndrome	Consumer	Paxil	PS	Glaxosmithkline	ORAL
31 WK		Emotional Disorder		Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day	2 MON	Mood Swings		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Twice		Weight Increased					
per day	100 WK			Prenatal Vitamins	C		
				Aloe Vera	C		

Date:02/09/04ISR Number: 4288074-2Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0441759A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Paxil	PS	Glaxosmithkline	ORAL
25MG Per day				Wellbutrin	SS	Glaxosmithkline	

Date:02/09/04ISR Number: 4288165-6Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424006A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bladder Disorder		Paxil	PS	Glaxosmithkline	ORAL
6 WK							



75MG Per day      Condition Aggravated      Wellbutrin      SS      Glaxosmithkline      ORAL  
 Dizziness  
 Drug Ineffective

Date:02/09/04ISR Number: 4288169-3Report Type:Periodic      Company Report #US-GLAXOSMITHKLINE-A0424111A  
 Age:      Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
20MG Per day	1 YR	Libido Decreased		Paxil	PS	Glaxosmithkline	ORAL
		Urticaria		Wellbutrin	SS	Glaxosmithkline	ORAL

Date:02/09/04ISR Number: 4288209-1Report Type:Periodic      Company Report #US-GLAXOSMITHKLINE-A0424887A  
 Age:47 YR      Gender:Female      I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anxiety		Paxil	PS	Glaxosmithkline	ORAL
40MG Per day	6 YR	Crying		Wellbutrin	SS	Glaxosmithkline	ORAL
Initial or Prolonged		Depression		Methadone	C	Glaxosmithkline	ORAL
10MG Three		Drug Ineffective					
times per day		Drug Withdrawal Syndrome		Vicodin	C		ORAL
		Gastric Disorder		Premarin	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/09/04ISR Number: 4288424-7Report Type:Periodic  
Age:25 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428277A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
20MG Per day	3 YR	Anger		Paxil	PS	Glaxosmithkline	ORAL
150MG Twice		Anxiety		Wellbutrin	SS	Glaxosmithkline	ORAL
per day	2 WK	Crying					
		Drug Withdrawal Syndrome		Clonazepam	C		
		Libido Decreased					
		Sexual Dysfunction					

Date:02/09/04ISR Number: 4288598-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0495970A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
		Rectal Haemorrhage					

Date:02/09/04ISR Number: 4288600-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0496061A  
Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Hallucination					
		Overdose					

Date:02/09/04ISR Number: 4288604-0Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0312102A  
Age:26 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Arthralgia	Health	Zyban	PS	Glaxosmithkline	ORAL
22 DAY							
Initial or Prolonged		Body Temperature	Professional	Methylprednisolone	C		
1TAB Single							

dose

Increased

Drug Hypersensitivity  
 Eyelid Oedema  
 Face Oedema  
 Hypersensitivity  
 Inflammation  
 Myalgia  
 Oedema Peripheral  
 Rash Maculo-Papular  
 Rash Pruritic  
 Urticaria

Date:02/09/04ISR Number: 4288607-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0317612A  
 Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams		Zyban	PS	Glaxosmithkline	
150MG Twice		Anaphylactic Reaction					
per day		Dry Mouth		Cipramil	C		
		Headache		Microgynon	C		
		Night Sweats					
		Restless Legs Syndrome					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/09/04ISR Number: 4288610-6Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0319175A  
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Vaginal Haemorrhage	Health Professional	Bupropion Hydrochloride No Concurrent Medication	PS C	Glaxosmithkline	

Date:02/09/04ISR Number: 4288621-0Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0321501A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Parkinsonism	Health Professional	Zyntabac	PS	Glaxosmithkline	

Date:02/10/04ISR Number: 4289751-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0442807A  
Age:9 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other	150MG Per day 1 WK	Depressed Level Of Consciousness		Wellbutrin	PS	Glaxosmithkline	ORAL
	20MG per day	Grand Mal Convulsion		Metadate Cd	C		ORAL

Date:02/10/04ISR Number: 4289769-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0496663A  
Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide Intentional Misuse		Wellbutrin	PS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	

Date:02/10/04ISR Number: 4289778-8Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0319051A  
Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG	Convulsion		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization -	Variable dose 12 DAY	Epilepsy					
Initial or Prolonged	200MG Twice	Muscle Contractions		Tramadol	C		ORAL
per day		Involuntary					
		Paralysis		Painkiller	C		
		Salivary Hypersecretion					

Date:02/10/04ISR Number: 4294044-0Report Type:Expedited (15-DaCompany Report #2002IC000296  
Age:30 YR Gender:Male I/FU:I

Outcome	PT
Death	Body Temperature
Hospitalization -	Increased
Initial or Prolonged	Completed Suicide
	Convulsion
	Disseminated
	Intravascular Coagulation
	Drug Interaction
	Hepatic Failure
	Multiple Drug Overdose
	Nystagmus
	Posturing

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Pupil Fixed Renal Failure Rhabdomyolysis	Literature	Librium	PS		ORAL
ORAL		Suicide Attempt	Health Professional	Regular Strength Tylenol	SS		ORAL
ORAL		Tachycardia		Extra Strength Tylenol Pm	SS		ORAL
ORAL				Effexor	SS		ORAL
ORAL				Wellbutrin Sr	SS		ORAL
ORAL				Ambien	SS		ORAL
ORAL				Skelaxin	SS		ORAL
ORAL				Midol	SS		ORAL
ORAL				Pyridium	SS		ORAL
ORAL				Aleve	SS		ORAL
ORAL				Relafen	SS		ORAL

Date:02/11/04ISR Number: 4290995-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431566A  
 Age: YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective Sexual Dysfunction		Paxil	PS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	ORAL

Date:02/11/04ISR Number: 4291306-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442159A  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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10MG Per day	1	MON	Headache	Paxil	PS	Glaxosmithkline	ORAL
			Hyperhidrosis	Wellbutrin Narcotic	SS C	Glaxosmithkline	ORAL

Date:02/11/04ISR Number: 4291321-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442597A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Personality Change		Paxil	PS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	ORAL
				Nytol	SS	Glaxosmithkline	

UNKNOWN

Date:02/11/04ISR Number: 4291322-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442624A  
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Libido Decreased		Paxil	PS	Glaxosmithkline	ORAL
20MG Per day		Myalgia		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Twice							
per day				Lipitor	SS		

Date:02/11/04ISR Number: 4292238-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0442630A  
 Age:18 YR Gender:Male I/FU:F

Outcome	PT
	Drug Ineffective
	Medication Error
	Pharmaceutical Product

Freedom Of Information (FOI) Report

Complaint

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	6 DAY	Health Professional	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day	1 WK		Wellbutrin	SS	Glaxosmithkline	ORAL

Date:02/11/04ISR Number: 4292243-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0496778A  
 Age: YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Medication Error		Wellbutrin Unspecified Medications	PS C	Glaxosmithkline	ORAL

Date:02/11/04ISR Number: 4292256-3Report Type:Expedited (15-DaCompany Report #FI-GLAXOSMITHKLINE-B0321644A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN Initial or Prolonged	16 DAY	Angina Pectoris Breast Disorder Dyspnoea Muscle Spasms Urticaria		Zyban	PS	Glaxosmithkline	

Date:02/11/04ISR Number: 4292540-3Report Type:Direct Company Report #CTU 212065  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Abnormal Behaviour Aggression Anxiety Crying		Wellbutrin	PS		



Fatigue  
Fear  
Murder  
Panic Attack  
Paranoia  
Psychotic Disorder

Date:02/11/04ISR Number: 4294327-4Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0420816A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Aphthous Stomatitis		Lamictal	PS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	ORAL
1TAB Twice							
per day				Lexapro	C		

Date:02/12/04ISR Number: 4293009-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420604A

Outcome  
PT  
Drug Interaction  
Migraine  
Nausea

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vomiting

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
20MG Per day	2 YR		Paxil	PS	Glaxosmithkline	ORAL
100MG Per day	2 YR		Wellbutrin	SS	Glaxosmithkline	ORAL

Date:02/12/04ISR Number: 4293153-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0412455A  
 Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Paxil	PS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	
				Lamictal	C	Glaxosmithkline	
				Lexapro	C		

Date:02/12/04ISR Number: 4293717-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0496783A  
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depressed Level Of		Wellbutrin	PS	Glaxosmithkline	ORAL
Disability		Consciousness		Depakote	C		
300MG Per day		Grand Mal Convulsion		Oxycontin	C		
		Incontinence					
		Tongue Biting					

Date:02/12/04ISR Number: 4293724-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0099526A  
 Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Astrocytoma		Zyban	PS	Glaxosmithkline	ORAL
29 DAY		Body Temperature		Prozac	C		
Hospitalization -		Increased					
Initial or Prolonged		Coma					

Dysphagia  
Eating Disorder  
Incontinence  
Infection  
Memory Impairment  
Sleep Disorder  
Vomiting

Date:02/12/04ISR Number: 4293743-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0320609A

Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Astrocytoma	Consumer	Zyban	PS	Glaxosmithkline	ORAL
28 DAY				Fluoxetine Hydrochloride	C		

UNKNOWN

Date:02/12/04ISR Number: 4293746-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0321954A

Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Creatine		Zyban	PS	Glaxosmithkline	ORAL
Other		Phosphokinase Increased					
150MG		Muscle Spasms					
Variable dose	12 DAY						

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/12/04ISR Number: 4293747-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0321971A  
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression	Consumer	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Alcohol Use					
per day		Intentional Self-Injury		Ethanol	SS		
UNKNOWN				Beclomethasone Dipropionate	C	Glaxosmithkline	
UNKNOWN				Salbutamol	C	Glaxosmithkline	
UNKNOWN				Erythromycin	C	Glaxosmithkline	
UNKNOWN	5ML Four						
times per day							

Date:02/12/04ISR Number: 4293754-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0322806A  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Psoriasis		Zyban	PS	Glaxosmithkline	ORAL
150MG See							
dosage text	15 DAY						

Date:02/12/04ISR Number: 4295020-4Report Type:Direct Company Report #CTU 212221  
 Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - PO BID		Anaphylactic Reaction		Zyban 150 Mg	PS		ORAL
Initial or Prolonged							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Renal Failure Acute	Health Professional	Lexapro(Escitalopram ) Wellbutrin (Bupropion Hydrochloride) Lipitor (Atorvastatin) Avapro (Irbesartan) Tricor (Fenofibrate) Amiodarone Androderm Viagra (Sildenafil Citrate) Multivitamin Saw Pallmetto	PS   SS  C C C C C C C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1 WK Initial or Prolonged		Convulsion  Dizziness Road Traffic Accident	Health  Professional	Wellbutrin  Prozac	PS  C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/13/04ISR Number: 4294654-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441481A

Age: YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Drug Screen Positive	Health	Wellbutrin	PS	Glaxosmithkline	
UNKNOWN	9000MG	Single					
Hospitalization - dose		Intentional Misuse	Professional				
Initial or Prolonged		Suicide Attempt					
Other							

Date:02/13/04ISR Number: 4294666-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0321957A

Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hypoaesthesia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	5	DAY					
UNKNOWN		Panic Attack					
		Swollen Tongue		Zoladex	C		

Date:02/13/04ISR Number: 4294674-6Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0322531A

Age:69 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dizziness		Zyban	PS	Glaxosmithkline	ORAL
14 DAY							
UNKNOWN		Dry Mouth		Methyl dopa	C		
UNKNOWN		Insomnia		Gliclazide	C		
		Loss Of Consciousness					

Date:02/13/04ISR Number: 4294675-8Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0322550A

Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fall		Zyban	PS	Glaxosmithkline	ORAL
150MG	Unknown 3 DAY						
		Loss Of Consciousness					

Date:02/13/04ISR Number: 4297197-3Report Type:Expedited (15-DaCompany Report #2004006886

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Agitation	Health	Geodon (Ziprasidone)	PS		ORAL
ORAL							
Hospitalization -		Aspiration	Professional	Bupropion			
Initial or Prolonged		Cardiac Arrest	Company	Hydrochloride			
Other		Cardiac Disorder	Representative	(Bupropion			
		Intentional Misuse		Hydrochloride)	SS		
		Loss Of Consciousness		All Other			
		Multiple Drug Overdose		Therapeutic Products	SS		
		Suicide Attempt		Ethanol (Ethanol)	SS		
		Tachycardia					

Date:02/13/04ISR Number: 4342622-2Report Type:Periodic Company Report #352771

Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Alopecia	Consumer	Pegasys			
Intervention to		Anaemia Haemolytic		(Peg-Interferon			
Prevent Permanent		Autoimmune		Alfa-2a) 180 Mcg/Ml	PS		OTHER
180 MCG OTHER							
Impairment/Damage				Copegus (Ribavirin)			
				90 Mg/Ml	SS		ORAL
1000 MG DAILY							

ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Wellbutrin  
(Bupropion  
Hydrochloride)

SS

ORAL

1 DOSE FORM

DAILY ORAL

Date:02/16/04ISR Number: 4294947-7Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0399865A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Lamictal	PS	Glaxosmithkline	ORAL
37.5MG Per							
day	55 DAY			Wellbutrin Allegra	SS C	Glaxosmithkline	ORAL

Date:02/16/04ISR Number: 4295075-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441718A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Abortion Spontaneous	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	1 YR	Drug Exposure During Pregnancy	Professional				
Other		Pregnancy					

Date:02/16/04ISR Number: 4295094-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0316252A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alcohol Poisoning	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
		Coma		Ethanol	SS		ORAL
		Completed Suicide		Fluoxetine	C		ORAL
		Depressed Level Of		Flurazepam	C		ORAL



UNKNOWN	Consciousness	Lorazepam	C	
UNKNOWN	Drug Toxicity	Ibuprofen	C	Glaxosmithkline
UNKNOWN	Overdose	Albuterol	C	Glaxosmithkline
RESPIRATORY (INHALATION)	Pulmonary Congestion			
	Pulmonary Oedema			

Date:02/16/04ISR Number: 4295107-6Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0043117A  
Age:34 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Unknown Initial or Prolonged	Blood Triglycerides Increased Gamma-Glutamyltransferase Increased Hepatic Function Abnormal		Zyban	PS	Glaxosmithkline	ORAL

Date:02/16/04ISR Number: 4295340-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0413091A  
Age:28 YR Gender:Female I/FU:F

Outcome	PT Chills Deja Vu Hallucination Hypokinesia Pollakiuria
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Stereotypy

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
600MG Unknown	1 YR		Lamictal	PS	Glaxosmithkline	ORAL
300MG Unknown	1 YR		Wellbutrin	SS	Glaxosmithkline	ORAL
UNKNOWN	90MG Weekly		Prozac	C		
UNKNOWN	60MG Unknown		Inderal La	C		
			Lo/Ovral 28	C		

Date:02/16/04ISR Number: 4295677-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0420604A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction		Paxil	PS	Glaxosmithkline	ORAL
20MG Per day	2 YR	Migraine		Wellbutrin	SS	Glaxosmithkline	ORAL
100MG Per day	2 YR	Nausea Vomiting					

Date:02/16/04ISR Number: 4295697-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0491314A  
 Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aphasia	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Disorientation		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Per day	2 WK	Feeling Abnormal					
75MG See		Headache		Paxil	SS	Glaxosmithkline	
dosage text	1 MON	Insomnia		Compazine	C	Glaxosmithkline	
		Libido Decreased		Lorazepam	C		
		Medication Error					
		Mood Altered					
		Muscle Tightness					

Obsessive-Compulsive  
 Disorder  
 Salivary Hypersecretion  
 Tongue Disorder  
 Visual Acuity Reduced  
 Weight Increased

Date:02/16/04ISR Number: 4295700-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0497973A

Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG Twice		Agitation Coma		Thorazine Wellbutrin Sr	PS SS	Glaxosmithkline Glaxosmithkline	ORAL
per day							

Date:02/16/04ISR Number: 4295913-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442083A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		Psychotic Disorder		Lamictal Wellbutrin Lithium	PS SS SS	Glaxosmithkline Glaxosmithkline Glaxosmithkline	ORAL ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/04ISR Number: 4296165-5Report Type:Periodic  
Age:43 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0424884A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
500MG Per day		Alopecia		Lamictal	PS	Glaxosmithkline	ORAL
2 MON		Anxiety		Wellbutrin	SS	Glaxosmithkline	ORAL
500MG Unknown		Delirium		Lamictal	SS	Glaxosmithkline	ORAL
		Depression		Neurontin	C		
		Irritability		Effexor	C		
		Libido Increased		Xanax	C		
		Sensory Disturbance					

Date:02/17/04ISR Number: 4296262-4Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0428105A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oropharyngeal Swelling		Lamictal	PS	Glaxosmithkline	ORAL
		Throat Tightness		Wellbutrin	SS	Glaxosmithkline	ORAL

Date:02/17/04ISR Number: 4296284-3Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429256A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
8 MON		Anorexia		Lamictal	PS	Glaxosmithkline	ORAL
5 WK		Anxiety		Wellbutrin	SS	Glaxosmithkline	ORAL
25MG Per day		Blood Pressure Decreased		Zoloft	C		
1MG Per day		Bradycardia		Ativan	C		
		Food Intolerance					
		Hypertension					
		Irritable Bowel Syndrome					
		Labile Blood Pressure					
		Nervousness					
		Rash					

Vertigo  
Weight Decreased

Date:02/17/04ISR Number: 4296311-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430279A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Photosensitivity Reaction		Lamictal	PS	Glaxosmithkline	
				Wellbutrin	SS	Glaxosmithkline	
				Topomax	C		

Date:02/17/04ISR Number: 4296568-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441124A  
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
1	MON						

Date:02/17/04ISR Number: 4296584-7Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0319062A  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated		Zyban	PS	Glaxosmithkline	ORAL
Other		Psoriasis					
150MG per day	8 DAY						

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/04ISR Number: 4296586-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0320504A  
 Age:53 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 2TABS Per day 7 DAY	Blister		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged Disability	Burning Sensation Eye Swelling Local Swelling Oedema Peripheral Psoriasis Rash Rash Macular Swelling Face Thrombosis Toxic Skin Eruption Tremor					

Date:02/17/04ISR Number: 4299207-6Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20040201456  
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly	Complications Of Maternal Exposure To Therapeutic	Health Professional	Topiramate (Topiramate) Tablets	PS		
TRANSPLACENTAL L	TRANSPLACENTA Drugs					
	Congenital Anomaly Maternal Drugs Affecting Foetus		Wellbutrin (Bupropion Hydrochloride) S	SS		
TRANSPLACENTAL	TRANSLACENTAL Pregnancy Premature Baby Talipes		Prenatal Vitamin (Prenatal Vitamins) Tylenol (Paracetamol)	C C		

Date:02/17/04ISR Number: 4299266-0Report Type:Expedited (15-DaCompany Report #2004006886  
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Life-Threatening ORAL	Agitation	Health	Geodon (Ziprasidone)	PS	ORAL
Hospitalization - Initial or Prolonged Other	Aspiration Cardiac Arrest Cardiac Disorder Intentional Misuse Loss Of Consciousness Suicide Attempt Tachycardia	Professional Company Representative	Bupropion Hydrochloride (Bupropion Hydrochloride) All Other Therapeutic Products Ethanol (Ethanol)	SS   SS SS	

Date:02/18/04ISR Number: 4297488-6Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0321846A  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Chest Pain Face Oedema Muscle Spasms Oedema Peripheral Pruritus Generalised Rash	Consumer	Zyntabac No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/18/04ISR Number: 4299901-7Report Type:Expedited (15-DaCompany Report #B0307360A  
 Age:38 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other 150 MG/PER DAY/ ORAL	Brain Stem Infarction Eyelid Ptosis Headache Hiccups Hypoaesthesia Photosensitivity Reaction Pupils Unequal Sensory Disturbance Vomiting Wallenberg Syndrome White Blood Cell Count Increased	Foreign Literature Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:02/18/04ISR Number: 4300222-4Report Type:Direct Company Report #CTU 212559  
 Age:60 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 150 MG PO BID Intervention to Prevent Permanent Impairment/Damage	Headache Pruritus		Bupropion	PS		ORAL

Date:02/18/04ISR Number: 4300397-7Report Type:Direct Company Report #CTU 212512  
 Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Medication Error		Bupropion Hci Er (Generic Wellbutrin Sr) Mfr -Eon Labs	PS	Eon Labs	



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cerebrovascular Accident Myocardial Infarction	Consumer	Topamax (Topiramate) Tablets	PS		ORAL
50 MG, 2 IN 1 DAY, ORAL				Wellbutrin (Bupropion Hydrochloride) Unspecified	SS		
				Flomax (Tamsulosin Hydrochloride)	C		
				Celexa (Citalopram Hydrobromide)	C		
				Amantadine (Amantadine)	C		
				Protonix (Pantoprazole)	C		
				Lipztor (Atorvastatin)	C		
				Atenolol (Atenolol)	C		
				Lisinopril (Lisinopril)	C		
				Imdur (Isosorbide			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Mononitrate) C  
 Ultracet  
 (Tramadol/Apap) C  
 Celebrex (Celecoxib) C  
 Nitroquik (Glyceryl  
 Trinitrate) C

Date:02/19/04ISR Number: 4299463-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439627A  
 Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error	Health	Wellbutrin	PS	Glaxosmithkline	
		Pharmaceutical Product	Professional	Wellbutrin	SS	Glaxosmithkline	ORAL
450MG Per day	5 WK	Complaint		Thyroid	C		
				Provigil	C		

Date:02/19/04ISR Number: 4299469-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0498181A  
 Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG See Initial or Prolonged dosage text		Abortion Spontaneous		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Drug Exposure During Pregnancy		Paxil Cr	SS	Glaxosmithkline	ORAL
37.5MG See dosage text		Overdose					
		Suicide Attempt		Geodon	C		
				Neurontin	C		
				Folate	C		

Date:02/19/04ISR Number: 4299477-4Report Type:Expedited (15-DaCompany Report #FI-GLAXOSMITHKLINE-B0321644A  
 Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 15MG Per day	16 DAY	Angina Pectoris	Health	Zyban	PS	Glaxosmithkline	ORAL

Initial or Prolonged      Breast Disorder      Professional  
 Breast Pain  
 Bronchospasm  
 Burning Sensation  
 Dyspepsia  
 Dyspnoea  
 Mucous Membrane Disorder  
 Muscle Spasms  
 Oral Soft Tissue Disorder  
 Urticaria

Date:02/19/04ISR Number: 4299481-6Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0322716A  
 Age:35 YR    Gender:Female      I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG See		Abdominal Distension		Zyban	PS	Glaxosmithkline	ORAL
dosage text	14	DAY					
10MG per day		Anxiety Depression		Singulair	C		ORAL
500MG per day	4	DAY		Tranexamic Acid	C		ORAL
		Panic Attack Suicidal Ideation					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/19/04ISR Number: 4299482-8Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0322717A  
 Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG per day	7 DAY		Zyban	PS	Glaxosmithkline	ORAL
Other		Skin Reaction		Miconazole	C		
				Timodine	C		
TOPICAL							
TOPICAL							

Date:02/19/04ISR Number: 4299490-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0323573A  
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day			Zyban	PS	Glaxosmithkline	ORAL
Other		Visual Acuity Reduced					

Date:02/19/04ISR Number: 4299491-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0323706A  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG See	9 DAY		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization -		Increased					
Initial or Prolonged dosage text		Chest Pain					
		Electrocardiogram St					
		Segment Elevation					
		Lung Infiltration					
		Pericarditis					
		Pharynx Discomfort					

Date:02/19/04ISR Number: 4301504-2Report Type:Expedited (15-DaCompany Report #S03-USA-02425-01  
 Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged 10 MG QAM PO	Coma Grand Mal Convulsion	Health Professional	Lexapro (Escitalopram)	PS	ORAL
	Libido Decreased Tendon Injury	Company Representative	Wellbutrin - Slow Release (Bupropion Hydrochloride)	SS	
150 MG BID			Wellbutrin (Bupropion Hydrochloride)	SS	
100 MG QD			Dilantin (Phenytoin Sodium)	C	
			Keppra (Levetiracetam)	C	

Date:02/20/04ISR Number: 4300366-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0498768A  
Age: YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	15 DAY	Medication Error Pharmaceutical Product Complaint		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/20/04ISR Number: 4300368-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0498848A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:02/20/04ISR Number: 4300381-3Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0322567A

Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression	Consumer	Zyntabac	PS	Glaxosmithkline	ORAL
		Alcohol Interaction		Alcohol	SS		
		Disorientation					

Date:02/20/04ISR Number: 4300383-7Report Type:Expedited (15-DaCompany Report #IE-GLAXOSMITHKLINE-B0322640A

Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain		Zyban	PS	Glaxosmithkline	
300MG per day	2 WK	Granulocytopenia		No Concurrent Medication	C		
		Syncope					

Date:02/20/04ISR Number: 4300384-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0322714A

Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Skin Lesion		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY			Lansoprazole	C		ORAL
15MG per day				Tranexamic Acid	C		ORAL
1.5G Per day				Norethisterone	C		ORAL
5MG Per day							

Date:02/23/04ISR Number: 4301837-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0443651A  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Muscle Twitching					
per day	YR	Pharmaceutical Product		Vitamins	C		
		Complaint		Calcium	C		
				Advil	C	Glaxosmithkline	

Date:02/23/04ISR Number: 4301839-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0495505A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	
				Wellbutrin	SS	Glaxosmithkline	

Date:02/23/04ISR Number: 4301842-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0499051A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion		Bupropion	PS	Glaxosmithkline	
150MG Per day		Drug Exposure Via Breast Milk					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/23/04ISR Number: 4301849-6Report Type:Expedited (15-DaCompany Report #TR-GLAXOSMITHKLINE-B0314678A  
 Age:30 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 27 DAY Initial or Prolonged	Genital Lesion		Zyban	PS	Glaxosmithkline	ORAL
	Headache Pruritus					

Date:02/23/04ISR Number: 4301851-4Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0320703A  
 Age:47 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 150MG per day	Amnesia Convulsion		Bupropion Hydrochloride	PS	Glaxosmithkline	
400MG per day	Muscle Twitching		Tramadol	C		
	Salivary Hypersecretion		Salbutamol / Ipratropium Salmeterol Xinafoate + Fluticasone Propionate	C	Glaxosmithkline	
RESPIRATORY (INHALATION)			Codeine	C		ORAL
30MG per day			Morphine Sulphate	C	Glaxosmithkline	ORAL
20MG per day						

Date:02/23/04ISR Number: 4301857-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0322707A  
 Age:53 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150MG per day 20 DAY Other UNKNOWN	Psoriasis		Zyban	PS	Glaxosmithkline	ORAL
	Rash		Serevent	C	Glaxosmithkline	



UNKNOWN Toxic Skin Eruption Salbutamol C Glaxosmithkline  
UNKNOWN Becotide C Glaxosmithkline

Date:02/23/04ISR Number: 4301859-9Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0323130A  
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour		Zyban	PS	Glaxosmithkline	
150MG Twice		Aggression					
per day		Amnesia		No Concurrent Medication	C		

Date:02/23/04ISR Number: 4301861-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0323980A  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Confusional State		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown		Malaise		Neuroleptic	C		
		Sudden Death					
		Tremor					

Date:02/23/04ISR Number: 4303165-5Report Type:Expedited (15-DaCompany Report #USA-2002-0002070  
Age:33 YR Gender:Female I/FU:F

Outcome  
Death  
Hospitalization -

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Initial or Prolonged  
Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anoxic Encephalopathy Apallic Syndrome Circulatory Collapse Depression	Consumer Health Professional Other	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		
MG		Drug Abuser Drug Dependence		Trazadone (Trazadone) Unknown	SS		
MG		Loss Of Consciousness Medication Error		Propoxyphene (Dextrop ropoxyphene) Unknown	SS		
MG		Multi-Organ Failure Overdose		Acetaminophen (Parace tamol) Unknown	SS		
MG		Respiratory Arrest		Bupropion (Amfebutamo ne) Unknown	SS		

Date: 02/23/04  
Age: 51 YR  
Gender: Female  
I/FU: F

ISR Number: 4303226-0  
Report Type: Expedited (15-Da  
Company Report #USA-2003-0007481

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death		Arteriosclerosis Coma Drug Toxicity Fatigue Pericardial Effusion Somnolence Toxicologic Test Abnormal	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553) (Oxycodone Hydrochloride) Bupropion (Amfebutamo ne) Caffeine (Caffeine)	PS SS SS		

Date: 02/23/04  
Age: 48 YR  
Gender: Female  
I/FU: I

ISR Number: 4304057-8  
Report Type: Periodic  
Company Report #PHEH2003US10311

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Affect Lability	Consumer	Zelnorm (Tegaserod)			

6 MG, BID,	Depression	Tablet	PS	ORAL
ORAL	Drug Interaction			
	Gastroenteritis	Wellbutrin		
	Helicobacter	(Bupropion		
	Mood Swings	Hydrochloride)	SS	
		Klonopin(Clonazepam)	SS	
		Synthroid	C	
		Prozac (Fluoxetine		
		Hydrochloride)	C	
		Zinc (Zinc)	C	
		Trileptal "Novartis"	C	

Date:02/24/04ISR Number: 4302973-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0442630A  
Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	6 DAY	Drug Ineffective	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Medication Error	Professional	Wellbutrin	SS	Glaxosmithkline	ORAL
per day	1 WK	Pharmaceutical Product					
		Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/24/04ISR Number: 4302982-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0499451A  
Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -							
150MG Per day	2 DAY			Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged							
		Hypertension		Remeron	C		
		Ill-Defined Disorder		Enalapril	C		
		Oral Intake Reduced		Amaryl	C		
		Oral Soft Tissue Disorder		Glucophage	C		
		Speech Disorder					
		Stomach Discomfort					

Date:02/24/04ISR Number: 4302985-0Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0315453A  
Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other							
150MG Twice				Zyban	PS	Glaxosmithkline	ORAL
per day	MON						
		Neutrophilic Dermatitis					
		Dermatitis		No Concurrent			
		Hypersensitivity		Medication	C		
		Oral Discomfort					
		Rash Erythematous					
		Rash Papular					

Date:02/24/04ISR Number: 4303010-8Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0043010A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other							
150MG Five				Zyban	PS	Glaxosmithkline	ORAL
times per day							
		Suicide Attempt	Professional				
				Alcohol	SS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Acute Pulmonary Oedema	Health	Oxycodone			
		Adenomyosis	Professional	Hdyrochloride			
		Alcoholism	Other	(Similar To Nda			
		Beta Haemolytic		20-553)(Oxycodone			
		Streptococcal Infection		Hydrochloride)	PS		
		Bulimia Nervosa		Hydrocodone			
		Cardio-Respiratory Arrest		Bitartrate (Similar			
		Coagulopathy		To Ind 59,175)			
		Coma		(Hydrocodone			
		Contusion		Bitartrate) Unknown	SS		
		Culture Positive		Phentermine			
		Fall		(Phentermine)	SS		
		Gastric Haemorrhage		Bupropion			
		Haematemesis		(Amfebutamone)	SS		
		Hand Fracture		Caffeine (Caffeine)	SS		
		Hepatic Steatosis		Citalopram			
		Hepatomegaly		(Citalopram)	C		
		Leiomyoma					
		Lung Disorder					
		Oedema Peripheral					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/24/04ISR Number: 4305970-8Report Type:Expedited (15-DaCompany Report #HQWYE040517FEB04

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Activities Of Daily Living Impaired Crying Drug Withdrawal Syndrome Suicidal Ideation	Consumer	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		ORAL
SEE IMAGE				Wellbutrin (Amfebutamone Hydrochloride, )	SS		ORAL
ORAL							

Date:02/25/04ISR Number: 4303859-1Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0499286A

Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blindness		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Unknown	11 MON			Paxil	C	Glaxosmithkline	ORAL
40MG Unknown		Blindness Unilateral					

Date:02/25/04ISR Number: 4303860-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0499416A

Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Haemorrhage		Wellbutrin Coumadin	PS C	Glaxosmithkline Glaxosmithkline	ORAL

Date:02/25/04ISR Number: 4303861-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0499457A

Age:17 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Self Mutilation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Lexapro C  
Lithium C Glaxosmithkline

Date:02/25/04ISR Number: 4303872-4Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0323086A

Age:27 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Abortion Spontaneous		Zyban	PS	Glaxosmithkline	
UNKNOWN	150MG Twice					
per day						
			No Concurrent Medication	C		

Date:02/25/04ISR Number: 4303875-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0323413A

Age:22 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Arthralgia
Initial or Prolonged	C-Reactive Protein
	Increased
	Drug Exposure During
	Pregnancy
	Face Oedema
	Joint Swelling
	Mobility Decreased
	Neutrophil Percentage

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Increased Odynophagia Oedema Pregnancy	Report Source	Product	Role	Manufacturer	Route
150MG See dosage text	10 DAY	Urticaria White Blood Cell Count Increased		Zyban	PS	Glaxosmithkline	ORAL

Date:02/25/04ISR Number: 4303880-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0324053A  
Age:13 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	1 DAY	Life-Threatening	Accidental Overdose	Zyban	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged Other		Convulsion Hallucination Pulmonary Oedema Tachycardia					

Date:02/25/04ISR Number: 4303882-7Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0043117A  
Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Unknown 6 DAY	Hospitalization - Initial or Prolonged	Asthenia Dizziness	Zyban Beer	PS C	Glaxosmithkline	ORAL ORAL
1.5L per day		Hepatic Enzyme Increased Hepatic Function Abnormal Hyperlipidaemia					

Date:02/26/04ISR Number: 4304613-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0425696A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



76 DAY Tinnitus Wellbutrin PS Glaxosmithkline ORAL

Date:02/26/04ISR Number: 4304614-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430054A  
 Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	7 DAY	Drug Ineffective		Verapamil	C		
		Pharmaceutical Product		Doxazosin	C		
		Complaint					
		Stool Analysis Abnormal					

Date:02/26/04ISR Number: 4304615-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430617A  
 Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Drug Ineffective					
300MG Per day	15 DAY						

Date:02/26/04ISR Number: 4304616-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431217A  
 Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Grand Mal Convulsion		Zoloft	C		
4 DAY				Alcohol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304617-4Report Type:Periodic  
Age:66 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLIN-A0431224A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	30 DAY	Cough		Wellbutrin	PS	Glaxosmithkline	ORAL
		Therapeutic Response		Lipitor	C		
UNKNOWN	20MG	Unknown		Prilosec	C		
20MG	Unknown	Unexpected		Claritin	C		
UNKNOWN				Ocupress	C		

Date:02/26/04ISR Number: 4304618-6Report Type:Periodic  
Age:57 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLIN-A0432151A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Per day 2 WK	Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
		Energy Increased		Ventolin Mdi	C	Glaxosmithkline	
		Heart Rate Increased		Synthroid	C	Glaxosmithkline	
				Trazodone	C		
				Depakote	C		

Date:02/26/04ISR Number: 4304619-8Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLIN-A0432580A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	300MG Per day 3 WK	Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Vitamins	C		

Date:02/26/04ISR Number: 4304620-4Report Type:Periodic  
Age:15 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLIN-A0432616A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Grand Mal Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Concerta	C		
		Antibiotic (Unspecified)	C		

Date:02/26/04ISR Number: 4304621-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432633A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304622-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432859A  
 Age:16 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
		Anger		Sam E	C		
		Insomnia		Vitamins	C		

Date:02/26/04ISR Number: 4304623-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432860A  
 Age:26 YR Gender:Female I/FU:F

Outcome	PT
Life-Threatening	Abdominal Pain
Other	Abdominal Pain Upper

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anorexia Chromaturia Hepatitis Acute					
300MG Per day	3 DAY	Jaundice		Wellbutrin	PS	Glaxosmithkline	ORAL
		Nausea		Echinacea	C		
		Thrombocytopenia		Mircette	C		

Date:02/26/04ISR Number: 4304624-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432864A  
 Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Drug Interaction Heart Rate Increased Hyperventilation		Wellbutrin Lipitor	PS SS	Glaxosmithkline	

Date:02/26/04ISR Number: 4304625-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432865A  
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Paraesthesia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304626-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432879A  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	4 DAY	Crying		Robaxin	C		
				Zoloft	C		
				Tagamet	C	Glaxosmithkline	
				Benadryl	C	Glaxosmithkline	
				Singulair	C		
				Humibid	C		
				Dalmane	C		

Ambien	C	
Phenergan	C	Glaxosmithkline
Zofran	C	Glaxosmithkline
Ativan	C	
Zithromax	C	
Estrogen Injections	C	

Date:02/26/04ISR Number: 4304627-7Report Type:Periodic  
 Age:55 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432886A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK						

Trazodone	C	
Ativan	C	

Date:02/26/04ISR Number: 4304629-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432888A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hyperhidrosis		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	6 DAY						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304630-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432893A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	2 DAY	Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
		Joint Swelling					

Date:02/26/04ISR Number: 4304631-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0432982A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Unknown	Crying		Wellbutrin	PS	Glaxosmithkline	ORAL
		Eating Disorder					
		Purging					
		Weight Increased					

Date:02/26/04ISR Number: 4304632-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432999A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Insomnia					
		Nausea					

Date:02/26/04ISR Number: 4304633-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0433115A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	Visual Disturbance		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304634-4Report Type:Periodic  
Age:43 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0433117A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
15 DAY		Restlessness		Trazodone	C		
50MG per day	2 DAY						

Date:02/26/04ISR Number: 4304635-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0433119A  
 Age: YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mania		Wellbutrin	PS	Glaxosmithkline	
Other							

Date:02/26/04ISR Number: 4304636-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0433126A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304637-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0433138A  
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day	1 MON			Risperdal	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Trazodone C

Date:02/26/04ISR Number: 4304638-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0433168A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Flatulence		Wellbutrin	PS	Glaxosmithkline	

Date:02/26/04ISR Number: 4304639-3Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0433281A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL
1 WK		Palpitations		Antihypertensive	C		

Date:02/26/04ISR Number: 4304640-XReport Type:Periodic  
Age:13 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0433289A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Eruption		Wellbutrin	PS	Glaxosmithkline	ORAL
5 DAY		Urticaria					

Date:02/26/04ISR Number: 4304641-1Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0433299A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Contusion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	4 WK	Libido Decreased		Paxil Cr	SS	Glaxosmithkline	
25MG per day	1 MON	Urticaria		Toprol XL	C		
				Micardis	C	Glaxosmithkline	
				Benadryl	C	Glaxosmithkline	



Date:02/26/04ISR Number: 4304642-3Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0433300A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day				Depakote	C		

Date:02/26/04ISR Number: 4304643-5Report Type:Periodic  
Age:15 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0433338A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304644-7Report Type:Periodic  
Age:25 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0433339A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Grand Mal Convulsion					
150MG Three				Sonata	C		
times per day 11	DAY			Ortho Cyclen	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304645-9Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0438798A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bedridden		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	8 DAY	Dizziness Feeling Cold Nausea Vertigo Vomiting		Zoloft	C		

Date:02/26/04ISR Number: 4304647-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0438804A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	7 DAY	Headache		Xanax	C		

Date:02/26/04ISR Number: 4304648-4Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0438808A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Wellbutrin	SS	Glaxosmithkline	

Date:02/26/04ISR Number: 4304649-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0438831A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:02/26/04ISR Number: 4304650-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0438873A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression Anger		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304651-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0438914A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Apathy		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Depressed Mood Drug Ineffective Lethargy					

Date:02/26/04ISR Number: 4304652-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0438925A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304654-XReport Type:Periodic  
 Age: YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0439013A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	450MG Per day	Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304655-1Report Type:Periodic  
 Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0439014A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	Bruxism		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304656-3Report Type:Periodic  
 Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0439025A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	

Date:02/26/04ISR Number: 4304657-5Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439032A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	450MG Per day	Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304658-7Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0439049A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304659-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439057A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:02/26/04ISR Number: 4304660-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439109A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304661-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439110A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gynaecomastia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304662-9Report Type:Periodic  
Age:80 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0439231A

Outcome	PT
Other	Hallucination, Visual Hypertension

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Insomnia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
			Wellbutrin	PS	Glaxosmithkline	ORAL
			Exelon	C		
			Beta-Blocker(Unspeci fied)	C		
			Ace-Inhibitor	C		

Date:02/26/04ISR Number: 4304664-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439342A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day							

Date:02/26/04ISR Number: 4304665-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439425A  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK			Thyroid Replacement	C		
				Temazepam	C		

Date:02/26/04ISR Number: 4304666-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439432A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304667-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439435A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Ear Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1	MON		Hormone Pill	C		
		Oedema Peripheral		Multivitamin	C		
				Calcium	C		
				Vitamin C	C	Glaxosmithkline	

Date:02/26/04ISR Number: 4304669-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439440A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1	MON		Lipitor	C		

Date:02/26/04ISR Number: 4304670-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439478A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Nausea					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304672-1Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439609A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY			Prozac	C		

Date:02/26/04ISR Number: 4304673-3Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439615A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MGM Per day	5 DAY	Insomnia					
5 WK		Myalgia		Zyban	SS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304675-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0439616A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	4 WK	Drug Ineffective					

Date:02/26/04ISR Number: 4304676-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439621A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Galactorrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day				Lexapro	C		



Date:02/26/04ISR Number: 4304678-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439624A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1	MON		Lamictal	C	Glaxosmithkline	
				Risperdal	C		

Date:02/26/04ISR Number: 4304679-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439629A  
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2	WK		Depakote	C		
		Visual Disturbance		Altace	C		
				Risperdal	C		

Date:02/26/04ISR Number: 4304680-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439643A  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	4	WK					
		Rash					
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304681-2Report Type:Periodic  
Age:65 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439656A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK	Nausea		Unknown	C		

Date:02/26/04ISR Number: 4304682-4Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0439663A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	5 DAY	Dry Mouth					
		Fatigue					
		Feeling Drunk					
		Headache					

Date:02/26/04ISR Number: 4304683-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439679A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304686-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439818A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4 WK			Prozac	C		
				Benadryl	C	Glaxosmithkline	

Date:02/26/04ISR Number: 4304687-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439837A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Swelling Face					

Date:02/26/04ISR Number: 4304688-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439849A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aphasia		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion					

Date:02/26/04ISR Number: 4304689-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439980A  
Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Libido Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	13 DAY			Toprol	C		
				Avapro	C		
				Pravachol	C		
				Claritin	C		
				Aspirin	C	Glaxosmithkline	
				Folic Acid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Multivitamin C

Date:02/26/04ISR Number: 4304690-3Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439984A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	16 DAY	Muscle Tightness		Zomig	C		

Date:02/26/04ISR Number: 4304691-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440000A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	10 DAY			Cozaar	C		
				Klonopin	C		

Date:02/26/04ISR Number: 4304692-7Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440007A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Arthritis		Wellbutrin	PS	Glaxosmithkline	ORAL
		Tendonitis					

Date:02/26/04ISR Number: 4304693-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440204A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	7 DAY	Dry Mouth					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	450MG Per day	Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
				Synthroid	C	Glaxosmithkline	
				Lipitor	C		
				Prozac	C		
				Glucotrol	C		
				Zestril	C		
				Provigil	C		
				Dilantin	C		
				Centrum	C		
UNKNOWN				Vitamin C	C	Glaxosmithkline	
				Tums Ex	C	Glaxosmithkline	
				Prilosec	C		
UNKNOWN				Vitamin E	C		
UNKNOWN				Vitamin B	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304695-2Report Type:Periodic  
Age:72 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440208A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	Hyperglycaemia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Valtrex	C	Glaxosmithkline	
				Avandia	C	Glaxosmithkline	
				Hyzaar	C		
				Dyazide	C	Glaxosmithkline	

Date:02/26/04ISR Number: 4304696-4Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440210A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Unknown	Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304697-6Report Type:Periodic  
Age:34 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0440234A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Panic Attack Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304698-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440251A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304699-XReport Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0440349A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304700-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0440380A  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other	300MG Per day						

Date:02/26/04ISR Number: 4304701-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0440389A  
 Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other	300MG Per day 4 WK			Ativan	C		ORAL
	.5MG Twice						
	per day			Ambien	C		ORAL
	5MG At night			Nortriptyline	C		ORAL
	50MG At night						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304702-7Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440390A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1	MON	Stool Analysis Abnormal	Celebrex	C		

Date:02/26/04ISR Number: 4304703-9Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440397A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other			Grand Mal Convulsion	Clonazepam	C		
450MG Per day	4	DAY		Paxil	C	Glaxosmithkline	

Date:02/26/04ISR Number: 4304704-0Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440401A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	6	DAY	Dizziness	Pepto Bismol	C		
			Ear Discomfort				
			Nausea				

Date:02/26/04ISR Number: 4304705-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0440440A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day			Dizziness				

Date:02/26/04ISR Number: 4304706-4Report Type:Periodic  
Age:16 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440448A



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304707-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0440449A  
Age:17 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304708-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0440569A  
Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Joint Swelling		Neurontin	SS		

Date:02/26/04ISR Number: 4304709-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0440600A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304710-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440602A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Jittery Insomnia Nervousness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304711-8Report Type:Periodic  
Age:65 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440609A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Headache Hypoaesthesia Muscular Weakness Nausea		Lipitor	C		

Date:02/26/04ISR Number: 4304712-XReport Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440611A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Xanax	C		
				Pegasys	C		
				Copegus	C		

Date:02/26/04ISR Number: 4304713-1Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440613A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 DAY			Levoxyl	C	Glaxosmithkline	
				Armour Thyroid	C		

Nexium	C	
Bactrim	C	Glaxosmithkline
Flumadine	C	
Allegra	C	

Date:02/26/04ISR Number: 4304714-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0440697A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Jittery Insomnia Nervousness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304715-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0440711A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304716-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440798A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	4 DAY	Abnormal Dreams	Wellbutrin	PS	Glaxosmithkline	ORAL
			Anxiety				
			Crying				
			Feeling Abnormal				
			Insomnia				

Date:02/26/04ISR Number: 4304718-0Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440832A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Alopecia	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304719-2Report Type:Periodic  
 Age: YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0440848A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	3 DAY	Urticaria	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304720-9Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440851A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Grand Mal Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
Other	1 MON			Lexapro	C		

Date:02/26/04ISR Number: 4304721-0Report Type:Periodic  
 Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440984A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day	2	WK	Abdominal Pain Upper	Wellbutrin	PS	Glaxosmithkline	ORAL
				Anxiety	Lipitor	C		
				Constipation				
				Headache				
				Listless				
				Nausea				

Date:02/26/04ISR Number: 4304724-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0440993A  
Age:62 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG Per day	3	WK	Pruritus	Wellbutrin	PS	Glaxosmithkline	ORAL
				Urticaria	Toprol Xl	C		
					Lipitor	C		
					Aspirin	C	Glaxosmithkline	
					Antivert	C		

Date:02/26/04ISR Number: 4304725-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441012A  
Age:50 YR Gender:Female I/FU:F

Outcome	PT
	Anxiety
	Capillary Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Insomnia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1 WK		Wellbutrin	PS	Glaxosmithkline	ORAL
			Ativan	C		
			Estrace	C		

Date:02/26/04ISR Number: 4304726-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441018A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day	4 WK			Clonazepam	C		
				Doxepin	C		

Date:02/26/04ISR Number: 4304727-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441021A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK						

Date:02/26/04ISR Number: 4304728-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441024A  
 Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK						

Date:02/26/04ISR Number: 4304729-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441029A  
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
2	DAY			Seroquel	C		
				Klonopin	C		
				Lexapro	C		

Date:02/26/04ISR Number: 4304730-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441094A  
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
5	YR	Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
		Blood Triglycerides Increased		Zyprexa	C		

Date:02/26/04ISR Number: 4304731-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441098A  
 Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1 DAY	Menstrual Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
				Ortho Tricyclen	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304732-5Report Type:Periodic  
Age:15 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0441104A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
20 DAY		Grand Mal Convulsion Loss Of Consciousness					

Date:02/26/04ISR Number: 4304733-7Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0441105A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304734-9Report Type:Periodic  
Age: YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0441106A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304735-0Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0441107A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304736-2Report Type:Periodic  
Age:27 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441116A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



150MG Per day 4 DAY Anxiety Wellbutrin PS Glaxosmithkline ORAL  
Stool Analysis Abnormal Thyroid C

Date:02/26/04ISR Number: 4304738-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441117A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Per day 5 DAY		Blood Pressure Increased		Prevacid	C		
				Zyrtec	C	Glaxosmithkline	
				Vioxx	C		
				Multi-Vitamin	C		

Date:02/26/04ISR Number: 4304739-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441158A  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 1 DAY							

Date:02/26/04ISR Number: 4304740-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441164A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Vision Blurred		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304741-6Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441165A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				St Johns Wort	C		
				Lovastatin	C		
				Atacand	C		
				Glucophage	C		
				Glucotrol	C		
				Actos	C		

Date:02/26/04ISR Number: 4304742-8Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0441186A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304743-XReport Type:Periodic  
Age:41 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441432A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK	Fatigue					
		Headache					
		Nervousness					

Date:02/26/04ISR Number: 4304744-1Report Type:Periodic  
Age:51 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441435A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK			None	C		

Date:02/26/04ISR Number: 4304745-3Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441444A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:02/26/04ISR Number: 4304746-5Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441461A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK			Glipizide	C		

Date:02/26/04ISR Number: 4304747-7Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441468A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Myalgia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Lexapro	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304749-0Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441483A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nephrolithiasis		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK			Glucosamine	C		

Date:02/26/04ISR Number: 4304750-7Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441494A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	21 DAY			Adderall	C		
Hospitalization - Initial or Prolonged Other							

Date:02/26/04ISR Number: 4304751-9Report Type:Periodic  
Age:68 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0441503A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Atrioventricular Block		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	4 DAY						
Hospitalization - Initial or Prolonged Disability Other		Complete Rash Syncope					

Date:02/26/04ISR Number: 4304752-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441533A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				Toprol XL	C		
UNKNOWN	100MG Per day						

UNKNOWN 100MCG Per

Lotrel C  
Levoxyl C Glaxosmithkline

day

Date:02/26/04ISR Number: 4304753-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441631A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day 2	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
		WK					

Date:02/26/04ISR Number: 4304754-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441635A  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day 2	Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
	MON	Angioneurotic Oedema					
		Anxiety					
		Chest Pain					
		Dry Mouth					
		Hypoacusis					
		Nausea					
		Rash Maculo-Papular					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304755-6Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441638A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304756-8Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441642A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypertensive Crisis		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:02/26/04ISR Number: 4304757-XReport Type:Periodic  
Age:30 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0441651A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				Adderall	C		

Date:02/26/04ISR Number: 4304758-1Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441658A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1 DAY	Hallucination		Thyroid Medication	C		
		Road Traffic Accident					

Date:02/26/04ISR Number: 4304760-XReport Type:Periodic  
Age:25 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0441743A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Grand Mal Convulsion Wellbutrin PS Glaxosmithkline ORAL  
300MG Per day 2 WK  
Road Traffic Accident Lexapro C

Date:02/26/04ISR Number: 4304761-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441760A  
Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1	MON					

Date:02/26/04ISR Number: 4304762-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441769A  
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	24	DAY					
				Pamelor	C		
				Xanax	C		
				Ambien	C		

Date:02/26/04ISR Number: 4304763-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441775A  
Age:42 YR Gender:Female I/FU:I

Outcome	PT
Other	Amnesia Convulsion

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Periorbital Haematoma					
		Tooth Fracture					
		Tooth Injury	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Toothache		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK			Celexa	C		

Date:02/26/04ISR Number: 4304764-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442005A  
 Age: YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Dysuria		Wellbutrin Xl	PS	Glaxosmithkline	
Dose							

Date:02/26/04ISR Number: 4304765-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442013A  
 Age:29 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Abdominal Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Dose		Anxiety		Oral Contraceptives	C		
150MG Per day	2 WK	Diarrhoea					
		Dysmenorrhoea					

Date:02/26/04ISR Number: 4304766-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442024A  
 Age: YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Dose				Effexor Xr	C		ORAL
300MG Per day							
300MG Per day							

Date:02/26/04ISR Number: 4304767-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442029A  
 Age: YR Gender:Male I/FU:I



Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Bruxism		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 MON	Jaw Disorder		Ritalin	C		

Date:02/26/04ISR Number: 4304768-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442034A  
 Age: Gender:Male I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Fatigue		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day							

Date:02/26/04ISR Number: 4304769-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442035A  
 Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Fatigue		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day							

Date:02/26/04ISR Number: 4304770-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442045A  
 Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304771-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442050A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
3	WK						

Date:02/26/04ISR Number: 4304772-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442052A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG	Per day			Wellbutrin	SS	Glaxosmithkline	

Date:02/26/04ISR Number: 4304773-8Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0442053A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Weight Loss Poor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304774-XReport Type:Periodic  
 Age:68 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442188A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Per day 2 DAY	Depression Dizziness Nausea Psychomotor Hyperactivity Tremor		Lipitor	C		

Date:02/26/04ISR Number: 4304775-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442317A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 DAY	Amnesia		Advair	C	Glaxosmithkline	
		Chest Pain		Singulair	C		
				Allegra	C		
				Unknown Medication	C		

Date:02/26/04ISR Number: 4304776-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442343A  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Fatigue		Claritin	C		
3 WK		Grand Mal Convulsion		Flonase	C	Glaxosmithkline	
		Headache		Oral Contraceptive	C		

Date:02/26/04ISR Number: 4304777-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442355A  
Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304778-7Report Type:Periodic  
Age:25 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442356A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	
12 DAY				None	C		

Date:02/26/04ISR Number: 4304779-9Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442360A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2 MON	Anxiety		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Per day		Drug Ineffective		Hormone Replacement Therapy	C		

Date:02/26/04ISR Number: 4304780-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442436A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Musculoskeletal Stiffness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304781-7Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442511A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304782-9Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442512A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK						

Date:02/26/04ISR Number: 4304783-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442521A  
Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day		Parosmia		Chemotherapy	C		

Date:02/26/04ISR Number: 4304784-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442528A  
Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Somnolence		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1 MON						

Date:02/26/04ISR Number: 4304785-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442549A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Wellbutrin	PS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304786-6Report Type:Periodic  
 Age:34 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442550A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	5 YR			Tylenol Sinus	C		

Date:02/26/04ISR Number: 4304787-8Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442610A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Overdose		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1 MON						

Date:02/26/04ISR Number: 4304788-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442612A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 MON						

Date:02/26/04ISR Number: 4304789-1Report Type:Periodic  
 Age:54 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442652A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorgasmia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 MON						
		Inadequate Lubrication		Wellbutrin	SS	Glaxosmithkline	ORAL
100MG Twice							
per day	2 YR	Libido Decreased					
				Combipatch	C		
				Dhea	C		
				Dong Quai Root	C		

Date:02/26/04ISR Number: 4304790-8Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442653A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	0 DAY			Wellbutrin	PS	Glaxosmithkline	ORAL
		Abdominal Pain Upper					
		Insomnia					
		Mouth Ulceration					
		Nervousness					
		Night Sweats					
		Pruritus					

Date:02/26/04ISR Number: 4304791-XReport Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0442727A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other				Wellbutrin	PS	Glaxosmithkline	
		Convulsion		Unspecified Medications	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304792-1Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0442730A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	
450MG Per day							

Date:02/26/04ISR Number: 4304793-3Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442764A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK			Effexor	C		

Date:02/26/04ISR Number: 4304794-5Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442777A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Headache		Zocor	C		
				Zestril	C		

Date:02/26/04ISR Number: 4304795-7Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442780A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 DAY	Dry Mouth		Topamax	C		
				Estrogen	C		
				Thyroid	C		



Date:02/26/04ISR Number: 4304796-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442797A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flushing Hot Flush		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304797-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0443028A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304798-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0443052A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Movement Disorder Tic		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304799-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443101A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:02/26/04ISR Number: 4304800-8Report Type:Periodic  
Age:77 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443171A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 2 WK							
		Restlessness		Tamoxifen	C		
		Tremor		Diovan	C		
				Prednisone	C		
				Neurontin	C		
				Nexium	C		
				Megace	C		

Date:02/26/04ISR Number: 4304801-XReport Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0443178A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Discomfort		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304802-1Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443207A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 2 WK							

Date:02/26/04ISR Number: 4304803-3Report Type:Periodic  
Age:20 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0443215A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304804-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0443243A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Unknown		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304805-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0443357A  
 Age:22 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day		Grand Mal Convulsion		Adderall Xr	C		
10MG Per day				Focalin	C		
				Ativan	C		
				Ortho Tri-Cyclen	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304806-9Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443471A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	6 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
		Paraesthesia		Hctz	C		

Date:02/26/04ISR Number: 4304807-0Report Type:Periodic  
Age:16 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443487A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day			Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion		Unknown	C		

Date:02/26/04ISR Number: 4304808-2Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443493A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	1 MON		Wellbutrin	PS	Glaxosmithkline	ORAL
		Anorexia					
		Fatigue					
		Headache					
		Insomnia					
		Somnolence					
		Weight Increased					

Date:02/26/04ISR Number: 4304809-4Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443501A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day			Wellbutrin	PS	Glaxosmithkline	ORAL
		Pruritus		Unknown	C		

Date:02/26/04ISR Number: 4304831-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443503A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1	WK		Effexor	C		

Date:02/26/04ISR Number: 4304832-XReport Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443506A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304833-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443531A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gastrooesophageal Reflux Disease		Wellbutrin	PS	Glaxosmithkline	ORAL
				Wellbutrin	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304834-3Report Type:Periodic  
 Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443569A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	5 DAY	Balance Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dizziness					

Date:02/26/04ISR Number: 4304835-5Report Type:Periodic  
 Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443571A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG In the morning		Balance Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Feeling Abnormal					
		Impaired Driving Ability					

Date:02/26/04ISR Number: 4304836-7Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443654A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 WK	Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
		Diarrhoea		Wellbutrin	SS	Glaxosmithkline	
		Energy Increased					
		Tearfulness					

Date:02/26/04ISR Number: 4304837-9Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443656A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304838-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0443664A  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK	Irritability		Zoloft	C		
				Insulin	C		

Date:02/26/04ISR Number: 4304839-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0443665A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK			Lithium	C	Glaxosmithkline	

Date:02/26/04ISR Number: 4304840-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0443670A  
Age:15 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Anaphylactic Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1 DAY	Dyspnoea					
Hospitalization -							
Initial or Prolonged							
Other							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304841-0Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443679A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Acne		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304842-2Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443680A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304843-4Report Type:Periodic  
 Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443758A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	1 MON	Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Allegra-D	C		
				Nasonex	C		

Date:02/26/04ISR Number: 4304844-6Report Type:Periodic  
 Age:75 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443797A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
450MG Per day	4 WK	Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Drug Ineffective		Valium	C		
				Zoloft	C		

Date:02/26/04ISR Number: 4304845-8Report Type:Periodic  
 Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443902A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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150MG Per day 3 DAY	Insomnia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Klonopin	C		
		Estrogen	C		
		Cozaar	C		
		Zocor	C		
		Atenolol	C		

Date:02/26/04ISR Number: 4304846-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0443905A  
 Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 5 DAY		Insomnia		Arthritis Medication	C		

Date:02/26/04ISR Number: 4304847-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0443926A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fatigue		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
50MG Per day 3 MON				Sleeping Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304848-3Report Type:Periodic  
 Age:21 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443944A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	2 WK	Tinnitus	Wellbutrin	PS	Glaxosmithkline	ORAL
				Ddavp	C		
				Cipro	C	Glaxosmithkline	

Date:02/26/04ISR Number: 4304850-1Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0443984A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Bradyphrenia	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304851-3Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0444029A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Adverse Event	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304852-5Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0444056A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Rash	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304853-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0444067A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Rash Pruritic	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304854-9Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0444102A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3	WK					

Date:02/26/04ISR Number: 4304855-0Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0444160A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus Rash Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304856-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0444169A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304857-4Report Type:Periodic  
Age:66 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0490776A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	5 WK						

Date:02/26/04ISR Number: 4304858-6Report Type:Periodic  
Age:35 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0490787A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Epistaxis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6 MON			Trileptal	C		ORAL
600MG At							
night				Ambien	C		ORAL
10MG At night				Vitamin Complex	C		ORAL

Date:02/26/04ISR Number: 4304859-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0490827A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Uveitis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304860-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0490852A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Back Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							
Initial or Prolonged		Insomnia		Restoril	C		
		Mouth Ulceration		Ambien	C		
				Blood Pressure Medication	C		

Date:02/26/04ISR Number: 4304861-6Report Type:Periodic  
Age:70 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0490856A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Thyroid Medication	C		

Date:02/26/04ISR Number: 4304862-8Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0490954A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	7 DAY			Xanax	C		
.25MG As							
required							

Date:02/26/04ISR Number: 4304863-XReport Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0490955A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Migraine		Ritalin	C		
				Kdur	C	Glaxosmithkline	
				Metadate	C		
				Allegra D	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nexium C  
Relpax C

Date:02/26/04ISR Number: 4304864-1Report Type:Periodic  
Age:27 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0490957A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3	WK	Drug Ineffective	Wellbutrin	PS	Glaxosmithkline	ORAL
Pharmaceutical Product Complaint							

Date:02/26/04ISR Number: 4304865-3Report Type:Periodic  
Age:21 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0490960A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Per day	2	MON	Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
Alcohol Unspecified Medications							
C							

Date:02/26/04ISR Number: 4304866-5Report Type:Periodic  
Age:38 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0490961A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
40MG Per day			Muscle Twitching	Wellbutrin	PS	Glaxosmithkline	
				Prozac	C		
20MG Per day				Prevacid	C		
				Hemocyte Plus	C		
10MG Per day				Zyrtec	C	Glaxosmithkline	

Date:02/26/04ISR Number: 4304867-7Report Type:Periodic  
Age:49 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0490976A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2	MON		Paxil Cr	C	Glaxosmithkline	ORAL
				Neurontin	C		
				Estrogen	C		

Date:02/26/04ISR Number: 4304868-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0490979A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Irritability Mood Altered		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304869-0Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0490980A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3	WK					
		Pain Pruritus					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304870-7Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0490982A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
3TAB Single		Dyspepsia					
dose		Lethargy					
		Overdose					

Date:02/26/04ISR Number: 4304871-9Report Type:Periodic  
 Age:56 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0491277A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Twitching		Wellbutrin Xl	PS	Glaxosmithkline	
150MG In the							
morning				Norvasc	C		
5MG Per day				Imitrex	C	Glaxosmithkline	
50MG As							
required				Multivitamin	C		
				Aleve	C		
				Dyazide	C	Glaxosmithkline	
				Estratest	C		

Date:02/26/04ISR Number: 4304872-0Report Type:Periodic  
 Age:51 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491312A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 WK	Nausea		Buspar	C		
				Ativan	C		



Date:02/26/04ISR Number: 4304873-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491331A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Dysgeusia Insomnia					

Date:02/26/04ISR Number: 4304874-4Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491334A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Distension		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	15 DAY						

Date:02/26/04ISR Number: 4304875-6Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491358A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:02/26/04ISR Number: 4304881-1Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491689A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG See							
dosage text	3	WK					

Date:02/26/04ISR Number: 4304882-3Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491691A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dry Mouth		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6	MON					
		Feeling Abnormal		Trileptal	C		
		Keratoconjunctivitis					
		Sicca					

Date:02/26/04ISR Number: 4304883-5Report Type:Periodic  
Age: YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0491704A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Agitation		Wellbutrin	PS	Glaxosmithkline	
		Anorexia					
		Insomnia					
		Panic Attack					
		Tremor					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304884-7Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491706A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Therapeutic Response Decreased		Wellbutrin	PS	Glaxosmithkline	

Date:02/26/04ISR Number: 4304885-9Report Type:Periodic  
Age: YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0491719A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sexual Dysfunction		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1	MON					

Date:02/26/04ISR Number: 4304886-0Report Type:Periodic  
Age:75 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491723A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Behaviour		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1	MON					
		Constipation		Wellbutrin	SS	Glaxosmithkline	ORAL
		Dissociative Identity		Ritalin	SS		ORAL
14	MON	Disorder					
		Gallbladder Operation					

Date:02/26/04ISR Number: 4304887-2Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491733A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day							

Date:02/26/04ISR Number: 4304888-4Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491745A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Foreign Body Trauma		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	6 WK	Insomnia		Clarinet	C		
				Copaxone	C		

Date:02/26/04ISR Number: 4304889-6Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491880A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Nervousness		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:02/26/04ISR Number: 4304890-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491884A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Neck Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
3 WK							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304891-4Report Type:Periodic  
 Age:27 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491886A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1	MON		None	C		
		Urticaria					

Date:02/26/04ISR Number: 4304892-6Report Type:Periodic  
 Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0491889A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flushing		Wellbutrin	PS	Glaxosmithkline	

Date:02/26/04ISR Number: 4304893-8Report Type:Periodic  
 Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491898A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3	WK					
		Agitation					
		Depression					
		Therapeutic Response					
		Unexpected					
		Weight Decreased					

Date:02/26/04ISR Number: 4304894-XReport Type:Periodic  
 Age:39 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0491909A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1	WK					
		Grand Mal Convulsion					

Date:02/26/04ISR Number: 4304895-1Report Type:Periodic  
Age:86 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491924A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Dizziness Nausea					

Date:02/26/04ISR Number: 4304896-3Report Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491926A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Contusion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 3 WK		Rash		Ritalin	C		

Date:02/26/04ISR Number: 4304897-5Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492006A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 2 WK		Dysuria Incontinence Micturition Urgency		Prozac Provigil	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304899-9Report Type:Periodic  
Age:17 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0492008A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	450MG Per day	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304900-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492012A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG Per day	Urine Odour Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304901-4Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492021A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day 3 DAY	Abdominal Distension		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304902-6Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492022A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day 1 MON	Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Visual Disturbance		Zoloft	C		

Date:02/26/04ISR Number: 4304903-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492025A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



300MG Per day 6 MON Headache Wellbutrin PS Glaxosmithkline ORAL

Date:02/26/04ISR Number: 4304904-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0492208A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Psychomotor Hyperactivity Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304905-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0492258A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Palpitations		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

150MG Per day

Date:02/26/04ISR Number: 4304906-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0492269A  
Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Panic Reaction Weight Decreased		Geodon	C		

150MG Per day

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304907-5Report Type:Periodic  
Age:41 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492408A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eye Irritation		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Keratoconjunctivitis					
		Sicca					
		Vision Blurred					

Date:02/26/04ISR Number: 4304908-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492410A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				Valium	C		ORAL
2MG Unknown				Prozac	C		ORAL
40MG Unknown				Seroquel	C		
UNKNOWN	100MG Unknown						

Date:02/26/04ISR Number: 4304909-9Report Type:Periodic  
Age:22 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492411A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK	Hyperhidrosis		Amoxicillin	C	Glaxosmithkline	
		Rash		Tylenol 3	C		

Date:02/26/04ISR Number: 4304910-5Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0492435A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Night Sweats		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304911-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492560A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 4 WK Initial or Prolonged		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
				Unknown Medication	C		

Date:02/26/04ISR Number: 4304912-9Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492578A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	

Date:02/26/04ISR Number: 4304913-0Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0492592A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 450MG Per day 2 MON		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304914-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492696A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Irritability		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304915-4Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0492709A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Intolerance		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304916-6Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0492884A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	

Date:02/26/04ISR Number: 4304917-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492896A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Disturbance In Attention		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Irritability		Prozac	C		
				Lexapro	C		

Date:02/26/04ISR Number: 4304918-XReport Type:Periodic  
Age:34 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0492897A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	12 DAY	Abnormal Dreams		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dizziness		Klonopin	C		

Middle Insomnia  
Nervousness  
Panic Attack  
Tremor

Date:02/26/04ISR Number: 4304919-1Report Type:Periodic  
Age:60 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492900A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Euphoric Mood		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	8 DAY			Plavix	C		
				Lopressor	C		
				Pravacol	C		
				Ativan	C		

Date:02/26/04ISR Number: 4304920-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492906A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Exfoliative		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Rash					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304921-XReport Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492913A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG Per day	Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304922-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492916A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG Per day	Sinus Tachycardia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Topamax	C		

Date:02/26/04ISR Number: 4304923-3Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493116A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day	Dry Mouth		Wellbutrin	PS	Glaxosmithkline	ORAL
		Energy Increased Feeling Jittery Palpitations					

Date:02/26/04ISR Number: 4304924-5Report Type:Periodic  
 Age:25 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493257A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304925-7Report Type:Periodic  
 Age:57 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493262A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	7 DAY	Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hyperhidrosis		Zocor	C		
		Insomnia		Vioxx	C		
		Nausea		Atenolol	C		
				Imdur	C		

Date:02/26/04ISR Number: 4304926-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0493276A  
Age: YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Menstrual Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304927-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0493281A  
Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	1 MON	Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Estroven	C		
				Multivitamins	C		
				Advair	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304928-2Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493313A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							
				Methadone	SS	Glaxosmithkline	
UNKNOWN							

Date:02/26/04ISR Number: 4304929-4Report Type:Periodic  
 Age:44 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0493452A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
150MG See							
dosage text	4	MON		Paxil	C	Glaxosmithkline	
80MG Per day	10	YR		Prozac	C		
80MG per day							

Date:02/26/04ISR Number: 4304930-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493464A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	10	DAY					
				Interferon	C		
				Adalat	C	Glaxosmithkline	

Date:02/26/04ISR Number: 4304931-2Report Type:Periodic  
 Age:28 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493492A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day							



Euphoric Mood  
Fatigue

Zoloft

C

Date:02/26/04ISR Number: 4304932-4Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493493A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4 DAY	Vision Blurred					

Date:02/26/04ISR Number: 4304933-6Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493559A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:02/26/04ISR Number: 4304934-8Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493562A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Withdrawal Syndrome		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304935-XReport Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493576A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304936-1Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493580A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304937-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493774A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304938-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493783A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety Fatigue Nervousness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304939-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493801A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304940-3Report Type:Periodic  
Age: YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0493938A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300TAB Per							
day				Prozac	C		

Date:02/26/04ISR Number: 4304941-5Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493941A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Herpes Simplex		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 MON	Urticaria		Vitamins	C		

Date:02/26/04ISR Number: 4304942-7Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493950A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Wellbutrin Xl	PS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304943-9Report Type:Periodic  
Age:68 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493966A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Insomnia					
		Tremor		Prilosec	C		
				Lipitor	C		
				Zestril	C		

Date:02/26/04ISR Number: 4304944-0Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493970A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Amnesia					
1TAB Per day	3	MON		Evista	SS		
		Arthralgia					
		Dry Mouth					
		Feeling Hot					
		Visual Disturbance					

Date:02/26/04ISR Number: 4304945-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494032A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
UNKNOWN							
		Drug Interaction		Wellbutrin	PS	Glaxosmithkline	ORAL
		Hypertension		Unspecified Drug	SS		

Date:02/26/04ISR Number: 4304946-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494236A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hypersensitivity					

Date:02/26/04ISR Number: 4304947-6Report Type:Periodic  
Age:71 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494238A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK	Disturbance In Attention		Wellbutrin	SS	Glaxosmithkline	ORAL
		Feeling Abnormal		Xanax	C		
		Feeling Drunk		Vioxx	C		
				Ambien	C		
				Vitamin C	C	Glaxosmithkline	
				Vitamine B12	C	Glaxosmithkline	

Date:02/26/04ISR Number: 4304948-8Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494250A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304950-6Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494268A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 MON			Gabitril	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Toprol XL C  
 Trazodone C  
 Fiber C  
 Oxazepam C

Date:02/26/04ISR Number: 4304951-8Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494273A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day	Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Cardura	C		
				Aspirin	C	Glaxosmithkline	
				Multivitamin	C		

Date:02/26/04ISR Number: 4304952-XReport Type:Periodic  
 Age: YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0494277A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304953-1Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0494322A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Muscle Twitching		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304954-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494333A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	2 WK	Lip Blister		Wellbutrin XL	PS	Glaxosmithkline	ORAL
		Rash					

Rash Pruritic

Date:02/26/04ISR Number: 4304956-7Report Type:Periodic  
Age: YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0494459A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Generalised		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304957-9Report Type:Periodic  
Age:26 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0494469A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY			Zoloft	C		

Date:02/26/04ISR Number: 4304958-0Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494490A

Outcome	PT
	Insomnia
	Menstrual Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Suicidal Ideation

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	20 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
			Xanax	C		
			Nexium	C		

Date:02/26/04ISR Number: 4304959-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0494492A  
 Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 DAY	Chest Discomfort		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dry Mouth					
		Headache					
		Palpitations					
		Tremor					

Date:02/26/04ISR Number: 4304960-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0494519A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dyspnoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Yawning					

Date:02/26/04ISR Number: 4304961-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0494534A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Pharmaceutical Product Complaint					



Date:02/26/04ISR Number: 4304962-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494549A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Breast Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Unknown		Nipple Pain Nipple Swelling					

Date:02/26/04ISR Number: 4304963-4Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494682A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 2 WK		Blood Pressure Increased		Actonel	C		
		Drug Ineffective		Calcium	C		
		Increased Appetite		Aspirin	C	Glaxosmithkline	
		Medication Error		Vitamin E	C		
		Sensory Disturbance					
		Stool Analysis Abnormal					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304965-8Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0494685A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Grand Mal Convulsion Incontinence Tongue Biting		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304966-XReport Type:Periodic  
Age:25 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494692A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day							

Date:02/26/04ISR Number: 4304967-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494697A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	2 WK	Anxiety Dry Mouth Dyspepsia Flushing Headache Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304968-3Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494708A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3 WK	Insomnia Rash Generalised		Wellbutrin Xl Vitamins	PS C	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304969-5Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494718A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day				Synthroid	C	Glaxosmithkline	
				Xanax	C		

Date:02/26/04ISR Number: 4304970-1Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494721A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day 3 MON		Tinnitus		Synthroid	C	Glaxosmithkline	
UNKNOWN		Weight Increased		Diovan	C		

Date:02/26/04ISR Number: 4304971-3Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494725A

Outcome	PT
	Anxiety
	Chest Pain
	Confusional State

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Impaired Driving Ability  
Nervousness

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3 DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Cipro	C	Glaxosmithkline	
			Desogen	C		
			Sudafed	C	Glaxosmithkline	

Date:02/26/04ISR Number: 4304972-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0494728A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Menstrual Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Lexapro	SS		
20MG Per day				Oxycontin	C		
				Benadryl	C	Glaxosmithkline	

Date:02/26/04ISR Number: 4304973-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0494875A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Suicidal Ideation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304974-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0494895A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Mydriasis		Wellbutrin	PS	Glaxosmithkline	ORAL
				Effexor	SS		

Date:02/26/04ISR Number: 4304975-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0495026A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Migraine		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 DAY			Neurontin	C		
				Seroquel	C		
				Effexor	C		

Date:02/26/04ISR Number: 4304976-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0495029A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vulvovaginal Dryness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304977-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0495039A  
 Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304978-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495042A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Ranitidine	C	Glaxosmithkline	

Date:02/26/04ISR Number: 4304979-8Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495059A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:02/26/04ISR Number: 4304980-4Report Type:Periodic  
Age:39 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0495270A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	9 DAY			Nicorette	C	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304981-6Report Type:Periodic  
Age:17 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0495274A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Hospitalization - 150MG In the Initial or Prolonged morning	20 DAY						

Date:02/26/04ISR Number: 4304982-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495333A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
1560MG Per day	6	WK	Abdominal Pain Upper Fluid Retention	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Headache	Thyroid	C		
			Insomnia	Estrogen	C		
			Jaw Disorder	Soma	C		
			Nausea	Lasix	C	Glaxosmithkline	
			Pain In Jaw				
			Pharmaceutical Product Complaint				
			Weight Increased				

Date:02/26/04ISR Number: 4304983-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0495337A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				Effexor Xr	C		

Date:02/26/04ISR Number: 4304984-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0495501A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Convulsion	Wellbutrin	PS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304985-3Report Type:Periodic  
Age:18 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495504A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4304986-5Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0495507A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	
300MG Per day	3 WK			Reglan	SS	Glaxosmithkline	
10MG Per day							

Date:02/26/04ISR Number: 4304987-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495547A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR			Microzide	C		ORAL

Date:02/26/04ISR Number: 4304988-9Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495551A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 MON						

Date:02/26/04ISR Number: 4304989-0Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0495567A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Amnesia

Wellbutrin Xl

PS

Glaxosmithkline

ORAL

Date:02/26/04ISR Number: 4304990-7Report Type:Periodic  
Age:46 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0495721A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK			Vitamin B Complex	C		
				Dostinex	C		

Date:02/26/04ISR Number: 4304992-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495722A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	5 WK	Irritability					

Date:02/26/04ISR Number: 4304993-2Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495724A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Chest Discomfort					
		Heart Rate Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4304994-4Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495725A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6	MON		Sinemet	C		
				Comtan	C		
				Mirapex	C		
				Flomax	C		
				Lasix	C	Glaxosmithkline	
				Synthroid	C	Glaxosmithkline	

Date:02/26/04ISR Number: 4304995-6Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495734A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Lithium	C	Glaxosmithkline	

Date:02/26/04ISR Number: 4304996-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495755A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Delirium		Tenormin	C		
300MG Per day	1	Neuralgia		Valium	C		
		Neuropathy					
		Vertigo					

Date:02/26/04ISR Number: 4304997-XReport Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495757A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Photosensitivity Reaction		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2	WK					

Lexapro C  
Ambien C  
Clonazepam C

Date:02/26/04ISR Number: 4304998-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0495760A  
Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Generalised		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3	DAY						

Date:02/26/04ISR Number: 4304999-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0495766A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flatulence		Wellbutrin Xl	PS	Glaxosmithkline	

Date:02/26/04ISR Number: 4305000-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0495773A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 MON	Sleep Disorder		Nexium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Celexa C  
 Blood Pressure  
 Medication C

Date:02/26/04ISR Number: 4305001-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495775A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hallucination		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:02/26/04ISR Number: 4305002-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495779A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Feeling Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	5 DAY			Zocor	C		

Date:02/26/04ISR Number: 4305003-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495783A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:02/26/04ISR Number: 4305004-5Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495805A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Rash					

Date:02/26/04ISR Number: 4305005-7Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495931A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Burning Sensation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	8 WK	Pruritus					
		Rash					

Date:02/26/04ISR Number: 4305006-9Report Type:Periodic  
Age:17 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495974A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day				None	C		

Date:02/26/04ISR Number: 4305007-0Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496028A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Adverse Event		Wellbutrin	PS	Glaxosmithkline	ORAL
		Rash					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4305008-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496270A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG per day	Confusional State		Wellbutrin	PS	Glaxosmithkline	ORAL
		Decreased Appetite Paranoia					

Date:02/26/04ISR Number: 4305009-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496278A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	2 MON	Crying Depression		Wellbutrin Wellbutrin	PS C	Glaxosmithkline Glaxosmithkline	ORAL
		Drug Ineffective					

Date:02/26/04ISR Number: 4305010-0Report Type:Periodic  
Age:44 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496282A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day 1 MON	Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	1 MON			Lexapro	SS		
				Advair	C	Glaxosmithkline	
				Flonase	C	Glaxosmithkline	

RESPIRATORY  
(INHALATION)

Date:02/26/04ISR Number: 4305011-2Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496332A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							

Date:02/26/04ISR Number: 4305012-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496515A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4305013-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496516A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Decreased Dizziness Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4305014-8Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496936A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Somnolence		Wellbutrin Xl Respordol Zoloft	PS C C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4305015-XReport Type:Periodic  
 Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497386A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Twice per day	Heart Rate Increased Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/26/04ISR Number: 4305016-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497387A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin Xl	PS	Glaxosmithkline	

Date:02/26/04ISR Number: 4305019-7Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497388A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl Wellbutrin Sr	PS C	Glaxosmithkline Glaxosmithkline	ORAL ORAL

Date:02/26/04ISR Number: 4305073-2Report Type:Expedited (15-Da  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499436A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:02/26/04ISR Number: 4305078-1Report Type:Expedited (15-Da  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499594A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL



Date:02/26/04ISR Number: 4305798-9Report Type:Direct  
Age:32 YR Gender:Female I/FU:I

Company Report #CTU 213269

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 500 MG/1 DAY	Arrhythmia		Wellbutrin-Glaxo Xl	PS	Glaxo	
Initial or Prolonged	Convulsion Influenza Like Illness Loss Of Consciousness Rash					

Date:02/26/04ISR Number: 4307414-9Report Type:Expedited (15-DaCompany Report #200410862BCC  
Age:15 YR Gender: I/FU:I

Outcome	PT
Hospitalization -	Coma
Initial or Prolonged	Convulsion Electrocardiogram Qt Prolonged Intentional Misuse International Normalised Ratio Decreased Lethargy Suicide Attempt Syncope

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tachycardia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
2640 MG, ONCE, ORAL		Health Professional Other	Aleve Tablet 220mg From Bayer Consumer Care (Naproxen Sodium)	PS		ORAL
3750 MG, ONCE, ORAL			Wellbutrin 150 Mg (Bupropion Hydrochloride)	SS		ORAL
12.5 MG, ONCE, ORAL			Risperdal M-Tab Disintegrating Tablet 0.5 Mg From Janssen Pharmaceutica	SS	Janssen Pharmaceutical	ORAL
ONCE, ORAL			Percocet (5/325) Endo Laboratories	SS	Endo Laboratories	ORAL

Date:02/27/04ISR Number: 4306881-4Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0323085A  
Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	19 DAY	Arthralgia C-Reactive Protein Increased Haematoma Headache Joint Swelling Myalgia		Zyban  No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

Nausea  
 Neutrophilia  
 Pyrexia  
 Rash Generalised  
 Stevens-Johnson Syndrome  
 Urticaria

Date:02/27/04ISR Number: 4306886-3Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0323610A  
 Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 8 DAY Initial or Prolonged		Deep Vein Thrombosis		Zyban	PS	Glaxosmithkline	ORAL

Date:02/27/04ISR Number: 4306888-7Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0323712A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death UNKNOWN per day	150MG Twice 4 DAY	Circulatory Collapse Dyspnoea Rash Status Asthmaticus		Zyban  Inhalers	PS  C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/27/04ISR Number: 4306895-4Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0324788A

Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Missed		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	22 DAY	Complications Of Maternal Exposure To Therapeutic Drugs Intra-Uterine Death Pregnancy On Oral Contraceptive		Oral Contraceptive	C		

Date:02/27/04ISR Number: 4307730-0Report Type:Direct

Company Report #CTU 213330

Age:15 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Overdose		Paxil 10 Mg Bid Wellbutrin 30 Mg	PS SS		

Date:02/27/04ISR Number: 4308727-7Report Type:Expedited (15-DaCompany Report #USA-2003-0011315

Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Accidental Overdose Cerebral Haemorrhage Coma Drug Screen Positive Drug Toxicity Hypertensive Heart Disease Somnolence	Health Professional Other	Morphine Sulfate (Similar To Nda 19-516)(Morphine Sulfate) Unknown Hydroxyzine (Hydroxyzine) Carisoprodol (Carisoprodol) Nicotine (Nicotine) Cannabis (Cannabis) Temazepam (Temazepam) Bupropion (Amfebutamone)	PS SS SS SS SS SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Decreased Drug Interaction		Glucophage Tabs 850 Mg	PS	Bristol-Myers Squibb Company	ORAL
dose				Neurontin	C		
increased				Vioxx	C		
from 200 to				Allegra	C		
400 mg 3.5				Lisinopril	C		
hours before				Prilosec	C		
				Provigil	I		ORAL
				Wellbutrin	I		ORAL
				Celexa	I		ORAL
				Avandia	I		
				Humalog	I		
INTRAMUSCULAR	unit	QAM/MD					

Freedom Of Information (FOI) Report

INTRAMUSCULAR unit QAM/PM

Humulin Regular I

Date:03/02/04ISR Number: 4309086-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0440829A  
 Age:14 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
		Claustrophobia		Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day		Confusional State		Xanax	C		
		Convulsion		Vitamins	C		
		Disorientation					
		Disturbance In Attention					
		Dizziness					
		Flashback					
		Loss Of Consciousness					
		Palpitations					
		Therapeutic Response					
		Unexpected					
		Weight Decreased					

Date:03/02/04ISR Number: 4309090-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0499621A  
 Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Twice							
per day							

Date:03/02/04ISR Number: 4309322-6Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0321501A  
 Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Parkinsonism	Health Professional	Zyntabac	PS	Glaxosmithkline	

Outcome	PT
Life-Threatening	Arrhythmia
Hospitalization -	Blood Bicarbonate
Initial or Prolonged	Decreased
Other	Blood Calcium Decreased
	Blood Glucose Increased
	Blood Potassium Decreased
	Body Temperature
	Increased
	Depressed Level Of
	Consciousness
	Drug Screen Positive
	Electrocardiogram Qt
	Corrected Interval
	Prolonged
	Nausea
	Prothrombin Time
	Prolonged
	Red Blood Cell
	Sedimentation Rate

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
	Increased					
	Respiratory Arrest					
	Syncope					
	Torsade De Pointes					
	Ventricular Extrasystoles	Study	Morphine Sulfate			
	Ventricular Tachycardia	Health	(Similar To Nda			
	Vomiting	Professional	19-516) (Morphine			
	White Blood Cell Count	Other	Sulfate)	PS		ORAL
ORAL	Increased		Vistaril (Hydroxyzine Embonate)	SS		
			Benadryl (Diphenhydramine Hydrochloride)	SS		
			Metformin (Metformin)	SS		
			Lipitor (Atorvastatin)	SS		
			Effexor (Venlafaxine Hydrochloride)	SS		
			Lasix (Furosemide)	SS		
			Potassium (Potassium)	SS		
			Reglan (Metoclopramide)	SS		
			Amfetamine (Amfetamine)	SS		
			Ambien (Zolpidem Tartrate)	SS		
			Acebutolol Hydrochloride (Acebutolol Hydrochloride)	SS		
			Baclofen (Baclofen)	SS		
			Nexium (Esomeprazole)	SS		
			Prilosec (Omeprazole)	SS		
			Quinine (Quinine)	SS		
			Promethazine (Promethazine)	SS		
			Klonopin (Clonazepam)	SS		
			Wellbutrin (Amfebutamone Hydrochloride)	SS		
			Celexa (Citalopram			



Hydrobromide)

C

Date:03/02/04ISR Number: 4311167-8Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 213582

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Prozac 20 Mg	PS		ORAL
20 MG BID							
ORAL							
100 MG QD				Wellbutrin Sr 100 Mg	SS		ORAL
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/03/04ISR Number: 4310527-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441482A  
 Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthritis	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice			Professional				
per day	5 WK			Minocycline	C		ORAL
100MG Twice							
per day							

Date:03/03/04ISR Number: 4310533-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0499844A  
 Age: YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Completed Suicide		Bupropion	PS	Glaxosmithkline	
Death		Intentional Misuse					

Date:03/03/04ISR Number: 4310543-7Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0314544A  
 Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Zyban	PS	Glaxosmithkline	
Other		Anxiety					
22 DAY		Headache					
		Nightmare					
		Panic Attack					
		Psychotic Disorder					
		Sleep Disorder					
		Syncope					

Date:03/03/04ISR Number: 4310550-4Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0323130A  
 Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Twice	Abnormal Behaviour	Health	Zyban	PS	Glaxosmithkline	
	per day	Aggression	Professional				
		Alcohol Poisoning		No Concurrent Medication	C		
	1 DAY	Amnesia					
		Thinking Abnormal					

Date:03/03/04ISR Number: 4310559-0Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0323958A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	18 DAY	Pericarditis		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Sensation Of Pressure					

Date:03/03/04ISR Number: 4311498-1Report Type:Direct Company Report #CTU 213716  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150 MG. DAILY	Amnesia		Wellbutrin 150 Mg.	PS		ORAL
		Convulsion					
							ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/03/04ISR Number: 4311592-5Report Type:Direct  
Age:32 YR Gender:Female I/FU:I

Company Report #CTU 213690

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent ORAL Impairment/Damage		Aphasia Vocal Cord Disorder		Wellbutrin Sr 100 Mg Glaxo	PS	Glaxo	ORAL
				Lexapro	C		

Date:03/04/04ISR Number: 4311279-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0433343A  
Age:2 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Accidental Exposure Convulsion	Health Professional	Avandia Wellbutrin Metformin Lipitor	PS SS SS SS	Glaxosmithkline Glaxosmithkline	ORAL ORAL ORAL ORAL

Date:03/04/04ISR Number: 4311280-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0499548A  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other TRANSDERMAL		Psoriasis Psoriatic Arthropathy 6 WK Rheumatoid Arthritis		Wellbutrin Nicoderm Cq 21mg	PS SS	Glaxosmithkline Glaxosmithkline	ORAL

Date:03/04/04ISR Number: 4311281-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0499807A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:03/04/04ISR Number: 4311282-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0499947A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Adverse Event Completed Suicide		Wellbutrin	PS	Glaxosmithkline	

Date:03/05/04ISR Number: 43112277-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0357840A

Age:37 YR Gender:Female I/FU:F

Outcome	PT
Life-Threatening	Abdominal Discomfort
Hospitalization -	Abortion Spontaneous
Initial or Prolonged	Aggression
Other	Agitation
	Anger
	Anxiety
	Complications Of Maternal
	Exposure To Therapeutic
	Drugs
	Confusional State
	Convulsion
	Dependence
	Dizziness
	Drug Dependence

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
5 MON		Drug Ineffective Drug Withdrawal Syndrome Influenza					
150MG Twice		Insomnia		Paxil	PS	Glaxosmithkline	
per day		Maternal Drugs Affecting Foetus		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
.5MG As		Motor Dysfunction Nausea		Effexor Ativan	C C		
required		Nightmare					
25MG Per day		Pregnancy		Atenolol	C		
		Psychotic Disorder Pyrexia Sensory Disturbance Suicidal Ideation Suicide Attempt Tremor Vomiting		Alcohol Valium	C C		ORAL

Date:03/05/04ISR Number: 4312298-9Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0319984A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Pain Upper Anorexia Balance Disorder Constipation Deafness Disturbance In Attention Dysgeusia Ear Disorder Feeling Abnormal Nausea Sleep Disorder Tinnitus		Zyban	PS	Glaxosmithkline	

Date:03/05/04ISR Number: 4312303-XReport Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0323131A  
Age:25 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Aggression		Zyban	PS	Glaxosmithkline	
UNKNOWN	150MG	Twice					
per day		Alcohol Use					
				Alcohol	SS		
				No Concurrent			
				Medication	C		

Date:03/05/04ISR Number: 4312305-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0323706A  
Age:41 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Antinuclear Antibody
Initial or Prolonged	Positive
	C-Reactive Protein
	Increased
	Chest Pain
	Electrocardiogram St
	Segment Elevation
	Leukocytosis
	Lung Infiltration

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG See dosage text	9 DAY	Neutrophil Count Increased Pericarditis Pharynx Discomfort Viral Pericarditis	Zyban	PS	Glaxosmithkline	ORAL

Date:03/05/04ISR Number: 4312309-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0324489A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2 WK Initial or Prolonged		PT Anaphylactic Reaction Dysphagia Dyspnoea Local Swelling Pharyngolaryngeal Pain Rash Pruritic	Zyban No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:03/05/04ISR Number: 4312330-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0325566A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Other 150MG Per day	5 DAY	PT Abortion Spontaneous	Zyban	PS	Glaxosmithkline	ORAL

Date:03/08/04ISR Number: 4313521-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0500831A  
Age: Gender:Male I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Other		PT Drug Abuser Medication Error	Wellbutrin	PS	Glaxosmithkline	NASAL



Date:03/08/04ISR Number: 4313524-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0501213A

Age: YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:03/09/04ISR Number: 4314396-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0501024A

Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2	MON		Wellbutrin	SS	Glaxosmithkline	

Date:03/09/04ISR Number: 4314412-8Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0324869A

Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Oropharyngeal Swelling		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	18	DAY		Nurofen	C	Glaxosmithkline	ORAL
		Swelling Face					
		Urticaria Generalised					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/09/04ISR Number: 4314413-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0324872A

Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice	Cardiac Arrest		Zyban	PS	Glaxosmithkline	ORAL
per day							

Date:03/10/04ISR Number: 4314967-3Report Type:Direct

Company Report #CTU 214172

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 60 MG AT		Grand Mal Convulsion Petit Mal Epilepsy		Remeron 60 Mg (Generic)	PS	Generic	
BEDTIME							
ONE TAB Q AM							
				Wellbutrin Xl 300 Mg	SS		
				Adderall Xr	C		
				Pamelor	C		
				Zyprexa	C		
				Trazodone	C		
				Ativan	C		

Date:03/10/04ISR Number: 4315134-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0501711A

Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 300MG Unknown	139 DAY	Hemiplegia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged Disability		Pain Paraesthesia Simple Partial Seizures Syncope					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Twice	Coma	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - per day	Initial or Prolonged	Renal Failure Acute	Professional	Lexapro	C		
Disability	10MG Per day			Avapro	C		
Other				Coumadin	C	Glaxosmithkline	
				Amiodarone	C		
				Lipitor	C		
				Saw Palmetto	C		
				Multivitamin	C		
				Tricor	C		
				Viagra	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	15MG Per day	Amnesia	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Other	40MG Per day	Electroencephalogram	Professional	Zyprexa	C		ORAL
		Abnormal		Celexa	C		ORAL
		Grand Mal Convulsion					
		Medication Error					
		Postictal State					
		Tongue Biting					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/11/04ISR Number: 4315625-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0502223A  
 Age:9 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:03/11/04ISR Number: 4315806-7Report Type:Direct Company Report #USP 56383  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG		Medication Error		Wellbutrin Xl	PS		
SUSTAINED							
RELEASE							
TABLET							

Date:03/11/04ISR Number: 4316643-XReport Type:Expedited (15-DaCompany Report #A0500156A  
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Condition Aggravated Depression Dizziness	Consumer	Nicorette Otc Gum 4 Mg (Nicotine Polacrilex)	PS		
4		Drug Dependence					
MG/TRANSBUCCA		Dry Mouth					
L		Feeling Abnormal Irritability		Nicoderm Cq Patch 7 Mg (Nicotine)	SS		
TRANSDERMAL	7	Lymphoma					
MG/TRANSDERMA		Neoplasm Recurrence					
L		Pallor Pharyngolaryngeal Pain		Wellbutrin (Formulation			

ORAL

Unknown) (Bupropion Hydrochloride)	SS
Atomoxetine Hydrochloride	C
Clonazepam	C

Date:03/12/04ISR Number: 4316190-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441497A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
50MG Per day		Abnormal Behaviour		Paxil Cr	PS	Glaxosmithkline	ORAL
		Amnesia		Contac	SS	Glaxosmithkline	ORAL
		Attention		Wellbutrin	SS	Glaxosmithkline	ORAL
		Deficit/Hyperactivity		Paxil	SS	Glaxosmithkline	ORAL
		Disorder		Xanax	C		
		Confusional State		Clonopin	C		
		Depression					
		Diarrhoea					
		Disturbance In Attention					
		Drug Interaction					
		Dysphoria					
		Fatigue					
		Initial Insomnia					
		Nausea					
		Obsessive-Compulsive					
		Disorder					
		Psychotic Disorder					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/12/04ISR Number: 4316206-6Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0323086A  
 Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day 16 DAY 6 DAY	31 DAY	Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus Pregnancy		Zyban  Bisolvin Linctus Klacid Xl	PS  C C	Glaxosmithkline	ORAL  ORAL ORAL

Date:03/12/04ISR Number: 4316750-1Report Type:Direct Company Report #USP 56332  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose TABLET TABLET		Medication Error		Wellbutrin Xl Wellbutrin Sr	PS SS	Glaxosmithkline	

Date:03/12/04ISR Number: 4317238-4Report Type:Expedited (15-DaCompany Report #KII-2003-0007821  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other  ORAL  ORAL		Agitation Anion Gap Abnormal Blood Alcohol Increased Confusional State Convulsion Depressed Level Of Consciousness  Dysarthria Hypertension	Study Health Professional Other	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet Ethanol (Ethanol) Fluoxetine (Fluoxetine)  Bupropion (Amfebutamone)	   PS SS  SS  SS		        ORAL  ORAL

Intention Tremor  
Multiple Drug Overdose  
Serotonin Syndrome  
Tachycardia  
Tremor  
Vomiting

Date:03/15/04ISR Number: 4317635-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0496783A

Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	21 DAY	Depressed Level Of Consciousness Dizziness Grand Mal Convulsion Incontinence Myalgia Tongue Biting Visual Disturbance		Depakote Ms Contin	C C	Glaxosmithkline	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/15/04ISR Number: 4317638-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0502146A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1	MON	Agitation	Wellbutrin	PS	Glaxosmithkline	
150MG Per day			Medication Error	Wellbutrin	SS	Glaxosmithkline	ORAL
			Pharmaceutical Product Complaint	Clonazepam	C		

Date:03/15/04ISR Number: 4317640-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0502570A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Twice			Medication Error	Wellbutrin	PS	Glaxosmithkline	
per day	2	YR	Pharmaceutical Product Complaint	Wellbutrin	SS	Glaxosmithkline	ORAL

Date:03/15/04ISR Number: 4317650-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0325523A

Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Pruritus Generalised	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice			Tongue Ulceration				
per day	21	DAY					

Date:03/15/04ISR Number: 4317651-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0325526A

Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Unintended Pregnancy	Zyban	PS	Glaxosmithkline	ORAL
150MG Per day							



UNKNOWN	7	YR	Microval	SS	
100MG Per day			Sertraline	C	ORAL
			Kapake	C	ORAL
TOPICAL			Ibuprofen	C	Glaxosmithkline

Date:03/15/04ISR Number: 4317653-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0325531A  
 Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	9 DAY	Abnormal Sensation In Eye		Zyban	PS	Glaxosmithkline	ORAL
2.5MG per day		Blister Oral Discomfort		Bendrofluazide	C	Glaxosmithkline	ORAL
UNKNOWN		Rash Erythematous Stevens-Johnson Syndrome Tongue Discolouration		Felodipine	C		

Date:03/15/04ISR Number: 4317693-XReport Type:Expedited (15-DaCompany Report #US-ROCHE-360924  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Drug Interaction		Klonopin Wellbutrin	PS I	Roche	ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/15/04ISR Number: 4318218-5Report Type:Expedited (15-DaCompany Report #GBWYE626508MAR04  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Complications Of Maternal Exposure To Therapeutic Drugs	Health Professional Other	Microval (Levonorgestrel, Tablet)	PS		ORAL
TAKEN		Drug Ineffective					
REGULARLY FOR		Drug Interaction					
SEVEN YEARS		Unintended Pregnancy		Zyban (Amfebutamone Hydrochloride, , 0)	SS		ORAL
150 MG				Sertraline (Sertraline)	C		
				Kapake (Codeine Phosphate/Paracetamo l)	C		
				Ibuprofen (Ibuprofen)	C		

Date:03/15/04ISR Number: 4318671-7Report Type:Expedited (15-DaCompany Report #B0325227A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Back Crushing Convulsion Spinal Fracture	Foreign Literature Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochoride)	PS		
UNKNOWN	UNK, UNK;						

UNKNOWN

Date:03/16/04ISR Number: 4317364-XReport Type:Direct Company Report #CTU 214424  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150 MG BID

Oedema Peripheral

Wellbutrin

PS

Pruritus

Date:03/16/04ISR Number: 4317366-3Report Type:Direct Company Report #CTU 214426

Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Wellbutrin Xl	PS		
150 MG DAILY		Pruritus					
		Tremor					

Date:03/16/04ISR Number: 4317826-5Report Type:Expedited (15-DaCompany Report #US-ROCHE-360924

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated		Klonopin	PS	Roche	ORAL
Other		Convulsion		Wellbutrin	I		ORAL
		Drug Interaction					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/16/04ISR Number: 4317932-5Report Type:Expedited (15-DaCompany Report #US-ROCHE-360924

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Klonopin	PS	Roche	ORAL
		Drug Interaction		Wellbutrin	I		ORAL

Date:03/16/04ISR Number: 4317936-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0493122A

Age:25 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG See							
Initial or Prolonged		Concussion					
dosage text	23 DAY						
Other		Fall		Clomipramine	C		
150MG At							
night		Grand Mal Convulsion					
		Medication Error		Seroquel	C		ORAL
25MG At night		Skull Fracture					

Date:03/16/04ISR Number: 4317960-XReport Type:Expedited (15-DaCompany Report #DK-GLAXOSMITHKLINE-B0325421A

Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Arthralgia		Zyban	PS	Glaxosmithkline	ORAL
21 DAY							
		Joint Swelling					
		Lymphadenopathy					
		Urticaria					

Date:03/16/04ISR Number: 4317963-5Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0325690A

Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Angioneurotic Oedema	Zyban	PS	Glaxosmithkline	ORAL
150MG Per day 8 DAY	Cheilitis	Serevent	C	Glaxosmithkline	
	Eye Disorder	Pulmicort	C		
RESPIRATORY	Pruritus				
(INHALATION)	Swelling Face				
	Urticaria				

Date:03/16/04ISR Number: 4318481-0Report Type:Direct Company Report #USP 214507  
 Age:47 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Choking		Wellbutrin Xl 300			
Hospitalization -	Confusional State		Mg, 150mg, Glaxo	PS	Glaxo	
300 MG PO Q						
Initial or Prolonged	Convulsion					
AM ADDITIONAL						
	Loss Of Consciousness					
150 MG QD X 2						
	Skin Discolouration					
D, THEN 300						
	Tongue Biting					
MG QD X 3D;						
			Lexapro	C		
			Ambien	C		
			Zyprexa	C		
			Protonix	C		
			Tylenol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/16/04ISR Number: 4319296-XReport Type:Expedited (15-DaCompany Report #GBWYE626508MAR04

Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complications Of Maternal Exposure To Therapeutic Drugs	Health Professional Other	Microval (Levonorgestrel, Tablet, 0)	PS		ORAL
TAKEN		Drug Ineffective					
REGULARLY		Drug Interaction					
OVER SEVEN		Pregnancy On Oral					
YEARS		Contraceptive Unintended Pregnancy		Zyban (Amfebutamone Hydrochloride, , 0)	SS		ORAL
150MG				Sertraline (Sertraline)	C		
				Kapake (Codeine Phosphate / Paracetamol)	C		
				Ibuprofen (Ibuprofen)	C		

Date:03/16/04ISR Number: 4320048-5Report Type:Expedited (15-DaCompany Report #USA-2003-0011528

Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Bronchitis Chronic Cerebral Infarction Chronic Obstructive Pulmonary Disease Coma Drug Screen Positive Emphysema Pleural Adhesion Pneumonitis Pyelonephritis Chronic Renal Disorder Scar	Health Professional Other	Morphine Sulfate (Morphine Sulfate) Hydrocodone Bitartrate (Hydrocodone Bitartrate) Dihydrocodeine/Caffeine/Acetaminophen (Dihydrocodeine, Caffeine, Paracetamol) Acetaminophen (Paracetamol) Olanzapine	PS SS SS SS		

(Olanzapine)	SS
Amitriptyline	
(Amitriptyline)	SS
Trazodone	
(Trazodone)	SS
Nicotine (Nicotine)	SS
Caffeine (Caffeine)	SS
Bupropion	
(Amfebutamone)	SS

Date:03/17/04ISR Number: 4318139-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0502264A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly	150MG Per day	35 DAY	Drug Exposure During Pregnancy	Bupropion	PS	Glaxosmithkline	
			Trisomy 18				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/17/04ISR Number: 4318142-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0502603A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Myasthenia Gravis	Health Professional	Wellbutrin Trazodone	PS SS	Glaxosmithkline	

Date:03/18/04ISR Number: 4319549-5Report Type:Direct Company Report #CTU 214745

Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 100 MG BID		Condition Aggravated		Wellbutrin 100	PS		
Initial or Prolonged 150 MG TID		Granuloma		Wellbutrin 150	SS		
		Mental Disorder Nephritis Interstitial Proteinuria Pyuria Renal Failure					

Date:03/19/04ISR Number: 4319737-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441348A

Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Antepartum Haemorrhage		Bupropion	PS	Glaxosmithkline	ORAL
300MG per day		Drug Exposure During Pregnancy					

Date:03/19/04ISR Number: 4319746-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0502845A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
1TAB Per day		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL



150MG Twice  
per day  
Pharmaceutical Product  
Complaint  
Wellbutrin  
SS  
Glaxosmithkline  
ORAL

Date:03/19/04ISR Number: 4319747-0Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0502949A  
Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Corneal Bleb		Zyban	PS	Glaxosmithkline	ORAL
150MG See		Drug Interaction					
dosage text		Eye Pain		Alcohol	SS		ORAL
8Z See dosage		Keratoconjunctivitis					
text		Sicca		Antihypertensives	C		

Date:03/19/04ISR Number: 4319749-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0502995A  
Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
4 WK		Dyspnoea					
		Loss Of Consciousness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/19/04ISR Number: 4319758-5Report Type:Expedited (15-DaCompany Report #PT-GLAXOSMITHKLINE-B0321513A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hyperaesthesia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
		Rash Erythematous					
per day	35	DAY					
		Rash Pruritic		Motilium	C		ORAL
		Urticaria		Sedoxil	C		ORAL

Date:03/19/04ISR Number: 4319769-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0326019A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	1	MON					
		Pruritus					
		Renal Failure					

Date:03/19/04ISR Number: 4319772-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0326277A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erythema		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	20	DAY					
		Face Oedema		Naxy	C		ORAL
500MG Per day	5	DAY					
		Laryngeal Oedema					
		Tracheitis					

Date:03/19/04ISR Number: 4319779-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0327034A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alanine Aminotransferase		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	3	WK					

Increased  
Dermatitis Bullous  
Porphyria  
Serum Ferritin Increased

Date:03/22/04ISR Number: 4321105-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0442426A  
Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	51 DAY	Abnormal Behaviour		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged		Anxiety Chest Pain Circulatory Collapse Disorientation Heart Rate Increased Ill-Defined Disorder Stress Supraventricular Tachycardia		Glucosamine Nexium	C C		ORAL ORAL

Date:03/22/04ISR Number: 4321116-4Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0323573A  
Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Visual Acuity Reduced		Zyban	PS	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

2UNIT Three  
 times per day  
 Euphytose C ORAL

Date:03/22/04ISR Number: 4321123-1Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0326155A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blindness Transient		Zyban	PS	Glaxosmithkline	

Date:03/22/04ISR Number: 4323073-3Report Type:Expedited (15-DaCompany Report #002#4#2004-00046 (0)  
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Grand Mal Convulsion	Health Professional	Reglan-10mg-Tablet (Metoclopramide Hcl)	PS		ORAL
Required			Other	Bupropion	SS		ORAL
10MG, 1 IN 1				Valsartan-Hct	C		
Intervention to				Amlodipine	C		
D, ORAL				Simvastatin	C		
Prevent Permanent				Diflunisal	C		
ORAL				Doxepin	C		
Impairment/Damage				Levothyroxine	C		
				Omeprazole	C		
				Venlafaxine	C		
				Fluoxetine	C		
				Hydrochloroquine	C		
				Folic-Acid	C		

Date:03/23/04ISR Number: 4322080-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0443907A  
 Age:61 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 37.5MG Per Initial or Prolonged day	4 WK	Burning Sensation Erythema Hypersensitivity Nerve Injury Pruritus Pyrexia Stevens-Johnson Syndrome Swelling Face	Consumer	Lamictal	PS	Glaxosmithkline	ORAL
300MG Per day				Wellbutrin Xl	SS	Glaxosmithkline	
				Flomax	C		
				Detrol	C		
				Neurontin	C		
				Propranolol	C		

Date:03/23/04ISR Number: 4322081-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0503391A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Wellbutrin Sr	SS	Glaxosmithkline	ORAL
150MG Twice							
per day		YR					

Date:03/23/04ISR Number: 4322112-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0326881A  
 Age:32 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Anxiety
Initial or Prolonged	Loss Of Consciousness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Malaise Nightmare Tetany	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Tremor		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	8 DAY						

Date:03/23/04ISR Number: 4322389-4Report Type:Direct Company Report #CTU 215050  
Age:31 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Anaphylactic Reaction		Bupropion 300 Mg	PS		ORAL
Dose							
Hospitalization -		Hypersensitivity					
300 MG DAILY							
Initial or Prolonged							
ORAL							

Date:03/23/04ISR Number: 4322459-0Report Type:Direct Company Report #CTU 215033  
Age: Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Activities Of Daily Living Impaired		Welbutin			
Dose		Anorexia		Glaxo?	PS	Glaxo	
Life-Threatening		Anxiety		Zoloft	SS		
		Feeling Of Despair					
		Feelings Of Worthlessness					
		Insomnia					
		Paranoia					
		Suicidal Ideation					

Date:03/23/04ISR Number: 4322462-0Report Type:Direct Company Report #CTU 215036  
Age: Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Cerebrovascular Accident		Serzone	PS		ORAL
Dose							
Hospitalization -		Condition Aggravated					
600 MG DAILY							
Initial or Prolonged							
ORAL							

ORAL

Depression

Wellbutrin

SS

ORAL

Drug Effect Decreased  
Hypersensitivity  
Mental Disorder  
Suicidal Ideation

Date:03/24/04ISR Number: 4322728-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0498181A

Age:41 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 50TAB per day		Abortion Spontaneous	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged 37.5MG See		Complications Of Maternal	Professional	Paxil Cr	SS	Glaxosmithkline	ORAL
dosage text		Exposure To Therapeutic					
80MG Per day		Drugs		Geodon	C		
900MG Per day		Drug Exposure During		Neurontin	C		
1MG Per day	40 DAY	Pregnancy		Folate	C		
		Overdose Suicide Attempt		Prenatal Vitamin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/24/04ISR Number: 4322751-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0326452A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	ORAL
1TAB Twice		Anger					
per day	2	WK					
		Headache					

Date:03/24/04ISR Number: 4322757-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0326619A

Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysgeusia		Zyban	PS	Glaxosmithkline	ORAL
1TAB See		Tremor					
dosage text	10	DAY					

Date:03/24/04ISR Number: 4322760-0Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0042959A

Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain Upper	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Anaemia	Professional				
per day		Fat Intolerance					
		Gamma-Glutamyltransferase					
		Increased					
		Lipase Increased					

Date:03/24/04ISR Number: 4322973-8Report Type:Direct Company Report #CTU 215158

Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Gun Shot Wound		Paxil	PS		



Intentional Self-Injury

Wellbutrin

SS

Date:03/24/04ISR Number: 4323228-8Report Type:Direct  
Age:23 YR Gender:Female I/FU:I

Company Report #CTU 215194

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150 MG 1 A	Suicide Attempt		Wellbutrin	PS		ORAL
DAY ORAL, 300							
MG 1 A DAY							
ORAL							

Date:03/24/04ISR Number: 4323238-0Report Type:Direct  
Age:22 YR Gender:Female I/FU:I

Company Report #CTU 215201

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	1 DAY MORNIN	Fall		Wellbutrin 75 Mg	PS	Glaxo	ORAL
Hospitalization -	1 DAY MORNIN	Head Injury		Glaxo			
Initial or Prolonged	ORAL	Intentional Self-Injury					
ORAL							
1 DAY EVENIN							
ORAL							
Nervousness							
Suicide Attempt							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/24/04ISR Number: 4323270-7Report Type:Direct  
Age:37 YR Gender: I/FU:I

Company Report #CTU 215183

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Hypertension		Welbutrin	PS		ORAL
20 MG 1 DAILY						
Hospitalization -	Suicidal Ideation					
ORAL						
Initial or Prolonged			Celexa 20 Mg Fore	SS	Fore	
Required						
Intervention to						
Prevent Permanent						
Impairment/Damage						

Date:03/24/04ISR Number: 4323349-XReport Type:Direct  
Age:25 YR Gender:Male I/FU:I

Company Report #CTU 215179

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Depression		Wellbutrin Xl 300 Mg	PS		ORAL
1 TABLET PER						
ORAL	Impaired Work Ability					
	Memory Impairment					
	Thinking Abnormal					

Date:03/24/04ISR Number: 4323807-8Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 215213

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Aggression		Wellbutrin Sr 150 Mg			
150 MG BID	Alcoholism		Bid, Glaxo Smith K	PS	Glaxo Smith K	
	Impaired Driving Ability					
	Imprisonment					
	Poisoning					
	Suicide Attempt					

Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin Sr	PS	Glaxosmithkline	
400MG Per day							
		Drug Interaction		Clozaril	SS		ORAL
575MG Per day							
		Drug Level Decreased		Dilantin	SS		
		Grand Mal Convulsion		Clopixol	C		
		Therapeutic Response		Cogentin	C		
		Decreased		Fluoxetine	C		
				Lamictal	C	Glaxosmithkline	
				Lamotrigine	C	Glaxosmithkline	
				Lectopam	C		
				Olanzapine	C		
				Prozac	C		
				Seroquel	C		
				Zyprexa	C		

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Arrhythmia		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/25/04ISR Number: 4323385-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0503856A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:03/25/04ISR Number: 4323386-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0504045A

Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Alcohol Problem		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN		Completed Suicide		Xanax	C		
				Alcohol	C		

Date:03/25/04ISR Number: 4323387-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0504061A

Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG See							
Initial or Prolonged		Joint Dislocation					
dosage text		Medication Error		Nicorette Gum	C	Glaxosmithkline	
		Upper Limb Fracture		Diovan	C		
				Stelazine	C	Glaxosmithkline	

Date:03/25/04ISR Number: 4323390-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0504190A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Suicidal Ideation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Suicide Attempt					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Chest Pain	Health	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	2 WK	Dizziness	Professional				
UNKNOWN	100MG Per day	Dyspnoea		Astrix	C	Glaxosmithkline	
UNKNOWN	600MG	Nausea		Caltrate	C	Glaxosmithkline	
UNKNOWN	25MG Four	Retching		Cortisone Acetate	C	Glaxosmithkline	
times per day							
UNKNOWN	.5PCT Twice			Eleuphrat	C		
per day							
UNKNOWN	50MG Twice			Fenac	C	Glaxosmithkline	
per day							
UNKNOWN	20MG Twice			Losec	C		
per day							
UNKNOWN	400MCG As			Nitrolingual	C	Glaxosmithkline	
required							
UNKNOWN	.1MG Unknown			Oroxine	C	Glaxosmithkline	
UNKNOWN				Panadeine Forte	C	Glaxosmithkline	
UNKNOWN				Panamax	C	Glaxosmithkline	
per day							
UNKNOWN				Prednisone	C		
UNKNOWN	.625MG Per			Premarin	C		
day							
UNKNOWN	300MG At			Quinate	C	Glaxosmithkline	
night							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

RESPIRATORY ( INHALATION)		Seretide	C	Glaxosmithkline
UNKNOWN	18MCG Per day	Spiriva	C	
UNKNOWN	10MG See	Temaze	C	
dosage text		Ventolin	C	Glaxosmithkline
UNKNOWN				

Date:03/25/04ISR Number: 4324665-8Report Type:Direct Company Report #CTU 215279  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complications Of Maternal		Paxil 20 Mg	PS		ORAL
ONCE DAILY		Exposure To Therapeutic					
Required							
ORAL							
Intervention to		Drugs		Wellburtin Sr 150 Mg	SS		ORAL
TWICE DAILY							
Prevent Permanent		Depression					
ORAL							
Impairment/Damage		Feeling Of Despair Post-Traumatic Stress Disorder Pregnancy Self-Injurious Ideation					

Date:03/26/04ISR Number: 4324768-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441497A  
 Age:25 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour	Health	Paxil Cr	PS	Glaxosmithkline	ORAL
50MG Per day							
		Alcohol Poisoning	Professional	Contac	SS	Glaxosmithkline	ORAL
		Amnesia		Wellbutrin	SS	Glaxosmithkline	ORAL
		Anxiety		Paxil	SS	Glaxosmithkline	ORAL

1	DAY	Attention	Alcohol	SS	ORAL
		Deficit/Hyperactivity Disorder	Xanax	C	
		Confusional State	Clonopin	C	
		Depression	Seroquel	C	
		Diarrhoea	Atenolol	C	
		Disturbance In Attention			
		Drug Interaction			
		Dysphoria			
		Fatigue			
		Headache			
		Initial Insomnia			
		Laceration			
		Mental Status Changes			
		Mydriasis			
		Nausea			
		Obsessive-Compulsive Disorder			
		Panic Attack			
		Psychotic Disorder			
		Toxicologic Test Abnormal			

Date:03/26/04ISR Number: 4324774-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0498860A  
Age:18 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
		Medication Error					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/26/04ISR Number: 4324780-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0503547A  
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300MG Per day		Choking		Wellbutrin	PS	Glaxosmithkline	ORAL
		Convulsion		Seroquel	C		
1 YR		Depressed Level Of Consciousness Disorientation					

Date:03/26/04ISR Number: 4324787-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0504270A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
2 YR		Suicidal Ideation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Suicide Attempt					
Other							

Date:03/26/04ISR Number: 4324788-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0504316A  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
2 YR		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Diabetes Mellitus		Wellbutrin	SS	Glaxosmithkline	
150MG Per day 1 MON		Inadequate Control		Reglan	C	Glaxosmithkline	
		Insomnia		Trental	C		
		Medication Error		Alpha Lipoic Acid	C		
				Chromium Picolinate	C		
				Carbamazepine	C		
				Evening Primrose Oil	C		
				Insulin	C		

Date:03/26/04ISR Number: 4324813-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0327761A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK	Haematoma Myalgia Oedema Peripheral Serum Sickness					

Date:03/26/04ISR Number: 4325033-5Report Type:Direct Company Report #CTU 215375  
Age:29 YR Gender:Female I/FU:I

Outcome	PT
Life-Threatening	Affect Lability
Hospitalization -	Amnesia
Initial or Prolonged	Antisocial Behaviour
Required	Confusional State
Intervention to	Conversion Disorder
Prevent Permanent	Hostility
Impairment/Damage	Migraine
	Myalgia
	Paranoia
	Suicidal Ideation
	Suicide Attempt
	Vertigo

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Weight Increased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1 ORAL			Prozac	PS		ORAL
2-3 ORAL			Zoloft	SS		ORAL
			Paxil	SS		
			Wellbutrin	SS		

Date:03/29/04ISR Number: 4325955-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0494894A  
 Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Electrocardiogram Qt	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice		Prolonged	Professional				
per day	6	MON		Neurontin	C		ORAL
1800MG per							
day				Strattera	C		ORAL
64 DAY				Zyprexa	C		ORAL
10MG At night				Trazodone	C		

Date:03/29/04ISR Number: 4325956-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0495776A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
5 WK		Somnolence	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
5 WK				Trazodone	C		

Date:03/29/04ISR Number: 4325957-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0499051A

Age:6 MON Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Exposure Via Breast	Health	Bupropion	PS	Glaxosmithkline	
150MG Per day		Milk	Professional				
		Grand Mal Convulsion					

Date:03/29/04ISR Number: 4325960-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0503470A

Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Exposure During		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Pregnancy		Antibiotic	C		
		Fungal Infection					
		Nephrolithiasis					
		Urinary Tract Infection					

Date:03/29/04ISR Number: 4325961-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0503536A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
4 DAY							
Initial or Prolonged		Flight Of Ideas					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/04ISR Number: 4325969-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0504565A

Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
13 MON		Headache Suicidal Ideation					

Date:03/29/04ISR Number: 4325973-7Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0312414A

Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Arrhythmia	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG See							
Hospitalization - dosage text		Bradycardia	Professional				
Initial or Prolonged Disability		Cardiac Enzymes Increased		Ascorbic Acid	C	Glaxosmithkline	
Other		Epilepsy		Aspirin Junior	C	Glaxosmithkline	
		Influenza Like Illness		Multivitamin With			
		Loss Of Consciousness		Zinc	C		
		Malaise		Selenium	C		
		Sudden Death		Glucosamine Sulphate	C		
		Syncope		Ventolin	C	Glaxosmithkline	
RESPIRATORY							
(INHALATION)		Ventricular Extrasystoles					
		Ventricular Fibrillation		Lysomucil	C	Glaxosmithkline	ORAL
		White Blood Cell Count Increased					

Date:03/29/04ISR Number: 4325987-7Report Type:Expedited (15-DaCompany Report #FI-GLAXOSMITHKLINE-B0327048A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG Twice		Dyspnoea		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	17 DAY	Eczema					
		Oedema Mucosal					

Pruritus  
Urticaria  
Visual Disturbance

Date:03/29/04ISR Number: 4325993-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0327617A  
Age:54 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 150MG See Hospitalization - dosage text 35 DAY Initial or Prolonged UNKNOWN	Suicide Attempt		Zyban	PS	Glaxosmithkline	ORAL
UNKNOWN			Fenofibrate	C		
UNKNOWN			Maxepa	C	Glaxosmithkline	
UNKNOWN			Kardegic	C		
UNKNOWN			Titanoreine	C		

Date:03/29/04ISR Number: 4325994-4Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0327623A  
Age:40 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Per day Initial or Prolonged	Abdominal Pain Oligomenorrhoea		Zyban	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/04ISR Number: 4326341-4Report Type:Direct  
Age:35 YR Gender:Female I/FU:I

Company Report #CTU 215449

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage	Bipolar I Disorder Suicide Attempt		Wellbutrin Glaxo	PS	Glaxo	

Date:03/29/04ISR Number: 4326342-6Report Type:Direct  
Age:41 YR Gender:Female I/FU:I

Company Report #CTU 215451

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening ORAL	Convulsion		Wellbutrin	PS		ORAL

Date:03/29/04ISR Number: 4330951-8Report Type:Expedited (15-DaCompany Report #B0325227A  
Age:33 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Required ORAL Intervention to Prevent Permanent Impairment/Damage	Blood Creatine Phosphokinase Increased Gliosis Grand Mal Convulsion Hepatic Enzyme Increased Hyperreflexia Inflammation Spinal Compression Fracture Tachycardia	Foreign Literature Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:03/29/04ISR Number: 4332258-1Report Type:Expedited (15-DaCompany Report #S04-USA-01540-01  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Petit Mal Epilepsy Tremor	Consumer	Lexapro (Escitalopram)	PS		ORAL
	10 MG QD PO			Wellbutrin - Slow Release (Bupropion Hydrochloride)	SS		
	150 MG QD			Prozac (Fluoxetine Hydrochloride)	C		

Date:03/29/04ISR Number: 4332262-3Report Type:Expedited (15-DaCompany Report #S04-USA-01545-01  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Amnesia Grand Mal Convulsion	Health Professional	Tessalon Perle (Benzonatate)	PS		ORAL
	100 MG BID PO			Wellbutrin - Slow Release (Bupropion Hydrochloride)	SS		
	200 MG BID			Wellbutrin - Slow Release (Bupropion Hydrochloride)	SS		
	300 MG QD						



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/04ISR Number: 4327145-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0314874A

Age:61 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG See Initial or Prolonged dosage text 38 DAY	Tendon Rupture	Health  Professional	Zyban	PS	Glaxosmithkline	ORAL
			Avandia Metformine	C C	Glaxosmithkline	ORAL ORAL
850MG Twice  per day  2UNIT per day			Novonorm	C		ORAL

Date:03/30/04ISR Number: 4327147-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0318887A

Age:37 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Per day 12 DAY Initial or Prolonged	Diplopia Headache Iiird Nerve Paralysis Strabismus	Health  Professional	Zyban	PS	Glaxosmithkline	ORAL

Date:03/30/04ISR Number: 4327148-4Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0320004A

Age:33 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Unknown 22 DAY	Anxiety Aptyalism Asthenia Cytolytic Hepatitis Cytomegalovirus Antibody Positive Dermatitis Exfoliative Epstein-Barr Virus Antibody Positive	Health  Professional	Zyban	PS	Glaxosmithkline	ORAL

Hypersensitivity  
Hypotension  
Inflammation  
Insomnia  
Leukopenia  
Lymphadenopathy  
Nausea  
Painful Respiration  
Pruritus  
Purpura  
Rash Erythematous  
Rash Maculo-Papular  
Rash Morbilliform  
Rash Scarlatiniform  
Thrombocytopenia  
Tremor

Date:03/30/04ISR Number: 4327155-1Report Type:Expedited (15-DaCompany Report #FI-GLAXOSMITHKLINE-B0326891A  
Age:32 YR Gender:Female I/FU:I

Outcome PT  
Hospitalization - Aggression  
Initial or Prolonged Agitation  
Confusional State  
Headache  
Hypertension

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Hyperventilation Sinus Tachycardia				
Dose	Duration		Report Source	Product	Role	Manufacturer
				Zyban	PS	Glaxosmithkline
10	DAY					ORAL

Date:03/30/04ISR Number: 4327159-9Report Type:Expedited (15-DaCompany Report #FI-GLAXOSMITHKLINE-B0327209A  
Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Feeling Abnormal		Zyban	PS	Glaxosmithkline	ORAL
150MG See		Hypoaesthesia					
dosage text		Insomnia		Formoterol	C		
6MCG Per day		Irritability		Budesonide	C		
400MCG Twice		Muscular Weakness					
per day		Paraesthesia		Ipratropium			
		Peroneal Nerve Palsy		Salbutamol	C		

Date:03/30/04ISR Number: 4327538-XReport Type:Direct Company Report #CTU 215519  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Agitation		Lexipro Police			
ONCE A DAY		Anxiety		Confiscated	PS		ORAL
ORAL		Completed Suicide					
ONCE A DAY		Intentional Self-Injury		Welbutrin Police			
ORAL				Confiscated	SS		ORAL
				Ambien	C		
				Blood Pressure			
				Medication	C		

Trazidone C  
 Maxalt C  
 Flonase C

Date:03/30/04ISR Number: 4332054-5Report Type:Expedited (15-DaCompany Report #B0327440A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cardiac Failure Dyspnoea Tachycardia	Foreign Other	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG/ORAL	1 WK						

Date:03/31/04ISR Number: 4327686-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0420530A  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
5MG Four times per day	1 YR	Dysphoria Tinnitus		Compazine	PS	Glaxosmithkline	ORAL
WK				Wellbutrin	SS	Glaxosmithkline	ORAL
				Vicodin	C		ORAL
				Lexapro	C		
				Unspecified Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/31/04ISR Number: 4328185-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0504801A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia		Paxil	PS	Glaxosmithkline	ORAL
20MG Per day	4 YR	Blood Pressure Increased		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Insomnia		No Concurrent			
		Irritability		Medication	C		
		Pain					
		Paraesthesia					
		Suicidal Ideation					
		Tinnitus					
		Vision Blurred					
		Weight Increased					

Date:03/31/04ISR Number: 4331004-5Report Type:Expedited (15-DaCompany Report #2004005972  
Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Health	Zoloft (Sertraline)	PS		ORAL
50 MG		Benign Bone Neoplasm	Professional				
(DAILY), ORAL		Bruxism		Bupropion			
		Drug Withdrawal Syndrome		Hydrochloride			
		Feeling Abnormal		(Bupropion			
		Hot Flush		Hydrochloride)	SS		
300 MG (BID)		Hyperphagia		Levothyroxine Sodium			
		Paraesthesia		(Levothyroxine			
		Pharmaceutical Product		Sodium)	C		
		Complaint		Clonazepam			
		Sleep Disorder		(Clonazepam)	C		
		Tongue Ulceration		Vitamins	C		
		Weight Increased					

Date:04/01/04ISR Number: 4329728-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0422196A  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Back Pain					
per day		Eye Pain		Clonazepam	C		
		Headache					
		Thinking Abnormal					
		Visual Acuity Reduced					

Date:04/01/04ISR Number: 4329729-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0439007A  
Age:78 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Balance Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Twice		Chills					
Initial or Prolonged		Depression		Flomax	C		
per day	1 MON	Erectile Dysfunction		Cozaar	C		
Other		Insomnia		Indapamide	C		
		Myalgia		Duragesic	C		
		Myoclonus					
		Tremor					
		Vestibular Disorder					



Date:04/01/04ISR Number: 4329741-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0505251A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG per day							

Date:04/01/04ISR Number: 4329744-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0505433A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Medication Error		Wellbutrin Psychotropic Medications	PS C	Glaxosmithkline	ORAL

Date:04/01/04ISR Number: 4330257-7Report Type:Direct Company Report #CTU 215809  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression Impaired Work Ability		Wellbutrin (Bupropion)	PS		ORAL
150 MG BID PO							
Paraesthesia							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/01/04ISR Number: 4330724-6Report Type:Direct  
Age:50 YR Gender:Male I/FU:I

Company Report #CTU 215660

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - INTRAVENOUS Initial or Prolonged DAILY	100MG IV	Dizziness		Venofer Iv 100mg	PS		
		Hypoaesthesia Mental Impairment Speech Disorder		Wellbutrin	SS		

Date:04/01/04ISR Number: 4331683-2Report Type:Direct  
Age:38 YR Gender:Male I/FU:I

Company Report #CTU 215803

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ONE DOSE PO QD		Rash Macular Rash Pruritic		Wellbutrin Xl 150 Mg	PS		ORAL

Date:04/02/04ISR Number: 4330638-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0443385A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 400MG per day		Dystonia	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
		Fatigue Grand Mal Convulsion Hypoaesthesia Impaired Driving Ability Loss Of Consciousness Paraesthesia Tongue Biting	Professional				

Date:04/02/04ISR Number: 4330639-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0495363A  
Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice		Drug Interaction	Professional				
per day	4	MON		Prednisolone	SS	Glaxosmithkline	

Date:04/02/04ISR Number: 4330651-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0505440A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN		Drug Interaction		Klonopin	SS		

Date:04/02/04ISR Number: 4330666-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0328013A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dry Mouth		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Hypersensitivity					
		Insomnia					
		Sleep Disorder					
		Suffocation Feeling					
		Throat Tightness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/02/04ISR Number: 4330669-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0328536A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Unknown	15 DAY		Antinuclear Antibody	Zyban	PS	Glaxosmithkline	ORAL
UNKNOWN			Positive	Foncitril 4000	C		
UNKNOWN			Arthralgia	Orocal	C	Glaxosmithkline	
UNKNOWN			Drug Toxicity	Sulfarlem	C		
UNKNOWN			Epidermal Necrosis	Imovane	C		
UNKNOWN			Inflammation	Aspirin	C	Glaxosmithkline	
			Photosensitivity Reaction				
			Rash Erythematous				
			Systemic Lupus				
			Erythematosis				
			Toxic Skin Eruption				

Date:04/02/04ISR Number: 4330670-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0328580A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
Initial or Prolonged			Abnormal Behaviour	Zyban	PS	Glaxosmithkline	ORAL
500MG Twice			Confusional State	Ciflox 500	SS		ORAL
per day	5 DAY		Delirium Tremens				
500MG Three			Hyponatraemia	Vitamine B1	C		ORAL
times per day			Pyelonephritis				
			Renal Failure Acute				

Date:04/02/04ISR Number: 4330675-7Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0043506A  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Suicide Attempt Zyban PS Glaxosmithkline ORAL

44TAB Single

dose

Date:04/05/04ISR Number: 4332094-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0421244A

Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction		Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Twice		Transient Ischaemic					
per day		Attack		Nortriptyline	SS		
				Ibuprofen	SS	Glaxosmithkline	

Date:04/06/04ISR Number: 4333832-9Report Type:Direct Company Report #CTU 216002

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Accidental Overdose		Prozac	PS		
Hospitalization -		Anger		Paxil	SS		
Initial or Prolonged		Anorexia		Zoloft	SS		
Disability		Cognitive Disorder		Effexor	SS		
Required		Depression		Wellbutrin	SS		
Intervention to		Fibromyalgia		Sertraline	SS		
Prevent Permanent		Suicide Attempt					
Impairment/Damage							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/06/04ISR Number: 4335551-1Report Type:Expedited (15-DaCompany Report #CHPA2004US01801  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anger	Consumer	Riteaid Step 1 21mg (Nch)(Nicotine)			
		Drug Interaction		Trans-Therapeutic-Sy stem	PS		
		Personality Change					
		Psychotic Disorder					
TRANSDERMAL	21 MG, QD,	QD,					
TRANSDERMAL		Social Avoidant Behaviour					
				Wellbutrin(Bupropion Hydrochloride)	SS		
				Thiothixene (Tiotixene)	C		
				Benztropeine	C		
	150 MG, QD,						

Date:04/07/04ISR Number: 4334519-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0327924A  
Age:22 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Overdose		Zyban	PS	Glaxosmithkline	ORAL
1500MG per							
Initial or Prolonged		Suicide Attempt					
day	1 DAY			Aleve	SS		ORAL
1320MG per							
day	1 DAY						

Date:04/07/04ISR Number: 4334524-2Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0328019A  
Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dyspnoea		Zyban	PS	Glaxosmithkline	
Initial or Prolonged				Clopidogrel	C		
75MG per day							
				Bisoprolol Fumarat	C		
2.5MG per day							

4MG per day

Perindopril  
Tert-Butylamine C

160MG per day

Acetylsalicylic Acid C Glaxosmithkline

Date:04/07/04ISR Number: 4334551-5Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0328585A

Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Supraventricular		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown		Tachycardia					

Date:04/07/04ISR Number: 4334755-1Report Type:Direct

Company Report #CTU 216162

Age: Gender:Female I/FU:I

Outcome	PT
Life-Threatening	Anhedonia
	Anxiety
	Cardiac Flutter
	Crying
	Dizziness
	Drug Ineffective
	Electroencephalogram
	Abnormal
	Fatigue
	Feeling Abnormal
	Feeling Jittery
	Head Injury

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route

Date:04/08/04ISR Number: 4335029-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0504190A  
 Age: Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route

Date:04/08/04ISR Number: 4335030-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0505092A  
 Age:26 YR Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route

Date:04/08/04ISR Number: 4335034-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0505954A  
 Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Fall Haemorrhage Head Injury Skull Fracture					

Date:04/08/04ISR Number: 4335035-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0505967A  
 Age:15 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
300MG See		Medication Error	Professional				
dosage text	3 WK						

Date:04/08/04ISR Number: 4336941-3Report Type:Expedited (15-DaCompany Report #2004UW06035  
 Age:46 YR Gender:Male I/FU:I

Outcome  
 Required  
 Intervention to

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Prevent Permanent  
Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abasia	Consumer	Seroquel	PS		
		Dysstasia		Wellbutrin	SS		
		Fall		Motrin	C		
		Rhabdomyolysis		Heparin	C		
				Ultram	C		
				Prednisone	C		
				Cellcept	C		
				Sleeping Pill	C		

Date:04/09/04ISR Number: 4335454-2Report Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-12493342  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complications Of Maternal Exposure To Therapeutic Drugs		Abilify	PS	Otsuka Pharmaceutical Company, Ltd.	ORAL
12.5 to 25		Intra-Uterine Death		Seroquel	SS		
mg/day as		Maternal Drugs Affecting Foetus					
needed		Pregnancy		Klonopin	SS		
				Lexapro	SS		
				Wellbutrin	SS		

Date:04/09/04ISR Number: 4335677-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0505989A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:04/09/04ISR Number: 4335678-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506077A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin Sr	PS	Glaxosmithkline	
150MG per day							

Date:04/09/04ISR Number: 4335679-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506101A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
175MG See							
dosage text							

Date:04/09/04ISR Number: 4335681-4Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0506123A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Exposure During		Bupropion	PS	Glaxosmithkline	
150MG per day							
		Pregnancy					
		Foetal Heart Rate					
		Abnormal					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/09/04ISR Number: 4335700-5Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0326452A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Affective Disorder	Health	Zyban	PS	Glaxosmithkline	ORAL
1TAB Twice		Aggression	Professional				
per day	2	WK					
		Anger		Hydroxychloroquine			
		Headache		Sulphate	C		ORAL
200MG Twice							
per day							

Date:04/09/04ISR Number: 4335701-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0327623A

Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	8	DAY					
		Abdominal Pain Lower	Professional	Mercilon	C		ORAL
		Back Pain		Revaxis	C		
UNKNOWN		1 DAY					
		Oligomenorrhoea					
		Paraesthesia					

Date:04/09/04ISR Number: 4335702-9Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0328342A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cerebrovascular Accident		Zyban	PS	Glaxosmithkline	ORAL
150MG Three							
times per day							

Date:04/09/04ISR Number: 4335707-8Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0328702A

Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG per day 4 WK Initial or Prolonged		Cerebral Infarction		Zyban	PS	Glaxosmithkline	
		Diplopia		Logimax	C		
		Dysarthria		Thyroxin Sodium	C	Glaxosmithkline	
		Headache		Metformin			
		Hemiparesis		Hydrochloride	C		
				Rosiglitazone			
				Maleate	C	Glaxosmithkline	

Date:04/09/04ISR Number: 4335708-XReport Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0328825A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Diabetic Neuropathy	Health Professional	Zyntabac Unspecified Anti-Diabetic	PS  C	Glaxosmithkline	ORAL

Date:04/09/04ISR Number: 4342734-3Report Type:Periodic Company Report #03P-163-0213874-00  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage PER ORAL		Convulsion	Health Professional Company Representative	Semisodium Valproate (Depakote) (Divalproex Sodium) (Divalproex Sodium)	PS		ORAL
PER ORAL			Other	Bupropion Hydrochloride	SS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/12/04ISR Number: 4336661-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0505546A

Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300MG Per day	3	MON	Abortion Spontaneous	Wellbutrin	PS	Glaxosmithkline	ORAL
			Drug Exposure During Pregnancy	Ritalin	C		
			Weight Increased				

Date:04/12/04ISR Number: 4336663-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506244A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Hypersensitivity Medication Error	Wellbutrin	PS	Glaxosmithkline	
				Wellbutrin	SS	Glaxosmithkline	ORAL

Date:04/12/04ISR Number: 4336665-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506288A

Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
			Rash	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Vasculitis				

Date:04/12/04ISR Number: 4336666-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506304A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged			Amnesia	Wellbutrin	PS	Glaxosmithkline	ORAL
			Chest Pain	Alcohol	SS		ORAL
			Depressed Level Of Consciousness				
			Suicide Attempt				

Date:04/12/04ISR Number: 4336673-1Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0328618A

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Duration Cerebrovascular Accident Monoparesis		Zyban	PS	Glaxosmithkline	

Date:04/12/04ISR Number: 4338131-7Report Type:Direct

Company Report #CTU 216425

Age:49 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG X 2DAILY ORAL	Duration Pruritus Skin Lesion		Wellbutrin	PS		ORAL

Date:04/13/04ISR Number: 4337523-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506364A

Age: Gender: I/FU:F

Outcome  
Death  
Life-Threatening  
Hospitalization -  
Initial or Prolonged  
Disability

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Other

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN			Overdose		Wellbutrin	PS	Glaxosmithkline	
			Respiratory Failure		Prozac	SS		

Date:04/13/04ISR Number: 4337524-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506389A  
 Age: Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:04/13/04ISR Number: 4337526-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506474A  
 Age: Gender: I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Convulsion Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:04/13/04ISR Number: 4337528-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506493A  
 Age: Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:04/13/04ISR Number: 4338091-9Report Type:Direct Company Report #CTU 216521  
 Age:11 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100 MG ONE BID PO			Abnormal Behaviour Aggression		Bupropion Sr 100 Mg	PS		ORAL

Pharmaceutical Product  
Complaint

Date:04/13/04ISR Number: 4338297-9Report Type:Direct  
Age:46 YR Gender:Female I/FU:I

Company Report #CTU 216488

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG BID	Hypersensitivity		Wellbutrin Sr 150mr	PS		ORAL
ORAL		Urticaria					

Date:04/14/04ISR Number: 4338487-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506365A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day 5 WK	Disturbance In Attention		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day		Dizziness		Calcium + Magnesium			
		Muscle Twitching		Supplement	C		
		Nephrolithiasis		Zoloft	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/14/04ISR Number: 4338489-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506833A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	5 DAY	Insomnia					
		Medication Error		Toprol	C		
		Nausea					

Date:04/14/04ISR Number: 4338491-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506950A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Medication Error		Wellbutrin	PS	Glaxosmithkline	
		Pharmaceutical Product		Bupropion			
		Complaint		Hydrochloride	SS	Glaxosmithkline	ORAL

Date:04/14/04ISR Number: 4338506-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0329071A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Anger		Zyban	PS	Glaxosmithkline	ORAL
		Psychotic Disorder		Quetiapine	C		
		Suicide Attempt					

Date:04/15/04ISR Number: 4339432-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506081B

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Drug Exposure During		Bupropion	PS	Glaxosmithkline	
150MG per day	78 DAY	Pregnancy					
		Sudden Infant Death Syndrome					

Date:04/15/04ISR Number: 4339439-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0507157A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	
150MG Twice							
per day							

Date:04/15/04ISR Number: 4339440-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0507194A

Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							
		Overdose		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:04/16/04ISR Number: 4339924-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0503515A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Arrhythmia		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/16/04ISR Number: 4339943-6Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0328825A

Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	58 DAY		Cerebral Atrophy	Zyntabac	PS	Glaxosmithkline	ORAL
4MG Per day			Difficulty In Walking	Avandia	SS	Glaxosmithkline	ORAL
1700MG per day			Dizziness	Dianben	SS		ORAL
			Polyneuropathy				
			Pyrexia	Adiro	SS	Glaxosmithkline	ORAL

Date:04/16/04ISR Number: 4339947-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0329259A

Age:36 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Unknown	2 MON	Affect Lability	Zyban	PS	Glaxosmithkline	ORAL
INTRAMUSCULAR	150MG per day		Anxiety	Depo-Provera	C		
			Crying				
			Drug Withdrawal Syndrome				
			Psychiatric Symptom				

Date:04/16/04ISR Number: 4339948-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0329261A

Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	26 DAY		Cheilitis	Zyban	PS	Glaxosmithkline	ORAL
150MG per day	1 MON		Dysphagia	Ranitidine	C	Glaxosmithkline	ORAL
			Glossodynia				
			Oral Pain				

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Drug Interaction Hyperglycaemia	Wellbutrin Hiv Treatments (Unspecified)	PS C	Glaxosmithkline	

Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Unknown 4 WK			Anti-Thyroid Antibody Positive Arthralgia Blood Cortisol Increased Chromosomal Mutation Cytolytic Hepatitis Dermatitis Bullous Hypothyroidism Porphyria Non-Acute Porphyrinuria Scar Serum Ferritin Increased	Zyban	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/19/04ISR Number: 4341156-9Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0329055A  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	15 DAY	Balance Disorder		Zyban	PS	Glaxosmithkline	ORAL
40MG Per day		Feeling Abnormal Paraesthesia Speech Disorder Vertigo Visual Acuity Reduced		Pravachol	C		

Date:04/19/04ISR Number: 4341163-6Report Type:Expedited (15-DaCompany Report #TR-GLAXOSMITHKLINE-B0329470A  
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly 13 DAY		Abortion Spontaneous		Zyban	PS	Glaxosmithkline	ORAL
		Drug Exposure During Pregnancy Pregnancy		No Concurrent Medications	C		

Date:04/19/04ISR Number: 4341545-2Report Type:Direct Company Report #CTU 216878  
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1 TABLET DAILY		Balance Disorder Confusional State Depression Fatigue Feeling Abnormal Hangover Headache Heart Rate Increased Hypersomnia Hypokinesia		Bupropion 150 Watson	PS	Watson	

Lethargy  
Pharmaceutical Product  
Complaint  
Sedation

Date:04/20/04ISR Number: 4341794-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0495331A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	75 DAY	Fatigue Muscle Contractions Involuntary Partial Seizures	Health Professional	Wellbutrin	PS	Glaxosmithkline	ORAL
.5MG At night 10MG Per day		Tremor		Paxil Ativan Zyrtec	SS C C	Glaxosmithkline Glaxosmithkline	ORAL ORAL ORAL

Date:04/20/04ISR Number: 4341796-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0507470A  
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 15 WK		Chest Pain Medication Error Nervousness Tremor		Wellbutrin Xl Zantac One A Day Vitamins	PS C C	Glaxosmithkline Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/20/04ISR Number: 4341808-0Report Type:Expedited (15-DaCompany Report #FI-GLAXOSMITHKLINE-B0329553A  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Postmenopausal		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Haemorrhage					
per day				Symbicort Turbohaler	C		
RESPIRATORY							
(INHALATION)	1U Twice per						
day							

Date:04/21/04ISR Number: 4343107-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506371A  
 Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 4	WK						
Initial or Prolonged		Medication Error		Gabitril	C		
Other							

Date:04/21/04ISR Number: 4343111-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506950A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error	Consumer	Wellbutrin	PS	Glaxosmithkline	
		Pharmaceutical Product		Bupropion			
		Complaint		Hydrochloride	SS	Glaxosmithkline	ORAL
150MG Per day							

Date:04/21/04ISR Number: 4343122-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0328013A  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 1TAB Twice Initial or Prolonged per day	8 DAY	Dry Mouth Hypersensitivity Oropharyngeal Swelling Pharyngolaryngeal Pain Sleep Disorder Suffocation Feeling Throat Tightness	Health Professional	Zyban	PS	Glaxosmithkline	ORAL
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Date:04/21/04ISR Number: 4344410-XReport Type:Direct Company Report #CTU 217029  
 Age:43 YR Gender:Female I/FU:I

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 1-3X DAILY	Hospitalisation		Wellbutrin	PS		
Hospitalization - 1 X DAILY			Prozac 20 Mg	SS		
Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage			Wellbutrin	C		

Date:04/22/04ISR Number: 4344063-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0507922A  
 Age:27 YR Gender:Female I/FU:I

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice	Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	Medication Error					
6 DAY Other			Oral Contraceptives	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/23/04ISR Number: 4345298-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506106A  
 Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Feeling Abnormal		Wellbutrin	SS	Glaxosmithkline	
150MG Per day		Libido Disorder		Paxil Cr	SS	Glaxosmithkline	
12.5MG See		Medication Error					
dosage text		Nausea		Xanax	SS		
		Smoker					

Date:04/23/04ISR Number: 4345300-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508064A  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG See		Pharyngolaryngeal Pain					
dosage text		Vertigo					

Date:04/23/04ISR Number: 4345895-5Report Type:Direct Company Report #CTU 217245  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying		Wellbutrin Sr 150 Mg			
Life-Threatening		Fear		Glaxo Smithkline	PS	Glaxo Smithkline	ORAL
1 DAILY ORAL		Suicidal Ideation					

Date:04/23/04ISR Number: 4346282-6Report Type:Direct Company Report #CTU 217296  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Life-Threatening	Arrhythmia	Wellbutrin	PS	ORAL
150 MG/ PO				
Hospitalization -	Chest Pain			
Initial or Prolonged	Dizziness			
	Dyspnoea			
	Heart Rate Increased			
	Pallor			
	Supraventricular			
	Tachycardia			

Date:04/23/04ISR Number: 4346342-XReport Type:Direct Company Report #CTU 217292  
 Age:33 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Chest Pain		Nicotine Patch 21mg			
Initial or Prolonged	Dizziness		Tdrm	PS		
TRANSDERMAL	1 PATCH QD					
	Dyspnoea		Bupropion Sr (Zyban)			
	Feeling Abnormal		150 Mg	SS		ORAL
1 TAB PO QD						
	Nausea					
	Palpitations					
	Vomiting					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/23/04ISR Number: 4348558-5Report Type:Expedited (15-DaCompany Report #USA-2004-0014626  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arteriosclerosis	Health	Oxycodone			
		Chronic Obstructive	Professional	Hydrochloride			
		Pulmonary Disease	Other	(Similar To Nda			
		Coronary Artery Occlusion		20-553)(Oxycodone			
		Drug Screen Positive		Hydrochloride)	PS		
		Granuloma		Hydrocodone			
		Hepatic Cirrhosis		Bitartrate (Similar			
		Hepatitis C		To Ind			
		Pleural Adhesion		59,175)(Hydrocodone			
		Tuberculosis		Bitartrate) Unknown	SS		
				Ethanol (Ethanol)	SS		
				Acetaminophen			
				(Paracetamol)	SS		
				Alprazolam			
				(Alprazolam)	SS		
				Bupropion			
				(Amfebutamone)	SS		
				Doxepin (Doxepin)	SS		
				Salicylic Acid			
				(Salicylic Acid)	SS		
				Citalopram			
				(Citalopram)	SS		
				Dextromethorphan			
				(Dextromethorphan)	SS		
				Lorazepam			
				(Lorazepam)	SS		
				Paroxetine			
				(Paroxetine)	SS		
				Temazepam			
				(Temazepam)	SS		
				Oxazepam (Oxazepam)	SS		

Date:04/26/04ISR Number: 4347073-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0314643A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion		Zyban	PS	Glaxosmithkline	ORAL
		Drug Level Increased		Librax	C		ORAL
		Overdose		Levothyrox	C	Glaxosmithkline	ORAL

Sudden Death

Catapressan  
Agsreal

C  
C

ORAL

Date:04/26/04ISR Number: 4347323-2Report Type:Direct  
Age:16 YR Gender:Female I/FU:I

Company Report #CTU 217372

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG TWICE		Convulsion Depression		Wellbutrin Sr 200 Mg Generic			ORAL
A DA ORAL		Pharmaceutical Product Complaint			PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/26/04ISR Number: 4347327-XReport Type:Direct  
Age:13 YR Gender:Male I/FU:I

Company Report #CTU 217373

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100 MG DAILY ORAL		Abnormal Behaviour Pharmaceutical Product  Complaint  Therapeutic Response Decreased		Wellbutrin Sr 100 Mg Gsm Non-Generic	PS	Gsm Non-Generic	ORAL

Date:04/26/04ISR Number: 4347331-1Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 217375

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 35 MG EVERY Initial or Prolonged NIGH ORAL Disability 300 MG 3X A DAY ORAL		Agitation  Anxiety  Depression  General Physical Health  Deterioration Hostility Hypomania Impulsive Behaviour Insomnia Irritability Panic Attack Suicide Attempt		Remeron 35 Mg   Wellbutrin	PS   SS		ORAL   ORAL

Date:04/26/04ISR Number: 4347463-8Report Type:Direct  
Age:50 YR Gender:Male I/FU:I

Company Report #CTU 217352

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose  2 A DAY		Aggression Dyskinesia		Zoloft 100 Mg Tablet Roe	PS	Roe	

3 ADAY

Wellbutrin Sr 150 Mg  
Tab Sa Gsk SS Gsk

Date:04/26/04ISR Number: 4349010-3Report Type:Direct Company Report #CTU 217384  
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression		Wellbutrin Sr			
100 MG 2 PO		Pharmaceutical Product		(Generic)	PS		ORAL
		Complaint					
BID							

Trazadone C  
Xanax C

Date:04/27/04ISR Number: 4348316-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0504299A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Laryngeal Oedema		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG In the							
Other		Rash					
morning	2 WK						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/27/04ISR Number: 4348321-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508351A  
Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
4500MG Single		Overdose					
dose				No Concurrent Medication	C		

Date:04/27/04ISR Number: 4348326-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508440A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Twice		Headache					
per day		Medication Error		No Concurrent Medication	C		

Date:04/27/04ISR Number: 4348343-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0329938A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Zyban	PS	Glaxosmithkline	
150MG See		Foetal Growth Retardation					
dosage text	1 WK			Salbutamol	C	Glaxosmithkline	
				Becotide	C	Glaxosmithkline	

Date:04/27/04ISR Number: 4348346-XReport Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0330195A  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death 150MG Twice Other per day	Myocardial Infarction	Zyban	PS	Glaxosmithkline	ORAL
20TABS per day	Overdose	Paroxetine	SS	Glaxosmithkline	ORAL

Date:04/27/04ISR Number: 4348970-4Report Type:Direct Company Report #CTU 217487  
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Feeling Hot		Wellbutrin 150 Mg Xl	PS		ORAL
1/4 TAB PO		Renal Pain					

Date:04/27/04ISR Number: 4349130-3Report Type:Direct Company Report #CTU 217441  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Drug Ineffective		Wellbutrin Xl 150 Mcg Glaxosmithkline	PS		ORAL
300 MCG DAILY		Erythema					
ORAL		Pruritus Self-Medication Urticaria		Levothyroxine	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/27/04ISR Number: 4350573-2Report Type:Expedited (15-DaCompany Report #2004-01566

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Myasthenia Gravis	Health Professional Other	Trazodone Hydrochloride (Watson Laboratories) (Trazodone Wellbutrin (Bupropion Hydrochloride)	PS  SS	Watson Laboratories	

Date:04/28/04ISR Number: 4349387-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508423A

Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG Twice				Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	8 DAY	Dyspnoea					
Disability		Medication Error					
Other		Rash					

Date:04/28/04ISR Number: 4349390-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508692A

Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Affect Lability		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							
Other		Crying Depressed Mood		No Concurrent Medications	C		

Date:04/28/04ISR Number: 4349877-9Report Type:Direct Company Report #CTU 217597

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Anxiety	Bupropion Sr 100 Mg	PS	ORAL
TWO Q AM PO +				
	Hyperhidrosis			
ONE Q PM PO				
	Insomnia	Fluoxetine	C	
	Palpitations	Wellbutrin Sr	C	
	Pharmaceutical Product			
	Complaint			
	Trismus			

Date:04/29/04ISR Number: 4350088-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508665A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Carbon Monoxide Poisoning		Wellbutrin	PS	Glaxosmithkline	ORAL
		Completed Suicide		Remeron	C		
		Drug Toxicity		Hydroxyzine	C		
				Ambien	C		

Date:04/29/04ISR Number: 4350090-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313902A  
 Age:45 YR Gender:Male I/FU:F

Outcome	PT
Death	Cardio-Respiratory Arrest
	Coma

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Completed Suicide Depressed Level Of Consciousness	Report Source	Product	Role	Manufacturer	Route
		Intentional Misuse		Wellbutrin	PS	Glaxosmithkline	ORAL
				Doxepin	SS		ORAL
				Clorazepate	SS		ORAL
				Paxil	SS	Glaxosmithkline	
UNKNOWN							

Date:04/29/04ISR Number: 4350091-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313906A  
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aggression		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - PARENTERAL		Cardio-Respiratory Arrest		Insulin	SS		
Initial or Prolonged UNKNOWN		Completed Suicide		Citalopram	SS		
UNKNOWN		Convulsion		Trazodone	SS		
UNKNOWN		Decerebration		Vecuronium Bromide	C		
		Depressed Level Of Consciousness Hypertension Hyperthermia Tachycardia					

Date:04/29/04ISR Number: 4350092-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313909A  
Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death UNKNOWN		Arrhythmia		Wellbutrin	PS	Glaxosmithkline	
Hospitalization - UNKNOWN		Bundle Branch Block Right		Methadone	SS	Glaxosmithkline	
Initial or Prolonged UNKNOWN		Cardio-Respiratory Arrest		Alprazolam	SS		
UNKNOWN		Completed Suicide		Diphenhydramine	SS		

UNKNOWN

Dyspnoea

Cocaine

SS

Electrocardiogram Qt  
Prolonged  
Hypotension  
Hypothermia  
Hypoxia  
Multi-Organ Failure  
Respiratory Depression

Date:04/29/04ISR Number: 4350093-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313947A

Age:20 YR Gender:Female I/FU:F

Outcome

PT

Death

Blood Potassium Decreased  
Brain Death  
Completed Suicide  
Convulsion  
Dialysis  
Disseminated  
Intravascular Coagulation  
Drug Abuser  
Heart Rate Increased  
Hypotension  
Multi-Organ Failure  
Mydriasis  
Overdose  
Pupillary Reflex Impaired

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pyrexia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
100TAB			Wellbutrin	PS	Glaxosmithkline	ORAL
cumulative dose			Theophylline	SS		ORAL
100TAB			Heroin	SS		NASAL

Date:04/29/04ISR Number: 4350094-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313949A

Age:76 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN		Completed Suicide		Atenolol	SS		
UNKNOWN		Hypotension		Amlodipine	SS		
		Lethargy					
		Overdose					

Date:04/29/04ISR Number: 4350097-2Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0325638A

Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Alanine Aminotransferase		Zyban	PS	Glaxosmithkline	ORAL
1TAB Twice		Increased					
per day	2 WK	Arthralgia		No Concurrent			
		Aspartate		Medication	C		
		Aminotransferase					

Increased  
 Blood Alkaline  
 Phosphatase Increased  
 Gamma-Glutamyltransferase  
 Increased  
 Insomnia  
 Myalgia  
 Night Sweats  
 Urticaria  
 Weight Decreased

Date:04/29/04ISR Number: 4350099-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0328585A  
 Age:27 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG See				Zyban	PS	Glaxosmithkline	ORAL
dosage text	13 DAY		Supraventricular				
			Tachycardia				

Date:04/29/04ISR Number: 4350103-5Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0330307A  
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG See				Zyban	PS	Glaxosmithkline	ORAL
dosage text			Visual Acuity Reduced				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/29/04ISR Number: 4350105-9Report Type:Expedited (15-DaCompany Report #SK-GLAXOSMITHKLINE-B0330434A  
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abulia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Aphasia					
per day	20	DAY					
INTRAVENOUS		Cardiopulmonary Failure		Saline Solution	C	Glaxosmithkline	
		2 DAY					
.125MCG Twice		Catatonia		Digoxin	C	Glaxosmithkline	ORAL
per day		Cerebral Ataxia					
		Convulsion		Moduretic	C		ORAL
		Decreased Activity					
		Dehydration					
		Dementia					
		Dizziness					
		Dyspnoea					
		Epistaxis					
		Fatigue					
		Immobile					
		Loss Of Consciousness					
		Urinary Incontinence					

Date:04/29/04ISR Number: 4350108-4Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0330518A  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Depressed Level Of		Zyban	PS	Glaxosmithkline	ORAL
150MG See		Consciousness					
Initial or Prolonged		Grand Mal Convulsion		Mepronizine	C		
dosage text	14	DAY					
UNKNOWN		Malaise		Avlocardyl	C		
UNKNOWN				Tranxene	C		
UNKNOWN				Theralene	C		
UNKNOWN							

UNKNOWN	Loxapac	C	
UNKNOWN	Levothyrox	C	Glaxosmithkline
UNKNOWN	Dihydan	C	

Date:04/29/04ISR Number: 4350110-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0330717A  
 Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aggression		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	22 DAY	Excitability					
		Illusion					
		Irritability					
		Malaise					
		Palpitations					
		Road Traffic Accident					
		Thinking Abnormal					
		Vertigo					

Date:04/29/04ISR Number: 4350111-4Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0331014A  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Blood Pressure Diastolic		Zyban	PS	Glaxosmithkline	ORAL
6750MG Single		Decreased					
Initial or Prolonged		Grand Mal Convulsion		Polaramine	SS		ORAL
dose		Intentional Misuse					
30MG Single		Suicide Attempt		Aerius	C		
dose		Vomiting					
UNKNOWN							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/29/04ISR Number: 4350127-8Report Type:Expedited (15-DaCompany Report #US-ROCHE-360924

Age:26 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Life-Threatening	Convulsion		Klonopin	PS	Roche	ORAL
DOSAGE TAKEN						
Hospitalization -	Drug Interaction					
AS REQUIRED						
Initial or Prolonged (PRN).						
			Wellbutrin	SS		ORAL
DOSAGE						

REPORTED AS:

"AM". 449 DAY

Date:04/30/04ISR Number: 4350917-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313782A

Age:51 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Blood Pressure Decreased		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN						
Other	Completed Suicide		Acetaminophen +			
	Depressed Level Of		Propoxyphene			
	Consciousness		Napsylate	SS		ORAL
	Drug Toxicity		Carbamazepine	SS		
UNKNOWN						
			Acetaminophen +			
			Aspirin + Caffeine	C		
UNKNOWN						

Date:04/30/04ISR Number: 4350918-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313828A

Age:25 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Agitation		Bupropion	PS	Glaxosmithkline	ORAL
	Completed Suicide		Venlafaxine	SS		ORAL
	Convulsion					
	Haemodynamic Instability					

Hostility  
Mental Impairment  
Pyrexia

Date:04/30/04ISR Number: 4350919-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313890A  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
60TAB		Bundle Branch Block Left					
cumulative		Cardiac Arrest					
dose		Completed Suicide		Lorazepam	SS		ORAL
		Convulsion		Diphenhydramine	SS		ORAL
		Hypothermia		Sertraline	C		ORAL
30TAB		Hypoxia					
cumulative		Injury					
dose		Loss Of Consciousness					
		Nervous System Disorder					
		Overdose					
		Pupil Fixed					
		Rhabdomyolysis					
		Ventricular Fibrillation					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/30/04ISR Number: 4350920-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313891A  
 Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Bupropion	PS	Glaxosmithkline	ORAL
67TAB		Completed Suicide					
cumulative		Convulsion					
dose				Methylenedioxymetham phetamine	SS		ORAL

Date:04/30/04ISR Number: 4350921-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313893A  
 Age:17 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest		Bupropion	PS	Glaxosmithkline	ORAL
		Completed Suicide		Risperidone	SS		ORAL
		Convulsion		Sertraline	SS		ORAL
		Overdose					

Date:04/30/04ISR Number: 4350922-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313895A  
 Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Wellbutrin	PS	Glaxosmithkline	ORAL
7TAB		Completed Suicide					
cumulative		Convulsion					
dose		Dysarthria		Sertraline	SS		ORAL
7TAB		Hyperhidrosis					
cumulative		Mental Status Changes					
dose		Muscle Rigidity		Caffeine	SS	Glaxosmithkline	ORAL
		Overdose					

Date:04/30/04ISR Number: 4350923-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313896A  
 Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Acidosis		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Areflexia		Citalopram	SS		ORAL
		Aspiration		Zolpidem	SS		ORAL
		Bradycardia		Diazepam	SS		ORAL
		Cardio-Respiratory Arrest					
		Coma					
		Completed Suicide					
		Convulsion					
		Depressed Level Of Consciousness					
		Electroencephalogram Abnormal					
		Hyperkalaemia					
		Hypotension					
		Pupil Fixed					
		Pyrexia					
		Rhabdomyolysis					
		Tachycardia					
		Thrombocytopenia					
		Ventricular Fibrillation					
		Vomiting					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/30/04ISR Number: 4350924-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313897A  
 Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aggression		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Aspiration		Zolpidem	SS		ORAL
		Blood Alcohol Increased		Ethanol	SS		ORAL
		Cardio-Respiratory Arrest		Tetrahydrocannabinol	C		
		Coma		Amphetamines	C		
		Completed Suicide					
		Depressed Level Of Consciousness					
		Foreign Body Aspiration					
		Hypotension					
		Oxygen Saturation Decreased					
		Sedation					
		Toxicologic Test Abnormal					

Date:04/30/04ISR Number: 4350925-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313905A  
 Age:28 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Creatinine		Bupropion	PS	Glaxosmithkline	
UNKNOWN	13TAB per day	Increased		Verapamil	SS		ORAL
Other	30TAB per day	Blood Pressure Decreased		Venlafaxine	SS		
UNKNOWN	1575MG per day	Blood Urea Increased					
UNKNOWN		Bundle Branch Block		Cocaine	SS		
		Cardiac Arrest		Benzodiazepine	C		
		Completed Suicide					
		Convulsion					
		Heart Rate Decreased					
		Respiratory Rate Decreased					

Date:04/30/04ISR Number: 4350926-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313977A  
Age:19 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest		Lamictal	PS	Glaxosmithkline	ORAL
		Completed Suicide		Wellbutrin	SS	Glaxosmithkline	ORAL
		Convulsion		Citalopram	SS		ORAL
		Electrocardiogram Qrs		Acetaminophen+Dextro			
		Complex Prolonged		meth.Hbr+Doxyl.Succ+			
		Hypotension		Pseudoephed.Hcl	SS		
UNKNOWN							

Date:04/30/04ISR Number: 4350934-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0327761A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Zyban	PS	Glaxosmithkline	ORAL
Other		Haematoma					
150MG See		Myalgia					
dosage text	39 DAY	Oedema Peripheral					
		Serum Sickness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/30/04ISR Number: 4350942-0Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0330365A  
Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	21 DAY	Cerebral Haemorrhage		Zyban	PS	Glaxosmithkline	ORAL

Date:04/30/04ISR Number: 4350947-XReport Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0330593A  
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Coagulopathy		Bupropion Hydrochloride Ascal Persantin	PS C C	Glaxosmithkline	ORAL

Date:04/30/04ISR Number: 4350948-1Report Type:Expedited (15-DaCompany Report #CH-GLAXOSMITHKLINE-B0330728A  
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG per day 1 DAY Initial or Prolonged		Grand Mal Convulsion Leukoencephalopathy Nervous System Disorder		Zyban	PS	Glaxosmithkline	ORAL

Date:04/30/04ISR Number: 4350949-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0330834A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged dosage text 2 MON		Headache Nausea Nystagmus Vertigo		Zyban	PS	Glaxosmithkline	

Date:04/30/04ISR Number: 4350951-1Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0331251A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams Aggression Irritability Personality Change		Zyban	PS	Glaxosmithkline	

Date:04/30/04ISR Number: 4351296-6Report Type:Direct Company Report #CTU 217762

Age:9 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening ONCE DAILY		Self Injurious Behaviour Suicidal Ideation		Welbutrin	PS		

Date:04/30/04ISR Number: 4352145-2Report Type:Direct Company Report #CTU 217711

Age:87 YR Gender:Female I/FU:I

Outcome	PT
	Dizziness Nausea Pharmaceutical Product

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Complaint

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
BID			Wellbutrin 150 Sr	PS		

Date:05/03/04ISR Number: 4351750-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508981A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	1 MON		Medication Error	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/03/04ISR Number: 4351751-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0509114A  
 Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day			Coma	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day			Convulsion	Wellbutrin	SS	Glaxosmithkline	ORAL
			Depressed Level Of Consciousness	Ativan Tylenol	C C	Glaxosmithkline	

Date:05/03/04ISR Number: 4351755-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313777A  
 Age:19 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Acidosis	Wellbutrin	PS	Glaxosmithkline	ORAL
			Completed Suicide	Acetaminophen +			
			Condition Aggravated	Aspirin + Caffeine	SS		ORAL
UNKNOWN			Hypotension	Aspirin	SS	Glaxosmithkline	
UNKNOWN			Ventricular Tachycardia	Metformin	SS		

UNKNOWN	Citalopram	SS
UNKNOWN	Acetaminophen + Oxycodone	SS
UNKNOWN	Acetaminophen Codeine	SS
UNKNOWN	Diethylpropion	SS
UNKNOWN	Acetaminophen + Dextromethorphan + Pseudoephedrine	SS
UNKNOWN	Unspecified Drugs	SS

Date:05/03/04ISR Number: 4351756-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313822A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose		Bupropion	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged		Cardio-Respiratory Arrest Death		Methadone Diazepam	SS SS	Glaxosmithkline	ORAL ORAL

Date:05/03/04ISR Number: 4351757-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313871A  
Age:43 YR Gender:Male I/FU:F

Outcome	PT
Death	Adverse Drug Reaction
Other	Bradycardia Cardiac Arrest

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Death Eye Movement Disorder Mental Impairment				
		Muscle Rigidity Pyrexia	Paxil Wellbutrin Risperidone	PS SS SS	Glaxosmithkline Glaxosmithkline	ORAL ORAL
UNKNOWN			Clonazepam	SS		
UNKNOWN						

Date:05/04/04ISR Number: 4352725-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0509222A  
Age: Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
Hospitalization - 150MG Per day 5 DAY Initial or Prolonged		Fluid Retention Joint Stiffness Muscle Disorder	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/04/04ISR Number: 4352728-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313884A  
Age:15 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
Death		Cardio-Respiratory Arrest Cardiopulmonary Failure Convulsion Drowning Drug Abuser	Bupropion	PS	Glaxosmithkline	ORAL

Date:05/04/04ISR Number: 4352730-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0313889A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
Death		Cardio-Respiratory Arrest Cardiopulmonary Failure	Bupropion Acetaminophen	PS SS	Glaxosmithkline Glaxosmithkline	ORAL ORAL
75000MG per						

Completed Suicide

day

Drowning	Alprazolam	SS	ORAL
Intentional Misuse	Naproxen	SS	
Multiple Drug Overdose	Ma Huang	SS	

Date:05/04/04ISR Number: 4352740-0Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0043570A

Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day 2 DAY			Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Cardiovascular Disorder					

Date:05/05/04ISR Number: 4353563-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0509115A

Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG See			Zyban	PS	Glaxosmithkline	ORAL
dosage text	3 WK	Disorientation					
		Myalgia					
		Petechiae					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/05/04ISR Number: 4353573-1Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0330693A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Bupropion			
		Drug Interaction		Hydrochloride	PS	Glaxosmithkline	
		Hallucination		Cholesterol Reducing			
		Nasopharyngitis		Agent Name Not Known	C		
		Tremor		Cardiac Medication	C		

Date:05/05/04ISR Number: 4353574-3Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0330912A

Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema		Bupropion			
		Exanthem		Hydrochloride	PS	Glaxosmithkline	ORAL
150MG per day	18 DAY						
		Nasopharyngeal Disorder		Metformine	C		ORAL
500MG per day							
		Pruritus		Diclofenac Sodium	C	Glaxosmithkline	ORAL
150MG per day							
		Urticaria		Pariet	C		ORAL
				Orgametril	C		ORAL
5MG per day							

Date:05/05/04ISR Number: 4354100-5Report Type:Direct Company Report #CTU 217981

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Menopause		Buproprion 75 Mg Bid	PS		ORAL
PO							
		Pharmaceutical Product		Fluoxetine	C		
		Complaint					

Date:05/06/04ISR Number: 4354221-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0509451A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	400MG Per day	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
		Medication Error		Unknown Medication	C		

Date:05/06/04ISR Number: 4354223-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0509551A  
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	450MG Unknown	Loss Of Consciousness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Tremor					
		Urinary Incontinence					

Date:05/06/04ISR Number: 4354226-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0319984A  
 Age: Gender:Female I/FU:F

Outcome	PT
Other	Abdominal Pain Upper
	Anorexia
	Balance Disorder
	Constipation
	Deafness
	Disturbance In Attention
	Dysgeusia
	Ear Disorder
	Feeling Abnormal

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Nausea Sleep Disorder Tinnitus							
Vomiting				Zyban Valium	PS C	Glaxosmithkline	ORAL
Dose	2MG Twice per day						

Date:05/06/04ISR Number: 4356319-6Report Type:Expedited (15-DaCompany Report #S03-NOR-03725-01  
Age:23 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	40 MG QD PO	Convulsion Drug Interaction	Foreign Other	Cipramil (Citalopram Hydrobromide)	PS		ORAL
	150 MG BID PO	Loss Of Consciousness Tongue Biting		Zyban (Bupropion Hydrochloride)	SS		ORAL
	150 MG QD PO			Zyban (Bupropion Hydrochloride)	SS		ORAL
				Zopiklon (Zopiclone)	C		

Date:05/07/04ISR Number: 4354833-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506288A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Rash Vasculitis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/07/04ISR Number: 4354834-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0507922A  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Initial or Prolonged Medication Error  
dosage text 6 DAY  
Other Tongue Biting  
Vomiting

Oral Contraceptives C

Date:05/07/04ISR Number: 4354836-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0509822A  
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG See		Confusional State					
dosage text		Medication Error					
		Nervousness		Alprazolam	C		

Date:05/07/04ISR Number: 4354842-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0326019A  
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
Other		Asthenia					
150MG		Blood Creatine Increased					
Variable dose 1	MON	Blood Iron Decreased					
		Haemoglobin S Decreased					
		Hypercholesterolaemia					
		Iron Deficiency Anaemia					
		Pruritus Generalised					
		Renal Failure					





Date:05/10/04ISR Number: 4355579-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0510000A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Balance Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	9 WK	Dysarthria		Depakote	C		
3000MG per		Judgement Impaired					
day		Road Traffic Accident					
		Sluggishness					

Date:05/10/04ISR Number: 4355581-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0315354A

Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anxiety		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	6 WK	Delirium		Corticoid Ointment	C		
Initial or Prolonged		Depression					
TRANSDERMAL		Hallucination					
		Paranoia					
		Persecutory Delusion					
		Psychotic Disorder					

Date:05/10/04ISR Number: 4355602-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0332102A

Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Consciousness		Zyban	PS	Glaxosmithkline	ORAL
16 DAY		Malaise					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/10/04ISR Number: 4356307-XReport Type:Direct  
 Age:49 YR Gender:Female I/FU:I

Company Report #CTU 218278

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG PO BID	Fatigue	Pharmaceutical Product Complaint	Bupropion Sr	PS		ORAL

Date:05/11/04ISR Number: 4356092-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0503424A  
 Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	300MG Single	Disorientation		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - dose		Headache					
Initial or Prolonged	180MG Per day	White Blood Cell Count		Allegra	C		ORAL
Other	300MG Per day	Increased		Avalide	C		ORAL
	10MG At night			Singulair	C		ORAL
				Ibuprofen	C	Glaxosmithkline	

Date:05/11/04ISR Number: 4356095-7Report Type:Expedited (15-DaCompany Report #04US01863  
 Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	300MG In the morning	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - morning	449 DAY	Drug Interaction					
Initial or Prolonged	UNKNOWN			Klonopin	SS		
	1MG Three times per day						

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Simple Partial Seizures		Requip	PS	Glaxosmithkline	ORAL
		Tremor		Wellbutrin	SS	Glaxosmithkline	ORAL
2 YR				Selective Serotonin Reuptake Inhibitor	C		

Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alanine Aminotransferase		Zyban	PS	Glaxosmithkline	ORAL
1TAB Twice per day	2 WK	Increased		No Concurrent Medication	C		
		Arthralgia					
		Aspartate Aminotransferase					
		Increased					
		Blood Alkaline Phosphatase					
		Increased					
		Gamma-Glutamyltransferase					
		Increased					
		Insomnia					
		Myalgia					
		Night Sweats					
		Urticaria					
		Weight Decreased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/11/04ISR Number: 4356105-7Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0329071A  
 Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Depression		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization - per day	18 DAY	Psychotic Disorder					
Initial or Prolonged		Suicide Attempt		Quetiapine	C		ORAL
Other				Zoloft	C		ORAL
				Panadol	C	Glaxosmithkline	
				Endep	C	Glaxosmithkline	
				Vioxx	C		

Date:05/11/04ISR Number: 4356117-3Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0331481A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Face Injury		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Hypotension					
per day		Loss Of Consciousness		Beta Blocker	C		

Date:05/11/04ISR Number: 4356119-7Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0331580A  
 Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aggression		Bupropion			
		Depersonalisation		Hydrochloride	PS	Glaxosmithkline	
		Food Interaction					
		Hostility					
		Memory Impairment					

Date:05/11/04ISR Number: 4358785-9Report Type:Expedited (15-DaCompany Report #T02-USA-01514-01  
 Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Aggression Agitation Blood Glucose Increased	Study Health Professional	Memantine (Open Label, Phase B) (Memantine)	PS		ORAL
20 MG QD PO		Blood Urine Present Coma Dilatation Ventricular		Mjenatine (Open Label, Phase B) (Memantine)	SS		ORAL
20 MG QD PO		Grand Mal Convulsion Heart Rate Increased Incontinence		Memantine (Unblinded, Phase A) (Memantine)	SS		ORAL
10MG BID PO		Loss Of Consciousness Oxygen Saturation Decreased		Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
150 MG QD PO				Aricept (Donepezil Hydrochloride)	C		

Date:05/12/04ISR Number: 4357036-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508692A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 300MG Per day Other		Affect Lability  Crying Depressed Mood	Health  Professional	Wellbutrin Xl  No Concurrent Medications	PS  C	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/12/04ISR Number: 4357038-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0510367A  
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous		Bupropion	PS	Glaxosmithkline	ORAL

Date:05/12/04ISR Number: 4357048-5Report Type:Expedited (15-DaCompany Report #FI-GLAXOSMITHKLINE-B0331679A  
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG per day 13 DAY	Contusion		Zyban	PS	Glaxosmithkline	ORAL
		Loss Of Consciousness					
		Skin Laceration					

Date:05/12/04ISR Number: 4357605-6Report Type:Direct Company Report #CTU 218475  
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	300 MG DAILY	Amnesia		Wellbutrin	PS		ORAL
Initial or Prolonged	ORAL	Convulsion					
		Fall		Zoloft	C		
		Road Traffic Accident		Xanax	C		

Date:05/12/04ISR Number: 4357677-9Report Type:Direct Company Report #CTU 218489  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	ONE TAB Q DAY	Apathy		Wellbutrin Xl	PS		ORAL
		Condition Aggravated					
		Crying					
		Decreased Interest					
		Depression					

Drug Effect Decreased  
Self Esteem Decreased  
Suicidal Ideation

Date:05/12/04ISR Number: 4357826-2Report Type:Expedited (15-DaCompany Report #HQWYE676403MAY04  
Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blindness Unilateral Condition Aggravated Corneal Abrasion Glaucoma	Consumer	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		ORAL
150 MG 1X PER 1 DAY, ORAL				Wellbutrin (Amfebutanmone Hydrochloride)	SS		
HER DOSE "HAD DOUBLED"				Xalantan (Latanoprost)	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/13/04ISR Number: 4357708-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0499234A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
2 WK		Feeling Abnormal Headache Insomnia Medication Error Nausea Vomiting		Yasmin	C		

Date:05/13/04ISR Number: 4357719-0Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0331771A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Death		Zyban	PS	Glaxosmithkline	ORAL
Death							

Date:05/13/04ISR Number: 4357721-9Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0331807A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blindness		Quomem	PS	Glaxosmithkline	
Other							

Date:05/14/04ISR Number: 4358507-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0510766A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Atrial Fibrillation		Wellbutrin	PS	Glaxosmithkline	
Other				Coumadin	C	Glaxosmithkline	

Date:05/14/04ISR Number: 4358517-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0331984A  
Age:35 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG	Per day	4 DAY	Depressed Mood	Zyban	PS	Glaxosmithkline	ORAL
				Self-Injurious Ideation	Logynon	C		ORAL
				Suicidal Ideation				
				Tearfulness				

Date:05/14/04ISR Number: 4358518-6Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0331986A  
Age:27 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1TAB	Per day	7 DAY	Loss Of Consciousness	Zyban	PS	Glaxosmithkline	ORAL
	200MCG	Twice		Road Traffic Accident	Beclomethasone	C	Glaxosmithkline	
		per day						

Date:05/17/04ISR Number: 4359019-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508256A  
Age:78 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening				Grand Mal Convulsion	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Hospitalization -				Medication Error	Wellbutrin Sr	SS	Glaxosmithkline	ORAL
Initial or Prolonged					Alcohol	SS		ORAL
Other	25MG	per day			Atenolol	C		ORAL

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Freedom Of Information (FOI) Report

.5TAB per day				Triamterene + Hydrochlorothiazide	C	Glaxosmithkline	ORAL
40MG per day	YR			Pravacol	C		ORAL

Date:05/17/04ISR Number: 4359027-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0510886A  
Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous		Bupropion	PS	Glaxosmithkline	
150MG per day							

Date:05/17/04ISR Number: 4359033-6Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0328475A  
Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Bite		Zyban	PS	Glaxosmithkline	
150MG Twice per day		Deafness Dizziness Feeling Abnormal Myoclonus Palpitations Tinnitus Tongue Biting					

Date:05/17/04ISR Number: 4360761-7Report Type:Expedited (15-DaCompany Report #A0509420A  
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Pressure Increased Cerebrovascular Accident	Consumer	Nicoderm Cq Patch 21 Mg (Nicotine)	PS		
TRANSDERMAL DAY/	21 MG PER	Dizziness					

TRANSDERMAL	Erythema				
	Fatigue	Wellbutrin			
	Overdose	(Bupropion			
300 MG PER	Pruritus	Hydrochloride)	SS		ORAL
	Rash Papular				
DAY/ ORAL	Sensory Disturbance	Tarka	C		
	Speech Disorder	Alprazolam	C		
		Diuretic	C		

Date:05/18/04ISR Number: 4359773-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0505954A  
Age:20 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Amnesia	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day						
Initial or Prolonged	Convulsion	Professional	Over Counter			
	Drug Interaction		Medicines	SS		
	Fall					
	Haemorrhage					
	Head Injury					
	Skull Fracture					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/18/04ISR Number: 4359776-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0510479A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Murder		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN		Psychotic Disorder					

Date:05/18/04ISR Number: 4359777-6Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0510705A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Visual Acuity Reduced		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Unknown		Visual Disturbance					

Date:05/18/04ISR Number: 4360198-0Report Type:Direct Company Report #CTU 218881

Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Intolerance		Wellbutrin Sr 150 Mg	PS		ORAL
150 MG BID		Obstructive Airways					
ORAL		Disorder		Wellbutrin Xr 300 Mg	SS		ORAL
300 MG QD		Swollen Tongue					
ORAL							

Date:05/19/04ISR Number: 4360531-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511270A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Pharmaceutical Product					
per day							

Complaint

Date:05/19/04ISR Number: 4360532-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511280A  
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	5 WK		Feeling Abnormal	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	10 DAY		Medication Error	Bupropion	SS	Glaxosmithkline	ORAL
			Sluggishness	Protonix	C		

Date:05/19/04ISR Number: 4360533-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511289A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death			Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
Life-Threatening			Death				
Other							

Date:05/19/04ISR Number: 4360540-0Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0332296A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Convulsion	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice			Transient Ischaemic				
per day	4 DAY		Attack				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/19/04ISR Number: 4360629-6Report Type:Direct  
Age:33 YR Gender:Female I/FU:I

Company Report #CTU 218970

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - INTRA VENOUS 400 MG X 1	Grand Mal Convulsion		Tequin 400 Mg	PS		
Initial or Prolonged INTRA VENOUS	Rhabdomyolysis					
Required 150 MG DAILY			Wellbutrin Er	SS		ORAL
Intervention to ORAL						
Prevent Permanent Impairment/Damage			Prilosec Motrin	C C		

Date:05/19/04ISR Number: 4364138-XReport Type:Expedited (15-DaCompany Report #200412076BCC  
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Alanine Aminotransferase Increased	Health Professional	Aleve (Naproxen Sodium)	PS		ORAL
ORAL	Aspartate	Other	Centrum Vitamins	SS		ORAL
ORAL	Aminotransferase		Acetaminophen	SS		ORAL
	Increased Blood Pressure Diastolic Decreased Lethargy Multiple Drug Overdose Nausea Prothrombin Time Prolonged Sinus Tachycardia Vomiting		Imodium (Loperamide Hydrochloride) Wellbutrin (Bupropion Hydrochloride)	SS  SS		

Date:05/20/04ISR Number: 4361260-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508256A  
Age:78 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 2U Per day		Grand Mal Convulsion	Health Professional	Wellbutrin Sr Alcohol	PS SS	Glaxosmithkline	ORAL
Initial or Prolonged 25MG per day				Atenolol	C		ORAL
Other				Triamterene + Hydrochlorothiazide	C	Glaxosmithkline	ORAL
.5TAB per day				Pravacol	C		ORAL
40MG per day	YR						

Date:05/20/04ISR Number: 4361261-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0509115A  
Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG See		Conjunctival Haemorrhage	Health Professional	Zyban	PS	Glaxosmithkline	ORAL
dosage text 3 WK		Deep Vein Thrombosis	Professional				
		Disorientation Myalgia Petechiae		Ortho Tri-Cyclen	C		

Date:05/20/04ISR Number: 4361269-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511389A  
Age:22 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 1 WK		Athetosis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Initial or Prolonged Other		Cognitive Disorder Dyskinesia					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/20/04ISR Number: 4361271-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511459A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day	Renal Failure		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/20/04ISR Number: 4361272-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511475A

Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 6G Single		Diarrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged dose	1 DAY	Dizziness					
		Drug Screen Positive		Klonopin	SS		
5MG Single dose		Muscle Spasms					
		Overdose Vomiting		Lithium	C	Glaxosmithkline	

Date:05/20/04ISR Number: 4361280-4Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0330593A

Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Coagulopathy	Health Professional	Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
				Ascal	C		
				Persantin	C		

Date:05/20/04ISR Number: 4361286-5Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0332456A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Accidental Exposure Nervous System Disorder		Zyban	PS	Glaxosmithkline	

Date:05/20/04ISR Number: 4363737-9Report Type:Direct  
Age:51 YR Gender:Female I/FU:I

Company Report #CTU 219172

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated		Wellbutrin Sr	PS		
150 BID		Depression Pharmaceutical Product Complaint					

Date:05/21/04ISR Number: 4361437-2Report Type:Periodic  
Age:67 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0114838A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective Nausea Vomiting	Consumer	Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361438-4Report Type:Periodic  
Age:60 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0114981A

Outcome	PT
	Drug Ineffective
	Dry Mouth

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nausea

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
			Zyban	PS	Glaxosmithkline	ORAL
			Mvi	C		
			Tamoxifen	C		

Date:05/21/04ISR Number: 4361439-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0407078A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Urine Present		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	1 DAY						

Date:05/21/04ISR Number: 4361441-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0407652A  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Tremor		Clarinet	C		
per day	6 DAY			Spironolactone	C		

Date:05/21/04ISR Number: 4361442-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0407700A  
 Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urticaria		Zyban	PS	Glaxosmithkline	ORAL
18 DAY				Iron Sulfate	C	Glaxosmithkline	
				Folate	C		
				Celexa	C		

Date:05/21/04ISR Number: 4361443-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0408416A  
Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness		Bupropion	PS	Glaxosmithkline	ORAL
1 WK		Tinnitus					

Date:05/21/04ISR Number: 4361444-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0408563A  
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Nausea		Zyban	PS	Glaxosmithkline	ORAL
		Pharmaceutical Product		Zantac	C	Glaxosmithkline	
		Complaint		Phenergan	C	Glaxosmithkline	
		Vomiting					

Date:05/21/04ISR Number: 4361445-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0408817A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL
per day	WK	Tremor					
UNKNOWN				Ambien	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361446-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0408818A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Smoker		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	4 MON						

Date:05/21/04ISR Number: 4361447-5Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0409189A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	WK						

Date:05/21/04ISR Number: 4361448-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0410046A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown		Pruritus					

Date:05/21/04ISR Number: 4361449-9Report Type:Periodic  
 Age:24 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0410740A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization -							
150MG Twice							
Initial or Prolonged							
per day	2 WK						
Other							

Date:05/21/04ISR Number: 4361450-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0410771A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown		Urticaria					

Date:05/21/04ISR Number: 4361451-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0410772A  
Age:24 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anaphylactic Reaction		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization - 150MG Twice		Convulsion					
Initial or Prolonged per day 7 DAY		Drug Ineffective		Antibiotic	C		
Other							

Date:05/21/04ISR Number: 4361452-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0411252A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day 4 DAY		Heart Rate Increased					
		Rash					



150MG Twice	Constipation	Zyban	PS	Glaxosmithkline	ORAL
per day	Ejaculation Delayed				
	Sexual Dysfunction	Stool Softener	C		
		Titralac	C		
TOPICAL		Nicoderm	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361457-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0413310A  
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361458-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0414038A  
 Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice	42 DAY			Neurontin	C		
UNKNOWN				Zoloft	C		ORAL
UNKNOWN				Zocor	C		
UNKNOWN				Atenolol	C		
UNKNOWN				Triamterene	C	Glaxosmithkline	
UNKNOWN				Xanax	C		ORAL
UNKNOWN				Soma	C		
UNKNOWN				Vicoprofen	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361459-1Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0414538A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75MG Twice		Chills		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1 YR	Headache					
		Myalgia		Wellbutrin	SS	Glaxosmithkline	ORAL
		Pyrexia		Advil	C	Glaxosmithkline	
		Tinnitus					

Date:05/21/04ISR Number: 4361460-8Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0415365A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Pain Upper		Zyban	PS	Glaxosmithkline	ORAL
		Tremor					

Date:05/21/04ISR Number: 4361461-XReport Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418607A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 DAY		Hyperhidrosis		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361462-1Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0418793A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL
per day	WK	Dry Mouth					

Date:05/21/04ISR Number: 4361463-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0419202A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361464-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0419716A  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hiccups		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Vomiting					
per day	2 WK			Celebrex	C		

Date:05/21/04ISR Number: 4361465-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0419717A  
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Zyban	PS	Glaxosmithkline	ORAL
150MG Single		Paraesthesia					
dose	1 DAY			Premarin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361466-9Report Type:Periodic  
Age:43 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0419928A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Anxiety					
per day		Disturbance In Attention		Xanax	C		
		Feeling Jittery		Valium	C		
		Insomnia		Ambien	C		
		Mental Impairment		Oral Contraceptive	C		
		Weight Decreased		Dalmane	C		

Date:05/21/04ISR Number: 4361467-0Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0420791A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nonspecific Reaction		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361468-2Report Type:Periodic  
Age:53 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420806A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3 WK			Nicotine	C	Glaxosmithkline	
				Unknown Medication	C		

Date:05/21/04ISR Number: 4361469-4Report Type:Periodic  
Age:55 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0420919A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							

per day 9 DAY

Lipitor C

UNKNOWN

Lopressor C

UNKNOWN

Date:05/21/04ISR Number: 4361471-2Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420937A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Zyban	PS	Glaxosmithkline	ORAL
		Insomnia					
		Nervousness					

2 DAY

Date:05/21/04ISR Number: 4361472-4Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0420938A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Zyban	PS	Glaxosmithkline	ORAL
		Insomnia					
		Nervousness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361473-6Report Type:Periodic  
Age:49 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0422160A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG In the morning	2 DAY	Constipation Dry Mouth Feeling Abnormal Hallucination Headache Hyperacusis Keratoconjunctivitis Sicca Nervousness Oily Skin Paraesthesia		Zyban  Chewing Tobacco	PS  C	Glaxosmithkline	ORAL  ORAL

Date:05/21/04ISR Number: 4361474-8Report Type:Periodic  
Age:29 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422305A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	3 WK	Oedema Peripheral Pruritus Rash Urticaria		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361476-1Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0422908A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
10 DAY		Tinnitus		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361477-3Report Type:Periodic  
Age:36 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0422909A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2	WK		Zyban	PS	Glaxosmithkline	ORAL
				Ortho Tri-Cyclen Xalatan	C C		

Date:05/21/04ISR Number: 4361478-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0422918A  
 Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other				Zyban	PS	Glaxosmithkline	ORAL
1	MON						

Date:05/21/04ISR Number: 4361479-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0423429A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Unknown		DAY		Zyban	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361480-3Report Type:Periodic  
 Age: YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0423602A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	17 DAY	Urticaria		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361481-5Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0423864A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abnormal Behaviour Thinking Abnormal		Bupropion	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361482-7Report Type:Periodic  
 Age:40 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0423979A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	8 WK	Agitation Anxiety Dizziness Drug Ineffective Flight Of Ideas Insomnia Mood Altered Restlessness		Zyban  Doxycycline	PS  C	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361483-9Report Type:Periodic  
 Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424102A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	6 DAY	Feeling Jittery Mood Swings		Zyban	PS	Glaxosmithkline	ORAL

Nausea

Ortho Evra

C

Date:05/21/04ISR Number: 4361485-2Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424327A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Local Swelling		Zyban	PS	Glaxosmithkline	ORAL
per day	2 WK	Pruritus					
				Vitamins	C		

Date:05/21/04ISR Number: 4361486-4Report Type:Periodic  
Age:38 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0424756A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Unknown		Adverse Drug Reaction		Zyban	PS	Glaxosmithkline	ORAL
		Face Oedema		Vistaril	C		
		Pruritus					
		Rash					
		Urticaria					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361487-6Report Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0424885A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dementia		Zyban	PS	Glaxosmithkline	ORAL
1TAB Per day	3 DAY	Skin Tightness					

Date:05/21/04ISR Number: 4361488-8Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0425327A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Myalgia					
per day	3 WK	Pruritus		Vitamins	C		
		Swelling Face					
		Urticaria					

Date:05/21/04ISR Number: 4361489-XReport Type:Periodic  
Age:67 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0425340A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Activities Of Daily		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Living Impaired					
per day	8 DAY	Arthralgia		Synthroid	C	Glaxosmithkline	
		Depression		Amitriptyline	C		
				Ibuprofen	C	Glaxosmithkline	
				Evista	C		
				Prozac	C		
				Indapamide	C		
				Muscle Relaxer	C		
				Catapres-Tts-1	C		

Date:05/21/04ISR Number: 4361490-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0425920A  
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Delusion		Zyban	PS	Glaxosmithkline	ORAL
2 WK		Dry Mouth Insomnia Nightmare		None	C		

Date:05/21/04ISR Number: 4361491-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0425936A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Panic Attack		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice per day	6 DAY	Tremor		Birth Control	C		

Date:05/21/04ISR Number: 4361492-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0427087A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown UNKNOWN		Mood Swings Personality Change		Celexa	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361493-1Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0427728A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Swelling		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361494-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0427879A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice per day		Tremor		Centrum Multivitamin	C		

Date:05/21/04ISR Number: 4361495-5Report Type:Periodic  
Age:66 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428257A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Zyban	PS	Glaxosmithkline	ORAL
10 DAY				Chemotherapy	C		

Date:05/21/04ISR Number: 4361496-7Report Type:Periodic  
Age:25 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428700A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Jittery		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	5 DAY						

Date:05/21/04ISR Number: 4361497-9Report Type:Periodic  
Age:30 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0429240A

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Convulsion		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361498-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0429268A  
Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Irritability		Zyban	PS	Glaxosmithkline	
UNKNOWN							

Date:05/21/04ISR Number: 4361499-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0429516A  
Age: Gender:Female I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Face Oedema		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day							
Muscle Rigidity							
Oedema Peripheral							
Paraesthesia Oral							

Date:05/21/04ISR Number: 4361500-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0429531A  
Age:54 YR Gender:Male I/FU:F

Outcome		PT
		Dizziness
		Energy Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Insomnia Sinusitis Vertigo	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Zyban	PS	Glaxosmithkline	ORAL
150MG Twice per day	3 MON			Xanax	C		

Date:05/21/04ISR Number: 4361501-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430039A  
Age: Gender:Female I/FU:F

		PT Urticaria	Report Source	Product	Role	Manufacturer	Route
Outcome Dose	Duration		Consumer	Zyban	PS	Glaxosmithkline	ORAL
2 WK				Patch (Unspecified)	SS		

Date:05/21/04ISR Number: 4361502-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430440A  
Age:47 YR Gender:Female I/FU:I

		PT Feeling Abnormal	Report Source	Product	Role	Manufacturer	Route
Outcome Dose	Duration			Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK						

Date:05/21/04ISR Number: 4361503-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430741A  
Age: Gender:Female I/FU:I

		PT Pyrexia Urticaria	Report Source	Product	Role	Manufacturer	Route
Outcome Dose	Duration			Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361504-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431778A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Zyban	PS	Glaxosmithkline	ORAL
1TAB Twice							
per day	2	WK					

Date:05/21/04ISR Number: 4361505-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432989A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361506-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439011A  
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	12	DAY		Oral Contraception	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361507-9Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439262A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL
per day	9 DAY			Prevacid	C		
				Actigall	C		

Date:05/21/04ISR Number: 4361508-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439427A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		Feeling Jittery		Zyban	PS	Glaxosmithkline	
		Insomnia					

Date:05/21/04ISR Number: 4361509-2Report Type:Periodic  
Age:20 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440622A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Constipation		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361510-9Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441190A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Dry Mouth		Zyban	PS	Glaxosmithkline	ORAL
per day	11 DAY	Fatigue					
		Neck Pain					

Date:05/21/04ISR Number: 4361511-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441528A  
Age: Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG	Unknown	Hypersensitivity		Zyban	PS	Glaxosmithkline	ORAL
		Pruritus					
		Urticaria					

Date:05/21/04ISR Number: 4361512-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442158A  
Age:42 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2	WK	Ageusia		Zyban	PS	Glaxosmithkline	ORAL
		Diarrhoea		Synthroid	C	Glaxosmithkline	
		Gastrooesophageal Reflux Disease					
		Nausea					
		Vomiting					

Date:05/21/04ISR Number: 4361513-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442534A  
Age:62 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2	DAY	Rash		Zyban	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361514-6Report Type:Periodic  
 Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442778A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL
2 WK				Lipitor	C		

Date:05/21/04ISR Number: 4361515-8Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443560A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dementia		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown		Homicidal Ideation					

Date:05/21/04ISR Number: 4361516-XReport Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0490990A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown		Incoherent Tremor					

Date:05/21/04ISR Number: 4361517-1Report Type:Periodic  
 Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493285A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Twitching		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day 2 DAY				None	C		

Date:05/21/04ISR Number: 4361518-3Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0493497A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression Fatigue		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361519-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0494821A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral Urticaria		Zyban Wellbutrin	PS C	Glaxosmithkline Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361520-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0496019B  
 Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cerebrovascular Accident Convulsion		Zyban Nicoderm Patch	PS C	Glaxosmithkline Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361521-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496342A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 WK		Anorexia Depression Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361522-5Report Type:Periodic  
 Age:41 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0497219A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	5 DAY	Myalgia		Zyban	PS	Glaxosmithkline	ORAL
				Zetia	C		

Date:05/21/04ISR Number: 4361523-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498011A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361524-9Report Type:Periodic  
 Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499841A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	4 DAY	Dysgeusia		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361525-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0500122A  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice			Professional				
per day	10	DAY					

Date:05/21/04ISR Number: 4361526-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0500153A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nicotine Dependence		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	3	MON					

Date:05/21/04ISR Number: 4361527-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0500837A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361528-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0500839A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Therapeutic Response					
per day	1	MON		Imitrex	C	Glaxosmithkline	NASAL
		Unexpected					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Amerge C Glaxosmithkline  
 Verapamil C

Date:05/21/04ISR Number: 4361529-8Report Type:Periodic  
 Age:41 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0501341A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Three		Arthralgia		Zyban	PS	Glaxosmithkline	ORAL
times per day	13	Fatigue					
	DAY	Myalgia					
		Pyrexia					
		Serum Sickness					
		Urticaria					

Date:05/21/04ISR Number: 4361530-4Report Type:Periodic  
 Age:46 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501412A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Pruritus	Consumer	Zyban	PS	Glaxosmithkline	ORAL
per day	4	Urticaria					
	DAY			Glucosamine + Chondroitin	SS		
1TAB Per day	5			Multivitamin	C		
	WK			Calcium Supplement	C		
				Vitamin C	C	Glaxosmithkline	
				Benicar	C		
				Magnesium Supplement	C		
				Zinc Supplement	C		

Date:05/21/04ISR Number: 4361531-6Report Type:Periodic  
 Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503009A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Twice	Drug Ineffective	Zyban	PS	Glaxosmithkline	ORAL
per day					
		Ortho Tri-Cyclen	C		
		Nicorette	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361532-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0503215A  
 Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Rash					
per day	2 WK	Rash Erythematous		Neurontin	C		
		Vomiting		Albuterol	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361533-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505136A  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization -				Hydrocodone	C		
Initial or Prolonged							
UNKNOWN							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361534-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505665A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 WK		Anger		Zyban	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361535-3Report Type:Periodic  
Age:76 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505968A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 WK	Anxiety Nervousness Tremor		Zyban	PS	Glaxosmithkline	ORAL
				Zocor	C		
				Xalatan	C		
				Aerobid	C		
				Floratil	C		

Date:05/21/04ISR Number: 4361536-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506110A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Depressed Mood Insomnia		Zyban	PS	Glaxosmithkline	ORAL
				Depo Provera	C		

Date:05/21/04ISR Number: 4361537-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506353A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 WK		Depression Drug Withdrawal Syndrome		Zyban	PS	Glaxosmithkline	
				Wellbutrin XL	SS	Glaxosmithkline	ORAL

Insomnia

Date:05/21/04ISR Number: 4361538-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0506388A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Zyban	PS	Glaxosmithkline	

Date:05/21/04ISR Number: 4361539-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506806A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Zyban	PS	Glaxosmithkline	ORAL
		Nervousness		Wellbutrin	SS	Glaxosmithkline	ORAL
		Tinnitus					

Date:05/21/04ISR Number: 4361540-7Report Type:Periodic  
Age:65 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0507937A

Outcome	PT
	Dermatitis Exfoliative
	Dry Mouth
	Dry Skin
	Haemorrhage Urinary Tract
	Rash



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Rash Papular

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	5 DAY		Zyban	PS	Glaxosmithkline	ORAL
			Altace	C		
			Atenolol	C		
			Zocor	C		
			Xanax	C		

Date:05/21/04ISR Number: 4361541-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508070A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Paraesthesia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice per day	2 WK			Depo-Provera	C		

Date:05/21/04ISR Number: 4361542-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508438A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day				Hctz	C		
				Accupril	C		
				Zocor	C		

Date:05/21/04ISR Number: 4361543-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508671A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Per day	7	DAY	Hypersensitivity	Zyban	PS	Glaxosmithkline	ORAL
			Nausea				
			Urticaria				
Date:05/21/04			ISR Number: 4361544-4	Report Type:Periodic	Company Report #US-GLAXOSMITHKLINE-A0508929A		
Age:	Gender:Male	I/FU:F					
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Zyban	PS	Glaxosmithkline	
24	DAY	Urticaria					
Date:05/21/04			ISR Number: 4361545-6	Report Type:Periodic	Company Report #US-GLAXOSMITHKLINE-A0509163A		
Age:	Gender:Female	I/FU:I					
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Zyban	PS	Glaxosmithkline	ORAL
3	WK						
Date:05/21/04			ISR Number: 4361547-X	Report Type:Periodic	Company Report #US-GLAXOSMITHKLINE-A0505370A		
Age: YR	Gender:	I/FU:F					
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	
19-Aug-2005 12:44 PM							
Page: 2593							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361548-1Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0505372A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1 WK			Amerge	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361549-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505378A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Psychotropic Medications	C		

Date:05/21/04ISR Number: 4361550-XReport Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505380A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice		Headache					
per day	3 WK	Insomnia		Celexa	C		
		Pollakiuria		Synthroid	C	Glaxosmithkline	
		Pruritus					

Date:05/21/04ISR Number: 4361551-1Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505385A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK			Covera	C		
				Lipitor	C		
				Diovan Hct	C		

Glucophage C  
Vicodin Es C  
Mobic C

Date:05/21/04ISR Number: 4361552-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505447A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
1	WK			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Accutane	C		

Date:05/21/04ISR Number: 4361553-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505448A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG	Unknown			Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361554-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505458A  
Age:37 YR Gender:Male I/FU:F

Outcome PT  
Hospitalization - Chest Pain  
Initial or Prolonged Hypertension

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Irritability

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
4	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Lotrel	C		
			Toprol Xl	C		
			Hctz	C		

Date:05/21/04ISR Number: 4361555-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505463A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Unknown	1	MON					

Date:05/21/04ISR Number: 4361556-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505547A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphagia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:05/21/04ISR Number: 4361557-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505573A  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gynaecomastia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day	4	YR		Wellbutrin Sr	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361559-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505591A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:05/21/04ISR Number: 4361560-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505607A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361561-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505672A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Myalgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 3 DAY				No Concurrent Medication	C		

Date:05/21/04ISR Number: 4361562-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505677A  
Age: Gender:Female I/FU:I

Outcome  
PT  
Anxiety  
Crying  
Depression

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hypoventilation

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			No Concurrent Medication	C		

Date:05/21/04ISR Number: 4361563-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505682A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3 WK	Oligomenorrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:05/21/04ISR Number: 4361564-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505687A  
 Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
				Prozac	C		

Date:05/21/04ISR Number: 4361565-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505697A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300TAB Per day		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361566-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505698A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menstruation Irregular		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361567-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505700A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 DAY	Swelling Face		Klonopin	C		

Date:05/21/04ISR Number: 4361569-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505932A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Local Swelling		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK	Pruritus		No Concurrent Medication	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361570-5Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0505940A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN							

Date:05/21/04ISR Number: 4361571-7Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505961A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	6 DAY	Abnormal Dreams		Wellbutrin	PS	Glaxosmithkline	ORAL
		Musculoskeletal Stiffness		No Concurrent Medication	C		

Date:05/21/04ISR Number: 4361572-9Report Type:Periodic  
Age:64 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505981A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1 DAY	Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Prinivil	C		
				Xanax	C		
				Ativan	C		

Date:05/21/04ISR Number: 4361573-0Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0505985A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	1 MON	Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361574-2Report Type:Periodic  
Age:34 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505998A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Somnolence		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY			Effexor	C		
				Diovan	C		

Date:05/21/04ISR Number: 4361575-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505999A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Paraesthesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6 WK			Thyroid Medication	C		

Date:05/21/04ISR Number: 4361576-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506016A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	30 MON	Dyspepsia					
		Hyperhidrosis					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361577-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506039A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	8 WK	Abdominal Distension		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361578-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506047A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361579-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506088A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety Heart Rate Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361580-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506090A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 WK		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361581-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506092A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 MON		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361582-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506093A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorgasmia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Libido Increased		Strattera	C		
				Zoloft	C		

Date:05/21/04ISR Number: 4361583-3Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506097A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other							
150MG Per day	108 DAY						

Date:05/21/04ISR Number: 4361584-5Report Type:Periodic  
Age:59 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0506128A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
		Depression		Guaifenesin	C		
		Weight Increased		Valium	C		ORAL
5MG As							
required							
				Kcl	C	Glaxosmithkline	ORAL
20MEQ Unknown				Librax	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Diovan Hct C

Date:05/21/04ISR Number: 4361585-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506129A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Somnolence		Wellbutrin Xl Effexor	PS C	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361586-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506131A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Disturbance In Attention Feeling Abnormal Memory Impairment		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361588-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506132A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Prolactin Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361589-4Report Type:Periodic  
Age:40 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0506133A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oedema Peripheral		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361590-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506134A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Swelling Face		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361591-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506240A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day	6 MON	Weight Increased		Bentyl	C		
				Premarin	C		

Date:05/21/04ISR Number: 4361592-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506243A  
 Age: Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361593-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506245A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	2	WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Anger					
		Anxiety		Valtrex	C	Glaxosmithkline	
		Herpes Simplex		Synthroid	C	Glaxosmithkline	
		Mood Altered		Zelnorm	C		

Date:05/21/04ISR Number: 4361595-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506252A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Agitation					
		Tremor					

Date:05/21/04ISR Number: 4361596-1Report Type:Periodic  
 Age:25 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506269A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300MG Per day				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Convulsion					

Date:05/21/04ISR Number: 4361597-3Report Type:Periodic  
 Age:70 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506289A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361599-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506368A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Nervousness		Zoloft	SS		
		Psychomotor Hyperactivity		Synthroid	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361600-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506390A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK						

Date:05/21/04ISR Number: 4361601-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506402A  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4 MON	Tremor					

Date:05/21/04ISR Number: 4361604-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506481A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1 MON						



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361605-XReport Type:Periodic  
 Age:90 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0506485A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1 DAY	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Balance Disorder		Aricept	C		
		Feeling Abnormal		Coumadin	C	Glaxosmithkline	
				Lopid	C		
				Vitamins	C		
				Actonel	C		
				Calcium	C		

Date:05/21/04ISR Number: 4361606-1Report Type:Periodic  
 Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506492A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	1 MON	Myoclonus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
250MG Per day				Wellbutrin Sr	SS	Glaxosmithkline	ORAL
				Celexa	C		

Date:05/21/04ISR Number: 4361607-3Report Type:Periodic  
 Age:44 MON Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506499A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Agitation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Tremor		Alcohol	C		
		Weight Decreased					

Date:05/21/04ISR Number: 4361608-5Report Type:Periodic  
 Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506501A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3 WK	Eye Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Zyprexa C  
Topamax C  
Klonopin C  
Nexium C  
Prozac C

Date:05/21/04ISR Number: 4361609-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506502A  
Age:30 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
300MG Per day		Stool Analysis Abnormal	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361610-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506503A  
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
300MG Per day	4 DAY	Nausea	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361611-5Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506516A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Blood Lactic Acid					
		Increased		Remeron	C		
				Ovcon	C		
				Vioxx	C		
				Wellbutrin Sr	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361612-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506523A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
1 MON		Tinnitus					
				Effexor	C		

Date:05/21/04ISR Number: 4361613-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506577A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
5 YR		Oedema Peripheral					
		Pruritus					

Date:05/21/04ISR Number: 4361614-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506578A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2 WK		Adverse Event					

Date:05/21/04ISR Number: 4361615-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506581A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361616-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506794A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other							

Date:05/21/04ISR Number: 4361617-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506825A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Premature Ejaculation		Wellbutrin	PS	Glaxosmithkline	

Date:05/21/04ISR Number: 4361618-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506832A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - 300MG Per day Initial or Prolonged							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361619-XReport Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506839A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3	WK					
		Mood Altered		Valtrex	C	Glaxosmithkline	
				Synthroid	C	Glaxosmithkline	
				Vitamins	C		
				Tylenol Pm	C		
				Effexor	C		

Date:05/21/04ISR Number: 4361620-6Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506842A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Heart Rate Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	12	DAY					
		Weight Decreased		Benicar	C		

Date:05/21/04ISR Number: 4361621-8Report Type:Periodic  
Age:29 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506843A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1	WK					
				Prilosec	SS		
				Centrum	C		

Date:05/21/04ISR Number: 4361622-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506845A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL
				Effexor	SS		

Date:05/21/04ISR Number: 4361623-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506849A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	6	MON					

Date:05/21/04ISR Number: 4361624-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506915A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Central Nervous System		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice		Stimulation		Wellbutrin Sr	C	Glaxosmithkline	
per day	5	Crying					
	YR	Emotional Distress					

Date:05/21/04ISR Number: 4361626-7Report Type:Periodic  
Age:59 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506916A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	6	Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	WK			Diovan	C		
				Synthroid	C	Glaxosmithkline	
				Prevacid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Zoloft C

Date:05/21/04ISR Number: 4361627-9Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506917A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3	MON	Drug Ineffective	Trazodone	C		

Date:05/21/04ISR Number: 4361628-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506942A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Mucous Membrane Disorder	Advair	C	Glaxosmithkline	
				Flovent	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361629-2Report Type:Periodic  
Age:51 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506945A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2	WK	Drug Ineffective				
			Insomnia				
			Somnolence				

Date:05/21/04ISR Number: 4361630-9Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0506956A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other			Convulsion				

Date:05/21/04ISR Number: 4361631-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0507051A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
16 DAY		Delusion		Elavil	C	Glaxosmithkline	ORAL
50MG At night		Paranoia					

Date:05/21/04ISR Number: 4361632-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0507076A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Faeces Hard		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
1 MON		Haemorrhoids					

Date:05/21/04ISR Number: 4361634-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0507160A  
Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysarthria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK	Tremor		Synthroid	C	Glaxosmithkline	
				Zestril	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361635-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507170A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	4	MON		Alcohol	C		

Date:05/21/04ISR Number: 4361636-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507175A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3	WK		Seroquel	C		
		Back Pain		Zoloft	C		
		Nausea		Zyrtec	C	Glaxosmithkline	
				Xanax	C		
				Mobic	C		
				Protonix	C		
				Albuterol Mdi	C	Glaxosmithkline	
				Neurontin	C		
				Trazodone	C		
				Compazine	C	Glaxosmithkline	
				Prednisone	C		

1 YR

Date:05/21/04ISR Number: 4361637-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507206A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Urinary Retention		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Oxycontin	SS		

Date:05/21/04ISR Number: 4361638-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507207A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Myalgia		Wellbutrin Xl Anti-Psychotic Medication	PS C	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361639-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0507254A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin Xl Effexor	PS C	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361640-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0507267A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other	300MG Per day			No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361641-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507269A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Feeling Jittery		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR	Overdose		Ambien	C		
				Trazec	C		
				Darvocet	C		

Date:05/21/04ISR Number: 4361642-5Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507275A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Panic Attack		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	10 DAY			Birth Control	C		
				Wellbutrin Sr	C	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361643-7Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507288A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Asthenia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 YR	Decreased Interest		Armour Thyroid	C		
				Strattera	C		
40MG Per day							

Date:05/21/04ISR Number: 4361644-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507440A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 MON	Psychomotor Hyperactivity					

Date:05/21/04ISR Number: 4361645-0Report Type:Periodic  
Age:12 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507444A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2	WK		Trazodone	C		
				Flonase	C	Glaxosmithkline	
				Clarinet	C		

Date:05/21/04ISR Number: 4361646-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507449A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361647-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507452A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1	MON					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361648-6Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507456A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	1 DAY	Anxiety	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Xanax	C		

Date:05/21/04ISR Number: 4361649-8Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0507465A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	6 WK	Other	Wellbutrin	PS	Glaxosmithkline	ORAL
			Convulsion				

Date:05/21/04ISR Number: 4361650-4Report Type:Periodic  
Age:77 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507469A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	3 MON	Dyspepsia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Dyspnoea	Hctz	C		
			Fatigue	Lopressor	C		
			Headache	Klonopin	C		
			Nausea				

Date:05/21/04ISR Number: 4361651-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507541A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Unknown		Insomnia	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361652-8Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0507542A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gingival Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361653-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0507559A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Wellbutrin Xl	PS	Glaxosmithkline	

Date:05/21/04ISR Number: 4361654-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0507566A  
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 MON	Serum Ferritin Increased					

Date:05/21/04ISR Number: 4361655-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0507567A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 MON			Vasotec	C		
				Lasix	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Birth Control Pills C

Date:05/21/04ISR Number: 4361656-5Report Type:Periodic  
Age:34 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507582A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	6	MON		None	C		
		Pyrexia					
		Vomiting					

Date:05/21/04ISR Number: 4361657-7Report Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507587A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2	MON		Hiv Treatments (Unspecified)	C		

Date:05/21/04ISR Number: 4361658-9Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507594A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	12	DAY		Premarin	C		
		Nausea		Xanax	C		
				Novastan	C		

Date:05/21/04ISR Number: 4361659-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507624A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

1 MON Muscle Spasms Wellbutrin Xl PS Glaxosmithkline ORAL

Date:05/21/04ISR Number: 4361663-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0507666A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4 WK	Back Pain					
		Headache		Metformin	C		
				Vitamins	C		

Date:05/21/04ISR Number: 4361664-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0507679A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Drug Ineffective					
				Zyprexa	C		
				Zoloft	C		
				Zocor	C		
				Hctz	C		
				Estradiol	C		
				Aspirin	C	Glaxosmithkline	
				Vitamins	C		
				B12 Vitamin	C	Glaxosmithkline	
				Xanax	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361665-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507700A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	

Date:05/21/04ISR Number: 4361666-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507702A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361667-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507843A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Crying Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361668-1Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507864A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	16 DAY	Aura Grand Mal Convulsion					

Date:05/21/04ISR Number: 4361669-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507896A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG	Twice per day	Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL
					Accupril	C		
Date:05/21/04ISR Number: 4361670-XReport Type:Periodic				Company Report #US-GLAXOSMITHKLINE-A0507899A				
Age:		Gender:Female		I/FU:F				
	150MG	Per day	Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
Date:05/21/04ISR Number: 4361671-1Report Type:Periodic				Company Report #US-GLAXOSMITHKLINE-A0507900A				
Age:		Gender:		I/FU:F				
	150MG	Per day	Myoclonus		Wellbutrin	PS	Glaxosmithkline	ORAL
Date:05/21/04ISR Number: 4361672-3Report Type:Periodic				Company Report #US-GLAXOSMITHKLINE-A0507921A				
Age:45 YR		Gender:Female		I/FU:I				
	150MG	Per day	Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL
		1 WK	Oedema		Lexapro	C		
			Rash					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361673-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507928A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
6 WK		Anger		Wellbutrin	PS	Glaxosmithkline	ORAL
		Feeling Abnormal Ill-Defined Disorder Suicidal Ideation					

Date:05/21/04ISR Number: 4361674-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507932A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urticaria		Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361675-9Report Type:Periodic  
Age:72 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507941A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	30 MON	Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Ineffective		Propranolol	C		
		Increased Tendency To Bruise		Atrovent	C		
		Insomnia		Prempro	C		
		Tremor		Bextra	C		
		Weight Decreased		Calcium	C		
				Multivitamin	C		
				Fish Oil	C		
				B12 Vitamin	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361676-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508019A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Memory Impairment

Wellbutrin Xl

PS

Glaxosmithkline

ORAL

Date:05/21/04ISR Number: 4361677-2Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0508054A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Lexapro	C		

Date:05/21/04ISR Number: 4361678-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0508055A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3 WK		Anger					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361679-6Report Type:Periodic  
 Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508061A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Night Sweats		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	4	MON		Celexa	C		

Date:05/21/04ISR Number: 4361680-2Report Type:Periodic  
 Age:59 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508067A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4	WK		Synthroid	C	Glaxosmithkline	
		Weight Decreased		Lamictal	C	Glaxosmithkline	
				Neurontin	C		
				Klonopin	C		

Date:05/21/04ISR Number: 4361681-4Report Type:Periodic  
 Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508099A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other			Health				
450MG Per day	10	WK	Professional	Zoloft	C		
200MG per day							

Date:05/21/04ISR Number: 4361682-6Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508187A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361683-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508332A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361684-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508337A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Cholesterol		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	4	MON		Lexapro	C		
		Increased		Zocor	C		

Date:05/21/04ISR Number: 4361685-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508341A  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	6	DAY		Toprol	C		
		Insomnia		Hctz	C		
		Swelling Face					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361686-3Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0508355A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	3 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
		Anger		Wellbutrin Sr	C	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361687-5Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508383A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Energy Increased Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361688-7Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508384A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ill-Defined Disorder Therapeutic Response Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361690-5Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508402A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Erythematous Rash Pruritic Swelling		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361691-7Report Type:Periodic  
 Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508421A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG	Per day	Abdominal Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
			Anorexia		Vitamin A	C		
			Menorrhagia		Vitamin C	C	Glaxosmithkline	
			Metrorrhagia		Ester C	C		
					Vitamin D	C		
					Vitamin E	C		
					Coenzyme Q10	C		
					Magnesium	C		
					Alpha Lipoic Acid	C		
					Flax Oil	C		
					Fish Oil	C		

Date:05/21/04ISR Number: 4361692-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508429A  
Age: Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG	Per day	Dry Skin		Wellbutrin	PS	Glaxosmithkline	ORAL
		5 MON	Pruritus		Synthroid	C	Glaxosmithkline	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361693-0Report Type:Periodic  
Age:14 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508449A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Jaw Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 DAY	Pain In Jaw		Neurontin	C		

Date:05/21/04ISR Number: 4361694-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508455A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
2 WK							

Date:05/21/04ISR Number: 4361695-4Report Type:Periodic  
Age:53 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0508458A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 MON	Nausea		Darvon	C		
				Oxycontin	C		

Date:05/21/04ISR Number: 4361696-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508466A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR			No Concurrent Medication	C		

Date:05/21/04ISR Number: 4361697-8Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508468A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6 DAY	Nausea					

Date:05/21/04ISR Number: 4361698-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508638A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice		Erythema					
per day	3 WK	Urticaria		Levothyroxine	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361699-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508653A  
 Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Deafness		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Dizziness		Hctz	C		
300MG Per day	0 DAY	Tinnitus		Atenolol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361700-5Report Type:Periodic  
Age:11 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508658A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other				Concerta	C		

Date:05/21/04ISR Number: 4361701-7Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0508662A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	
Other							

Date:05/21/04ISR Number: 4361702-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508672A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							

Date:05/21/04ISR Number: 4361703-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508691A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other							

Date:05/21/04ISR Number: 4361704-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508732A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Unknown Medication	C		

Date:05/21/04ISR Number: 4361705-4Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508748A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY	Muscle Spasms		Zoloft	C		
		Rash Papular		Trazodone	C		

Date:05/21/04ISR Number: 4361706-6Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508757A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK	Flatulence		Vioxx	C		
				Reglan	C	Glaxosmithkline	
				Nexium	C		
				Flonase	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361707-8Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508767A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	4	WK		Wellbutrin	PS	Glaxosmithkline	ORAL
				Thyroid Medication	C		

Date:05/21/04ISR Number: 4361708-XReport Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508774A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1	MON		Wellbutrin	PS	Glaxosmithkline	ORAL
				Vitamins	C		
				Xanax	C		

Date:05/21/04ISR Number: 4361709-1Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0508948A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening				Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				Adderall	C		
				Lidocaine	C		

Date:05/21/04ISR Number: 4361710-8Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0508950A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening				Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -							
Initial or Prolonged							

Date:05/21/04ISR Number: 4361711-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508953A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vomiting		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361712-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508977A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Exfoliative		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	8	WK					

Date:05/21/04ISR Number: 4361713-3Report Type:Periodic  
Age:43 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0509122A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1	WK		Advil	C	Glaxosmithkline	
		Drug Ineffective					
		Headache					
		Menstrual Discomfort					
		Nausea					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361714-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509136A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	4 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
		Dizziness					
		Nausea					
		Pruritus					

Date:05/21/04ISR Number: 4361715-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509157A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Pruritus					

Date:05/21/04ISR Number: 4361716-9Report Type:Periodic  
Age:50 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0509158A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	450MG Per day	53 DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other		Urticaria		Ultram	C		
				Celebrex	C		
				Ambien	C		

UNKNOWN

Date:05/21/04ISR Number: 4361718-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509198A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Constipation					

Date:05/21/04ISR Number: 4361719-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0509206A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	
UNKNOWN							

Date:05/21/04ISR Number: 4361720-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0509553A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							
Dizziness Hyperhidrosis Nausea							

Date:05/21/04ISR Number: 4361721-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0509601A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Coordination Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361722-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0510023A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Crying		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Therapeutic Response		Namenda	SS		
		Decreased					

Date:05/21/04ISR Number: 4361724-8Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500838A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Stool Analysis Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	7	WK		Effexor	C		

Date:05/21/04ISR Number: 4361725-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500844A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3	WK					
		Headache					
		Nausea					

Date:05/21/04ISR Number: 4361726-1Report Type:Periodic  
Age:61 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500861A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abnormal Dreams		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
1TAB Per day	6	WK		No Concurrent Medication	C		
		Anxiety					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	5 DAY	Agitation		Provigil	C		
		Dizziness		Valium	C		
				Prevacid	C		
				Tricor	C		
				Zanaflex	C		
				Darvocet	C		
				Climara Patch	C		
				Actonel	C		
				Levoxyl	C	Glaxosmithkline	
				B12 Injections	C	Glaxosmithkline	
				Betaferon	C		
				Niaspan	C		

UNKNOWN

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Back Pain					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361729-7Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501034A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 WK	Tinnitus					

Date:05/21/04ISR Number: 4361730-3Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501035A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK	Drug Ineffective Panic Attack					

Date:05/21/04ISR Number: 4361731-5Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0501037A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 19 DAY		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Convulsion Tongue Biting Tremor Urinary Incontinence					

Date:05/21/04ISR Number: 4361732-7Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501054A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4 DAY	Dry Mouth		Zoloft	C		

Date:05/21/04ISR Number: 4361733-9Report Type:Periodic  
Age:29 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501057A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3	MON		Vicodin	C		
				Soma	C		

Date:05/21/04ISR Number: 4361734-0Report Type:Periodic  
Age:14 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0501065A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
2	MON						

Date:05/21/04ISR Number: 4361735-2Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501069A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3	WK					
		Nausea					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361736-4Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501181A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6	MON					

Date:05/21/04ISR Number: 4361737-6Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501186A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Epistaxis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361738-8Report Type:Periodic  
Age:27 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501189A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharyngolaryngeal Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Wk					
		Rash					
		Stomach Discomfort					

Date:05/21/04ISR Number: 4361739-XReport Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501199A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day							
		Pharyngolaryngeal Pain					
		Rash					

Date:05/21/04ISR Number: 4361740-6Report Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501204A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	23 DAY			Allegra D	C		

Date:05/21/04ISR Number: 4361741-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0501211A  
 Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Lymphadenopathy		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG In the morning		Pain					
				Lexapro	SS		ORAL
10MG In the morning							

Date:05/21/04ISR Number: 4361742-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0501226A  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	7 DAY	Neck Pain		Neurontin	C		
		Tinnitus		Vicodin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361743-1Report Type:Periodic  
 Age:58 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501229A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
150MG Per day				Unknown	C		

Date:05/21/04ISR Number: 4361744-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501332A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
1 WK							

Date:05/21/04ISR Number: 4361745-5Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501342A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3 WK							

Date:05/21/04ISR Number: 4361746-7Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0501346A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Unknown							

Date:05/21/04ISR Number: 4361747-9Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501413A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Per day	2	DAY	Insomnia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Date:05/21/04ISR Number: 4361749-2Report Type:Periodic				Company Report #US-GLAXOSMITHKLINE-A0501420A			
Age: YR Gender: I/FU:I							
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Palpitations		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:05/21/04ISR Number: 4361750-9Report Type:Periodic				Company Report #US-GLAXOSMITHKLINE-A0501456A			
Age:21 YR Gender:Female I/FU:I							
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 2 MON							
		Palpitations					
		Weight Decreased					

Date:05/21/04ISR Number: 4361751-0Report Type:Periodic				Company Report #US-GLAXOSMITHKLINE-A0501599A			
Age:49 YR Gender:Male I/FU:I							
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Depression	Consumer	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 3 MON							
				Beer	SS		ORAL
10CANS Per							
day							
				Lexapro	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361752-2Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501605A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
7 DAY		Tongue Black Hairy		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361753-4Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501609A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	6 WK	Migraine		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Zyrtec-D	C		
				Armour Thyroid	C		

Date:05/21/04ISR Number: 4361754-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501613A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	6 WK	Amenorrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361755-8Report Type:Periodic  
Age:69 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0501615A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	4 DAY	Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK	Medication Error		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
				Diovan Hct	C		
				Allegra	C		
				Nasonex	C		
				Vioxx	C		
				Vitamins	C		

Date:05/21/04ISR Number: 4361756-XReport Type:Periodic  
Age:29 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501631A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Micturition Urgency		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6 DAY	Pollakiuria		Effexor	C		
		Urinary Incontinence					

Date:05/21/04ISR Number: 4361757-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501933A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 DAY						

Date:05/21/04ISR Number: 4361758-3Report Type:Periodic  
Age:52 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0501940A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK	Dyspnoea		Lexapro	C		
		Paraesthesia					
		Rash					
		Sleep Disorder					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361759-5Report Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501941A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	300MG Per day	1 YR		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Oral Contraceptives	C		

Date:05/21/04ISR Number: 4361760-1Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501943A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	300MG Per day	3 MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Seroquel	C		
				Klonopin	C		

Date:05/21/04ISR Number: 4361761-3Report Type:Periodic  
Age:70 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501954A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	300MG Per day	3 MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Norvasc	C		
				Digitex	C	Glaxosmithkline	
				Accupril	C		
				Aspirin	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361762-5Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501957A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Per day			Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361763-7Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501971A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK			Ambien	C		

Date:05/21/04ISR Number: 4361764-9Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501973A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hypoaesthesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK	Paraesthesia		Synthroid	C	Glaxosmithkline	
		Paraesthesia Oral		Oral Contraceptives	C		

Date:05/21/04ISR Number: 4361765-0Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501978A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Increased Appetite		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	5 WK	Weight Increased		Levoxyl	C	Glaxosmithkline	
				Verapamil	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361766-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502006A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nightmare		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361767-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502026A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthritis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
9 DAY		Hypersensitivity Oedema Peripheral Rash Pruritic Swelling Face Urticaria					

Date:05/21/04ISR Number: 4361770-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502030A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361771-6Report Type:Periodic  
Age:15 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502139A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK			Concerta	C		

Date:05/21/04ISR Number: 4361772-8Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502169A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG Per day	26 DAY	Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Nausea		K-Dur	C	Glaxosmithkline	
					Hctz	C		
					Endocet	C		
					Zocor	C		
					Xanax	C		
					Elavil	C	Glaxosmithkline	
					Flexeril	C		
					Valium	C		
					Albenza	C	Glaxosmithkline	
					Ultram	C		
					Tagamet	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361773-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0502176A  
Age:47 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG Per day	3 WK	Dysgeusia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	20MG Per day		Irritability		Paxil	C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361774-1Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502356A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Depression		Lexapro	SS		ORAL
20MG Per day	5 YR	Hangover					
		Nausea					
		Vertigo					

Date:05/21/04ISR Number: 4361775-3Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0502359A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	

Date:05/21/04ISR Number: 4361776-5Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502370A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	17 WK	Nervousness		Imipramine	C		
		Swelling		Klonopin	C		
				Hormone Replacement	C		
				Indocin	C		
				Neurontin	C		
				Prilosec	C		

Date:05/21/04ISR Number: 4361777-7Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502371A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	5 DAY						

Toprol Xl

C

Date:05/21/04ISR Number: 4361778-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502419A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2	WK						

Date:05/21/04ISR Number: 4361779-0Report Type:Periodic  
Age:52 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0502422A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300MG At Other night		Amnesia Confusional State Grand Mal Convulsion		Wellbutrin Xl Trileptal	PS C	Glaxosmithkline	ORAL ORAL
1MG At night				Risperidal	C		ORAL
25MG In the morning				Paxil Cr	C	Glaxosmithkline	ORAL
.1MG In the morning				Synthroid	C	Glaxosmithkline	ORAL
2MG Per day				Estrace	C		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361780-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502423A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2	MON						

Date:05/21/04ISR Number: 4361781-9Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502468A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Deafness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other							

Date:05/21/04ISR Number: 4361782-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502469A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other							

Date:05/21/04ISR Number: 4361783-2Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502562A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3	WK	Anxiety Tension					

Date:05/21/04ISR Number: 4361784-4Report Type:Periodic  
 Age:42 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0502565A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	19 DAY						

Fatigue  
Somnolence

Atenolol  
Nexium

C  
C

Date:05/21/04ISR Number: 4361785-6Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502573A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3	MON					

Date:05/21/04ISR Number: 4361786-8Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502577A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day	4	WK		Benadryl	SS	Glaxosmithkline	
UNKNOWN		Dysphagia					
		Insomnia		Lipitor	C		
				Antihypertensive	C		
				Arthritis Medication	C		ORAL
				Flonase	C	Glaxosmithkline	



300MG Per day	8	WK	Anxiety	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Mouth Ulceration	Valtrex	C	Glaxosmithkline	
			Swollen Tongue	Oral Contraceptives	C		
			Therapeutic Response				
			Unexpected				

Date:05/21/04ISR Number: 4361792-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0502852A  
 Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	4	WK	Increased Appetite				
			Weight Increased	Prozac	C		
				Topamax	C		
				Motrin	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361793-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0502854A  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4	DAY	Rash Generalised				
				Guaifenesin	C		
				Albuterol	C	Glaxosmithkline	
				Advil	C	Glaxosmithkline	



1	WK	Insomnia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Libido Disorder	Paxil	SS	Glaxosmithkline	ORAL
		Nervousness				

Date:05/21/04ISR Number: 4361801-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0502929A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Wellbutrin Sr	SS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361802-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0502960A  
 Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	9 WK			Vitamins	C		
				Aspirin	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361803-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0502969A  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK			Paxil	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Trileptal C  
Lorazepam C

Date:05/21/04ISR Number: 4361804-7Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502983A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK			Prozac	C		
				Neurontin	C		
				Ativan	C		
				Ambien	C		

Date:05/21/04ISR Number: 4361805-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502989A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Feeling Jittery					

Date:05/21/04ISR Number: 4361806-0Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502993A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Crying		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	23 DAY	Dizziness		Tylenol Sinus	C		
		Fatigue					
		Irritability					
		Nausea					
		Vertigo					

Date:05/21/04ISR Number: 4361807-2Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0503004A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash Vesicular		Wellbutrin Xl	PS	Glaxosmithkline	

Date:05/21/04ISR Number: 4361808-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0503011A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3 DAY	Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361809-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0503014A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	4 WK	Arthralgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Joint Stiffness		None	C		
		Myalgia					
		Oedema Peripheral					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361813-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503115A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Skin Discolouration		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361814-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503116A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361815-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503160A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abnormal Behaviour Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361816-3Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503199A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG Per day	7 WK	Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361817-5Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0503214A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 4 DAY Initial or Prolonged		Confusional State Convulsion Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361818-7Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503225A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361819-9Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503228A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK	Dizziness					
		Irritability					

Date:05/21/04ISR Number: 4361820-5Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503230A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Wellbutrin Sr	SS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361821-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503346A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	1TAB Twice	Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	per day	MON					

Date:05/21/04ISR Number: 4361822-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503440A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
	30 DAY						

Date:05/21/04ISR Number: 4361823-0Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503453A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361824-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503464A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	7 MON	Libido Decreased					
		Paranoia					

Date:05/21/04ISR Number: 4361825-4Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503465A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 MON			Birth Control	C		

Date:05/21/04ISR Number: 4361826-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0503493A  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
6 DAY							

Date:05/21/04ISR Number: 4361827-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0503530A  
 Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	

Date:05/21/04ISR Number: 4361828-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0503540A  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Wellbutrin Ultram	SS C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361829-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503546A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	2	MON					

Date:05/21/04ISR Number: 4361830-8Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503818A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Stool Analysis Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	4	DAY					

Date:05/21/04ISR Number: 4361831-XReport Type:Periodic  
Age:27 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0503840A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3	WK					
		Anxiety		Paxil Cr	SS	Glaxosmithkline	ORAL
25MG Per day	3	WK					
		Erectile Dysfunction		Hydroxyzine Pamoate	C		
				Trazodone	C		
				Vistaril	C		
				Percocet	C		
				Oxycontin	C		

Date:05/21/04ISR Number: 4361832-1Report Type:Periodic  
Age:63 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503853A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Throat		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6	WK					
		Hyperhidrosis		Ativan	C		

Thermal Burn  
Tremor

Diovan  
Synthroid  
Amitriptyline

C  
C  
C

Glaxosmithkline

Date:05/21/04ISR Number: 4361833-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503859A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chills		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	16 DAY	Diarrhoea		Asthma Medication	C		
		Influenza Like Illness					
		Oral Mucosal Blistering					
		Pain					
		Pyrexia					
		Rash					
		Swelling Face					
		Tongue Disorder					

Date:05/21/04ISR Number: 4361834-5Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503862A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Fluctuation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL



Pollakiuria  
 300MG Per day 15 DAY

Date:05/21/04ISR Number: 4361840-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504082A  
 Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Migraine		Wellbutrin	PS	Glaxosmithkline	

Date:05/21/04ISR Number: 4361841-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504084A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day Initial or Prolonged		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361842-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504091A  
 Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Insomnia		Wellbutrin	PS	Glaxosmithkline	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361843-6Report Type:Periodic  
 Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504099A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Twice		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 WK						
				Synthroid	C	Glaxosmithkline	
				Oxycontin	C		
				Amitriptyline	C		
				Xanax	C		
				Soma	C		

Date:05/21/04ISR Number: 4361844-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504206A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
1 WK		Sleep Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361845-XReport Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504207A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361846-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504216A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
6 MON		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Skin Exfoliation					

Date:05/21/04ISR Number: 4361847-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504288A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2	MON	Anger					

Date:05/21/04ISR Number: 4361848-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504293A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Lexapro	C		
				Topamax	C		

Date:05/21/04ISR Number: 4361849-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504297A  
Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK			Oral Contraceptives	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361850-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504303A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 WK	Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hyperhidrosis		Colace	C		
		Nausea		Serzone	C		
		Tremor					

Date:05/21/04ISR Number: 4361851-5Report Type:Periodic  
 Age:17 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504309A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 WK		Hypersensitivity		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Rash					

Date:05/21/04ISR Number: 4361852-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504317A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 MON	Disturbance In Attention		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Nervousness		Vitamins	C		

Date:05/21/04ISR Number: 4361853-9Report Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504320A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	45 DAY	Dysmenorrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Mood Altered		Celexa	C		
				Zyrtec	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361854-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504324A  
Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Discomfort Hot Flush		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361855-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504330A  
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day	4 MON			Provigil	C		
				Topamax	C		
				Hytrin	C		
				Levoxyl	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361856-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504334A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361857-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504455A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361858-8Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504561A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	5 WK	Drug Ineffective		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
150MG Twice		Feeling Of Despair					
per day				Birth Control Pill	C		

Date:05/21/04ISR Number: 4361859-XReport Type:Periodic  
Age:68 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504563A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hair Texture Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK						

Date:05/21/04ISR Number: 4361860-6Report Type:Periodic  
Age:72 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504572A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	12 DAY	Insomnia		Xanax	C		
		Nausea					

Date:05/21/04ISR Number: 4361861-8Report Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504577A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anger		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	0 DAY	Dyspnoea		Hytrin	C		
		Nervousness		Triavil	C		
				Mevacor	C		
				Valium	C		
				Cipro	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361865-5Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504586A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Joint Swelling		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK	Urticaria					

Date:05/21/04ISR Number: 4361866-7Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504597A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Jittery		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Somnolence		Maxalt	C		
				Flonase	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361867-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504677A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG	Unknown						

Date:05/21/04ISR Number: 4361868-0Report Type:Periodic  
Age: Gender:I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504704A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361869-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504705A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Pruritic		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG	Per day 3 WK						

Date:05/21/04ISR Number: 4361870-9Report Type:Periodic  
Age:75 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504781A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG	Per day 5 DAY			Restoril	C		
				Hctz	C		
				Lipitor	C		
				Captopril	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361871-0Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504791A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	7 DAY	Swelling		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Urticaria		Oral Contraceptive	C		
				Hydroxyzine	C		
				Prednisone	C		

Date:05/21/04ISR Number: 4361872-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504793A  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				No Concurrent Medication	C		

Date:05/21/04ISR Number: 4361873-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504799A  
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3 DAY	Nasopharyngitis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Pharyngolaryngeal Pain		Levoxyl	C	Glaxosmithkline	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361874-6Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504807A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Neck Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Wellbutrin Sr	C	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361875-8Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0504824A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361876-XReport Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504871A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other	300MG Per day 4 MON						

Date:05/21/04ISR Number: 4361877-1Report Type:Periodic  
 Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504874A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Behaviour		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2 DAY		Hallucination Hypertension Insomnia Night Sweats Paranoia Thinking Abnormal Tremor		Antibiotic	C		

Date:05/21/04ISR Number: 4361878-3Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504876A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Balance Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	17 DAY	Headache		Nexium	C		
		Lethargy		Ibuprofen	C	Glaxosmithkline	
		Mood Altered		Tylenol	C	Glaxosmithkline	
		Nausea		Pseudoephedrine	C		
		Palpitations		Tums	C	Glaxosmithkline	
		Tremor					

Date:05/21/04ISR Number: 4361879-5Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0504881A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361880-1Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504891A

Outcome	PT
	Drug Ineffective
	Dry Mouth

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Hepatic Enzyme Increased				
		Insomnia				
Dose	Duration	Memory Impairment	Report Source	Product	Role	Manufacturer
150MG Per day		Pharmaceutical Product		Wellbutrin Xl	PS	Glaxosmithkline
		Complaint		Wellbutrin Sr	SS	Glaxosmithkline
				Klonopin	C	
				Soy Supplement	C	
				Vitamins	C	

Date:05/21/04ISR Number: 4361881-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504895A  
 Age: YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer
Outcome	Duration	Agitation		Wellbutrin Xl	PS	Glaxosmithkline
Dose	150MG Per day 2 WK	Insomnia				ORAL

Date:05/21/04ISR Number: 4361882-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504896A  
 Age:32 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer
Outcome	Duration	Anxiety		Wellbutrin Xl	PS	Glaxosmithkline
Dose	150MG Per day	Dizziness		Coumadin	C	Glaxosmithkline
		Drug Ineffective		Amoxicillin	C	Glaxosmithkline
		Dyspnoea				
		Pharyngitis Streptococcal				

Date:05/21/04ISR Number: 4361883-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504915A  
 Age:44 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer
Outcome	Duration	Agitation		Wellbutrin Xl	PS	Glaxosmithkline
Dose	150MG Per day 2 WK	Anorexia		Darvocet	C	
				Plaquenil	C	

Date:05/21/04ISR Number: 4361884-9Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504919A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day	1	MON		Adderall	C		
				Xanax	C		
				Seroquel	C		

Date:05/21/04ISR Number: 4361885-0Report Type:Periodic  
Age:47 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0505025A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Macular		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other		Rash Vesicular		Ritalin	C		ORAL
150MG Unknown	7	DAY					
10MG Three		Skin Exfoliation					
times per day		Stevens-Johnson Syndrome					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361886-2Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0505093A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:05/21/04ISR Number: 4361887-4Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505107A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Myalgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK	Nausea		Augmentin	C	Glaxosmithkline	
		Tinnitus		Zyrtec	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361888-6Report Type:Periodic  
Age:33 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0505141A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lymphadenopathy		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK	Throat Tightness		Vitamins	C		
				Wellbutrin Sr	C	Glaxosmithkline	
150MG Per day	1 YR						

Date:05/21/04ISR Number: 4361889-8Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0505227A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:05/21/04ISR Number: 4361890-4Report Type:Periodic  
Age:65 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0505228A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Lexapro	C		
				Atenolol	C		

Date:05/21/04ISR Number: 4361891-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505229A  
 Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Confusional State		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 14 DAY		Convulsion		Allegra	C		
Initial or Prolonged		Disorientation					
		Tremor					

Date:05/21/04ISR Number: 4361892-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505238A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other							
2 MON							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361893-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505258A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
8	WK						

Date:05/21/04ISR Number: 4361894-1Report Type:Periodic  
 Age:52 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505259A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysuria		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Hytrin	C		
				Proscar	C		

Date:05/21/04ISR Number: 4361896-5Report Type:Periodic  
 Age: Gender:I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505260A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361898-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505261A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Affect Lability		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Unknown	4 DAY	Drug Ineffective					
		Irritability					
		Paraesthesia					

Date:05/21/04ISR Number: 4361909-0Report Type:Periodic  
 Age:42 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0441775A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG	Per day 2 WK	Amnesia	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
			Convulsion	Professional	Celexa	C		
			Epistaxis					
			Fall					
			Periorbital Haematoma					
			Tooth Fracture					
			Tooth Injury					
			Toothache					

Date:05/21/04ISR Number: 4361910-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0490957A  
 Age:27 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Drug Ineffective	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
			Pharmaceutical Product Complaint	Professional				

Date:05/21/04ISR Number: 4361911-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0491689A  
 Age:36 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG	See dosage text 3 WK	Convulsion	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
				Professional				

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Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361912-0Report Type:Periodic  
Age:52 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0491733A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	4 DAY	Health	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other		Dyspnoea Paraesthesia Rash Stomach Discomfort	Professional				

Date:05/21/04ISR Number: 4361913-2Report Type:Periodic  
Age:19 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0492592A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	450MG Per day	2 MON	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Amnesia Grand Mal Convulsion Tongue Biting	Professional	Ritalin Sr	C		

Date:05/21/04ISR Number: 4361914-4Report Type:Periodic  
Age:25 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0493257A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	450MG Per day	112 DAY	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Other	Hospitalization - Initial or Prolonged 20MG Per day	Grand Mal Convulsion	Professional	Lexapro	C		ORAL
				Norvir Invirase	C C		ORAL

Date:05/21/04ISR Number: 4361915-6Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0494685A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Amnesia Grand Mal Convulsion Incontinence	Professional				

Tongue Biting

Date:05/21/04ISR Number: 4361916-8Report Type:Periodic  
 Age:18 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0495501A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 22 DAY		Disorientation	Health	Wellbutrin	PS	Glaxosmithkline	
Initial or Prolonged		Fatigue	Professional	Advair	C	Glaxosmithkline	
		Grand Mal Convulsion		Singulair	C		
				Alesse	C		
				Zyrtec	C	Glaxosmithkline	
				Pulmicort	C		
				Proventil	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361917-XReport Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0495775A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG Per day		Hallucination	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
			Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361918-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496051A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Incontinence		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK	Pollakiuria					

Date:05/21/04ISR Number: 4361919-3Report Type:Periodic  
 Age:71 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496060A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	6 DAY			Toprol Xl	C		
				Norvasc	C		
				Accupril	C		
				Metformin	C		
				Lipitor	C		
				Amaryl	C		
				Xalatan	C		

Date:05/21/04ISR Number: 4361920-XReport Type:Periodic  
 Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496065A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4 WK	Dry Mouth		Zithromax	C		
		Headache					
		Nausea					

Date:05/21/04ISR Number: 4361921-1Report Type:Periodic  
 Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496300A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Fatigue  
Tremor

Date:05/21/04ISR Number: 4361922-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496303A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Health Professional	Wellbutrin Xl Proton Pump Inhibitor	PS SS	Glaxosmithkline	ORAL
UNKNOWN							

Date:05/21/04ISR Number: 4361923-5Report Type:Periodic  
Age:75 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496304A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	0 DAY			Celexa Xanax	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361924-7Report Type:Periodic  
Age:48 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496309A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 MON	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Constipation		Lasix	C	Glaxosmithkline	
		Sexual Dysfunction		Buspar	C		

Date:05/21/04ISR Number: 4361925-9Report Type:Periodic  
Age:16 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496315A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Peripheral Coldness					

Date:05/21/04ISR Number: 4361929-6Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496324A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 MON		Migraine		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361930-2Report Type:Periodic  
Age:36 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496336A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	6 DAY	Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Obsessive-Compulsive Disorder		Lexapro	C		
				Trazodone	C		
				Birth Control Pill	C		

Date:05/21/04ISR Number: 4361931-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0496665A  
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	26 DAY	Bursitis		Prevacid	C		
		Joint Swelling					
		Oedema Peripheral					

Date:05/21/04ISR Number: 4361932-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0496670A  
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pollakiuria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2 WK							

Date:05/21/04ISR Number: 4361933-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0496672A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Life-Threatening		Pharmaceutical Product					
300MG Per day	3 WK	Complaint					
Hospitalization -							
Initial or Prolonged							
Other							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361935-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496886A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Unknown 1	MON	Drug Ineffective	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361936-3Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496899A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day		Dizziness	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361937-5Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496901A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day		Nausea	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361938-7Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496903A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Adverse Event	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Apathy	Wellbutrin Sr	SS	Glaxosmithkline	ORAL
			Sexual Dysfunction				

Date:05/21/04ISR Number: 4361939-9Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496912A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Drug Ineffective	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361940-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496933A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Paraesthesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Weight Decreased					

Date:05/21/04ISR Number: 4361941-7Report Type:Periodic  
Age:25 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496953A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Suppressed Lactation		Wellbutrin	PS	Glaxosmithkline	ORAL
5 MON							

Date:05/21/04ISR Number: 4361942-9Report Type:Periodic  
Age:51 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496956A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Rash Pruritic Urticaria					



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Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361943-0Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496958A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pollakiuria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361944-2Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496959A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361945-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496961A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urine Amphetamine Positive		Wellbutrin Tylenol	PS C	Glaxosmithkline Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361946-6Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496988A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urticaria		Wellbutrin Adderall	PS C	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361947-8Report Type:Periodic  
 Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497062A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Drug Ineffective		Wellbutrin Xl Effexor	PS C	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361948-XReport Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497070A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Blood Glucose Increased	Wellbutrin Xl Glipizide	PS C	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361949-1Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497071A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Headache	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	5 WK		Nausea				

Date:05/21/04ISR Number: 4361951-XReport Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497075A

Outcome	PT
	Anorexia
	Chest Discomfort
	Feeling Jittery
	Heart Rate Increased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3 WK	Nausea Palpitations Retching		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
37.5MG Per day	3 WK	Somnolence Tinnitus Tremor		Effexor	SS		
				Valium	C		

Date:05/21/04ISR Number: 4361952-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497099A  
 Age:35 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	5 MON	Blood Pressure Increased Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361953-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497101A  
 Age:35 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	8 MON	Agitation Insomnia Libido Increased Smoker Weight Loss Poor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Vicodin	C		
				Nasonex	C		
				Allegra	C		

Date:05/21/04ISR Number: 4361954-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497105A  
 Age:31 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	10 DAY	Chest Discomfort Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Aciphex	C		

Dry Mouth  
Dyspnoea  
Headache  
Neck Pain  
Pain In Jaw

Yasmin C  
Nasacort Aq C  
Amoxicillin C Glaxosmithkline

Date:05/21/04ISR Number: 4361955-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497173A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eczema		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361956-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497175A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Papular		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361957-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497193A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flushing		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Urticaria		Combivent	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361958-2Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497198A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Urticaria		Benadryl	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4361959-4Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497215A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:05/21/04ISR Number: 4361960-0Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0497346A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Interaction		Strattera	SS		ORAL

Date:05/21/04ISR Number: 4361961-2Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0497367A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tic		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice				Wellbutrin Sr	C	Glaxosmithkline	ORAL
per day							

Date:05/21/04ISR Number: 4361962-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0497369A

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361963-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497376A  
Age: Gender: I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG	Per day 1 WK						

Date:05/21/04ISR Number: 4361964-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497538A  
Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Heart Rate Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
2	DAY	Initial Insomnia					

Date:05/21/04ISR Number: 4361965-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497561A  
Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG	Unknown	Nausea					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361966-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497582A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3 WK		Suicidal Ideation					

Date:05/21/04ISR Number: 4361967-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497585A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Spasms Muscle Twitching		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361968-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497588A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 2 WK		Flushing		Wellbutrin Sr	C	Glaxosmithkline	ORAL
150MG Twice per day 3 MON		Formication					

Date:05/21/04ISR Number: 4361969-7Report Type:Periodic  
Age:32 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0497602A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG per day 7 DAY		Drug Ineffective Feeling Jittery Suicidal Ideation		Paxil Cr	C	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361970-3Report Type:Periodic  
Age:59 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0497607A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day		Decreased Appetite		Xanax	C		
		Dry Mouth		Ambien	C		
		Nausea		Duragesic	C		
		Tachycardia		Oxycodone	C		

Date:05/21/04ISR Number: 4361971-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497719A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Foreign Body Trauma		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361973-9Report Type:Periodic  
Age:12 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0497721A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK						



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361974-0Report Type:Periodic  
Age:8 YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0497726A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
25MG Per day		Drug Ineffective		Strattera	C		
		Grand Mal Convulsion					
		Tongue Biting					

Date:05/21/04ISR Number: 4361975-2Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497728A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	4 WK	Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Xanax	C		

Date:05/21/04ISR Number: 4361976-4Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0497732A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				Lexapro	C		
				Topamax	C		

Date:05/21/04ISR Number: 4361977-6Report Type:Periodic  
Age:32 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497734A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	1 MON	Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Palpitations		Remeron	C		
		Paranoia		Buspar	C		

Date:05/21/04ISR Number: 4361978-8Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497736A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Ultram	C		

Date:05/21/04ISR Number: 4361979-XReport Type:Periodic  
Age:10 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497742A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Muscle Twitching Tremor					

Date:05/21/04ISR Number: 4361980-6Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497749A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Hypertension Tachycardia		None	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361981-8Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497759A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hypertension		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4	MON		Librax	C		
				Microzide	C		

Date:05/21/04ISR Number: 4361982-XReport Type:Periodic  
Age:55 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497764A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day	1	WK		Neurontin	C		
		Nausea		Celexa	C		
				Pravachol	C		
				Celebrex	C		

Date:05/21/04ISR Number: 4361983-1Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497791A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	

Date:05/21/04ISR Number: 4361984-3Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497799A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	
		Insomnia					

Date:05/21/04ISR Number: 4361985-5Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497807A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	4	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Anxiety					
		Dysgeusia		Buspar	C		
		Stomach Discomfort		Eskalith Cr	C	Glaxosmithkline	
		Tinnitus		Metformin	C		
				Vitamins	C		
				Calcium	C		
				Premarin	C		
				Unspecified			
				Medication	C		
				Prilosec	C		
				Lipitor	C		
				Klonopin	C		

Date:05/21/04ISR Number: 4361986-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497816A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Rash					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361987-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497849A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dissociation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dizziness		Zyban	SS	Glaxosmithkline	ORAL
		Insomnia					

Date:05/21/04ISR Number: 4361988-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497890A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Unknown	DAY	Pharyngolaryngeal Pain					

Date:05/21/04ISR Number: 4361989-2Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497971A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 1	MON			Climara	C		

Date:05/21/04ISR Number: 4361990-9Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498001A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 2	WK						

Date:05/21/04ISR Number: 4361991-0Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498007A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Adverse Event		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:05/21/04ISR Number: 4361992-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498010A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Wellbutrin	PS	Glaxosmithkline	
4	WK			Ambien	C		
				Celexa	C		
				Zestril	C		
				Diuretic	C		
				Lipitor	C		
				Tylenol With Codeine	C		

Date:05/21/04ISR Number: 4361993-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498016A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety		Wellbutrin	PS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4361994-6Report Type:Periodic  
 Age:35 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0498025A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	6 DAY	Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
		Heart Rate Increased					

Date:05/21/04ISR Number: 4361995-8Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498026A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4361996-XReport Type:Periodic  
 Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498027A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown		Chills		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN	30MG Unknown	Dizziness		Celexa	SS		
		Dry Mouth					
		Fatigue					
		Feeling Abnormal					
		Hyperhidrosis					
		Insomnia					
		Irritability					
		Libido Decreased					
		Migraine					
		Myalgia					
		Nausea					
		Vision Blurred					

Date:05/21/04ISR Number: 4362001-1Report Type:Periodic  
 Age:8 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498029A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
35	DAY						

Date:05/21/04ISR Number: 4362002-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498041A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hypersensitivity					

Date:05/21/04ISR Number: 4362003-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498153A  
 Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3 MON	Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Nausea		Lamictal	C	Glaxosmithkline	
				Lorazepam	C		
				Aspirin	C	Glaxosmithkline	
				Excedrin	C		
				Pepto-Bismol	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4362004-7Report Type:Periodic  
Age:52 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0498156A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	12 DAY	Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Alopecia		Relpax	C		
				Dhe	C		
				Depakote	C		

Date:05/21/04ISR Number: 4362005-9Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498161A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Wellbutrin Sr	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4362006-0Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498165A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dry Mouth		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362007-2Report Type:Periodic  
Age:34 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0498201A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG In the							
Initial or Prolonged							
morning	135 DAY			Xanax	C		
.5MG Three							
times per day				Herbal Medicine	C		ORAL

Date:05/21/04ISR Number: 4362008-4Report Type:Periodic  
Age:18 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498347A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK						

Date:05/21/04ISR Number: 4362009-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498349A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Orthostatic Hypotension		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Lexapro	C		

Date:05/21/04ISR Number: 4362010-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498361A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Unknown		Chest Discomfort		Effexor	SS		
UNKNOWN		Dyspnoea					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4362011-4Report Type:Periodic  
 Age:45 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498507A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3 DAY	Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Nausea					

Date:05/21/04ISR Number: 4362012-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498508A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	5 MON	Heart Rate Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Pruritus Generalised		Celexa	C		
		Rash					

Date:05/21/04ISR Number: 4362013-8Report Type:Periodic  
 Age:14 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498516A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Antisocial Personality		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Disorder		None	C		
		Crying					
		Depressed Mood					
		Dissociation					
		Pallor					

Date:05/21/04ISR Number: 4362014-XReport Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498519A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362015-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498524A  
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash		Wellbutrin	PS	Glaxosmithkline	
2 WK				Effexor Xr	C		

Date:05/21/04ISR Number: 4362017-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498527A  
Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Aura		Effexor Xr	C		
150MG Per day		Fatigue		Trazodone	C		
25MG As		Grand Mal Convulsion					
required							

Date:05/21/04ISR Number: 4362018-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498540A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4362019-9Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498544A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Arthralgia Joint Stiffness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362020-5Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498545A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Swelling Face		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362021-7Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498554A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Advair	C	Glaxosmithkline	
				Benadryl	C	Glaxosmithkline	
				Ortho Tri-Cyclen	C		

Date:05/21/04ISR Number: 4362022-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498602A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
6 MON		Pruritus Generalised		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Zyprexa	C		

Date:05/21/04ISR Number: 4362023-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498614A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362024-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498737A  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150TAB Per		Tremor		Prilosec	C		
day	1 DAY			Klonopin	C		

Date:05/21/04ISR Number: 4362025-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498767A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Glucophage	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4362026-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498777A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day 2	MON	Abdominal Pain Upper	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Nausea	Lamictal	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4362027-8Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498781A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Nausea	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362028-XReport Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0498792A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	5 MON		Alopecia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Depression	Neurontin	C		
			Hypotrichosis				

Date:05/21/04ISR Number: 4362029-1Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498842A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Hospitalization - Initial or Prolonged Other		Cerebrovascular Accident Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362030-8Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499053A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lymphadenopathy		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362031-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499084A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362032-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499202A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362033-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499205A  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3	MON					





300MG Per day 3 WK	Crying	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	Dizziness	Ortho-Novum 777	C		
	Nausea				
	Salivary Hypersecretion				
	Throat Tightness				

Date:05/21/04ISR Number: 4362039-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499593A  
 Age:54 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
300MG Per day 3 WK	Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	Pruritus		Evista	C		
	Urticaria		Caltrate	C	Glaxosmithkline	
			Multivitamin	C		

Date:05/21/04ISR Number: 4362040-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499597A  
 Age:50 YR Gender:Male I/FU:F

Outcome	PT
	Anorexia
	Headache
	Hypersomnia
	Nausea
	Neck Pain

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pain

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	36 DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Lipitor	C		

Date:05/21/04ISR Number: 4362041-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499599A  
 Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Orthostatic Hypotension		Wellbutrin Xl	PS	Glaxosmithkline	

Date:05/21/04ISR Number: 4362042-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499610A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 DAY						

Date:05/21/04ISR Number: 4362043-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499624A  
 Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362044-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499641A  
 Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Galactorrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2 WK							

Date:05/21/04ISR Number: 4362045-XReport Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499813A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4	MON		Lorazepam	C		

Date:05/21/04ISR Number: 4362046-1Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499818A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Somnolence		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6	MON		Zoloft	C		

Date:05/21/04ISR Number: 4362047-3Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499830A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1	YR		Lamictal	SS	Glaxosmithkline	ORAL
3	WK	Rash Generalised		Valproic Acid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4362048-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499836A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3 DAY			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Medication Error					
		Stomach Discomfort					
		Vertigo					

Date:05/21/04ISR Number: 4362049-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499846A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation	Consumer	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Impatience		Caffeinated Beverage	SS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362050-3Report Type:Periodic  
Age:60 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0499853A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
450MG Per day	3 DAY			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Blood Pressure Increased	Health				
			Professional	Premarin	C		
				Synthroid	C	Glaxosmithkline	
				Meridia	C		
				Klonopin	C		
				Hyzaar	C		

Date:05/21/04ISR Number: 4362051-5Report Type:Periodic  
Age:26 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499854A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	6 DAY			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Heart Rate Increased					
		Insomnia		Nexium	C		
		Tremor		Lexapro	C		
				Depakote	C		

Date:05/21/04ISR Number: 4362052-7Report Type:Periodic  
Age:11 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499868A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accidental Exposure		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Agitation		Nortriptyline	C		
		Palpitations		Concerta	C		
				Depakote	C		
				Ritalin	C		
				Clonidine	C		

Date:05/21/04ISR Number: 4362053-9Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500113A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	16 DAY						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4362054-0Report Type:Periodic  
Age:45 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0500115A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
600MG Per day	5 DAY	Tremor		Paxil	C	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362055-2Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500125A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Adderall	C		

Date:05/21/04ISR Number: 4362056-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500143A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK	Palpitations		Effexor	C		

Date:05/21/04ISR Number: 4362057-6Report Type:Periodic  
Age:14 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500146A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 MON	Feeling Jittery Mood Swings					

Date:05/21/04ISR Number: 4362058-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500149A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK						

Date:05/21/04ISR Number: 4362060-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0500152A  
 Age:12 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK						
		Somnolence		Wellbutrin	SS	Glaxosmithkline	ORAL
100MG Per day							
				Strattera	C		
				Vitamins	C		

Date:05/21/04ISR Number: 4362061-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0500181A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4362062-XReport Type:Periodic  
Age:36 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0500182A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG In the morning							

Date:05/21/04ISR Number: 4362063-1Report Type:Periodic  
Age:82 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500360A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Per day							
		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Blood Pressure Increased		Lexapro	SS		ORAL
		Decreased Appetite		Aspirin	C	Glaxosmithkline	
		Drug Ineffective		Zocor	C		
		Stress		Prilosec	C		
		Weight Decreased					

Date:05/21/04ISR Number: 4362064-3Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500361A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
450MG Per day 2 MON							
		Yawning		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Provigil	C		
				Oral Contraceptive	C		

Date:05/21/04ISR Number: 4362065-5Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500362A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Swelling		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362066-7Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500365A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	6 DAY			Levoxyl	C	Glaxosmithkline	

Date:05/21/04ISR Number: 4362067-9Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500366A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK			Asacol	C	Glaxosmithkline	
				Protonix	C		
				Birth Control	C		

Date:05/21/04ISR Number: 4362069-2Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500367A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fall		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -	2 WK	Grand Mal Convulsion		Glucophage	C		
Initial or Prolonged		Loss Of Consciousness					
500MG Twice		Tongue Biting		Zantac	C	Glaxosmithkline	
per day							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4362071-0Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500372A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362072-2Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500378A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	

Date:05/21/04ISR Number: 4362073-4Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500387A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362074-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500443A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362075-8Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0500562A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash Generalised		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362076-XReport Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0500607A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphasia		Wellbutrin Xl Multiple Medications	PS C	Glaxosmithkline	ORAL

Date:05/21/04ISR Number: 4362077-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0500617A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice		Crying		Wellbutrin Sr	C	Glaxosmithkline	ORAL
per day	3 YR						

Date:05/21/04ISR Number: 4362080-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0500637A  
 Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1 MON	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Anxiety		Lexapro	C		
		Therapeutic Response		Xanax	C		
		Unexpected		Vaseretic	C		
				Zocor	C		
				Ibuprofen	C	Glaxosmithkline	
800MG Unknown							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

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Date:05/21/04ISR Number: 4362081-3Report Type:Periodic  
Age:49 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLIN-A0500669A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	1 WK	Balance Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Crepitations		Synthroid	C	Glaxosmithkline	
		Gait Disturbance		Lasix	C	Glaxosmithkline	
		Tremor		Atenolol	C		
				Imuran	C	Glaxosmithkline	
				Nexium	C		
				Prednisone	C		

Date:05/21/04ISR Number: 4362082-5Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLIN-A0500671A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice		Rash		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
per day	MON	Urticaria					
		Weight Decreased		Vitamins	C		
				Calcium	C		

Date:05/21/04ISR Number: 4362083-7Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLIN-A0500676A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	1 YR	Somnolence		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Tinnitus		Bupropion Sr	SS	Glaxosmithkline	ORAL
				Paxil	SS	Glaxosmithkline	ORAL
				Diovan 80	C		
				Vitamins	C		

Date:05/21/04ISR Number: 4362084-9Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500677A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	8 DAY						

Date:05/21/04ISR Number: 4362085-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500679A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anaphylactic Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
3 WK							

Date:05/21/04ISR Number: 4362086-2Report Type:Periodic  
Age:55 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500694A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 MON						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/04ISR Number: 4362087-4Report Type:Periodic  
 Age:31 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500737A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3	MON	Weight Increased	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
UNKNOWN				Lexapro	C		
						10MG Unknown	

Date:05/21/04ISR Number: 4362444-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511243A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death							
			Completed Suicide Depressed Mood Suicidal Ideation	Wellbutrin Sr	PS	Glaxosmithkline	

Date:05/24/04ISR Number: 4363402-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511608A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day			Medication Error	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/24/04ISR Number: 4363403-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511715A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day Initial or Prolonged							
			Aphasia  Cerebrovascular Accident Transient Ischaemic Attack	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/24/04ISR Number: 4363404-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511742A  
Age:17 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 0 DAY		Intentional Misuse		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Self-Medication					

Date:05/24/04ISR Number: 4363406-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511755A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day	4 DAY	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
		Medication Error		Lexapro Risperdal	C C		

Date:05/24/04ISR Number: 4363418-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0323573A  
Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day	MON	Visual Acuity Reduced	Health Professional	Zyban	PS	Glaxosmithkline	ORAL
2UNIT Three times per day				Euphytose	C		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/24/04ISR Number: 4363421-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0330307A

Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Myopia	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG See		Visual Acuity Reduced	Professional				
dosage text	3	WK					

Date:05/24/04ISR Number: 4363427-2Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0331481A

Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Interaction	Professional				
per day	30	DAY		Tenoretic	SS		
		Face Injury					
		Hypotension					
		Loss Of Consciousness					
		Syncope Vasovagal					
		Transient Ischaemic					
		Attack					

Date:05/24/04ISR Number: 4363440-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0332749A

Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	7	DAY		Propranolol	C		ORAL
80MG Per day		Gingival Pain					
		Heart Rate Increased		Zopiclone	C		ORAL
3.75MG As		Insomnia					
required		Tinnitus					

Date:05/24/04ISR Number: 4363444-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0333138A  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Mouth Ulceration		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Rash Vesicular					
per day	21	DAY					
50MG Three		Vasculitis		Diclofenac	C		ORAL
times per day	5	DAY					

Date:05/25/04ISR Number: 4364384-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0493781A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Weight Loss Poor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/25/04ISR Number: 4364385-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0494320A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/25/04ISR Number: 4364386-9Report Type:Periodic  
 Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0500121A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Unspecified Drugs	C		

Date:05/25/04ISR Number: 4364435-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508981A  
 Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Ineffective					
per day	1 MON	Medication Error					
		Somnolence					

Date:05/25/04ISR Number: 4364437-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511270A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Pharmaceutical Product					
per day		Complaint					

Date:05/25/04ISR Number: 4364438-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511391A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abortion Spontaneous		Wellbutrin	PS	Glaxosmithkline	
Other							

Date:05/25/04ISR Number: 4364444-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511866A  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Atrial Fibrillation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	64 DAY			Paxil	C	Glaxosmithkline	ORAL
20MG Per day				Synthroid	C	Glaxosmithkline	
UNKNOWN				Zocor	C		
UNKNOWN				Tricor	C		
UNKNOWN				Allegra	C		
UNKNOWN							

Date:05/25/04ISR Number: 4364448-6Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0321501A  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Parkinson'S Disease		Zyntabac	PS	Glaxosmithkline	
19 DAY							

Date:05/25/04ISR Number: 4364454-1Report Type:Expedited (15-DaCompany Report #HU-GLAXOSMITHKLINE-B0328691A  
Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Hyperaemia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
15 DAY							
Initial or Prolonged		Oedema Pruritus Rash Erythematous Rash Maculo-Papular					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/25/04ISR Number: 4364458-9Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0331807A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blindness		Quomem	PS	Glaxosmithkline	
		Conjunctivitis		Zyntabac	C	Glaxosmithkline	

Date:05/25/04ISR Number: 4364465-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0332761A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Aggression		Zyban	PS	Glaxosmithkline	ORAL
		Amnesia		Alcohol	SS		
		Drug Interaction		Zoloft	C		
		Grand Mal Convulsion					
		Head Injury					
		Loss Of Consciousness					

Date:05/25/04ISR Number: 4364471-1Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0332974A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	ORAL
		Decreased Appetite					
		Insomnia					
		Mood Altered					
		Obsessive Thoughts					

Date:05/26/04ISR Number: 4364806-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0500390A

Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
1MG Per day	1 YR	Diarrhoea		Lotronex	PS	Glaxosmithkline	ORAL
				Wellbutrin Xl	SS	Glaxosmithkline	ORAL
				Nicoderm	SS	Glaxosmithkline	

UNKNOWN

Date:05/26/04ISR Number: 4365515-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506925A  
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Eye Haemorrhage	Bupropion	PS	Glaxosmithkline	ORAL
75MG See			Retinal Haemorrhage				
dosage text			Vision Blurred				
			Vitreous Floaters				

Date:05/26/04ISR Number: 4365521-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0512065A  
Age:85 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -			Confusional State	Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 3 WK							
Initial or Prolonged			Insomnia	Lasix	C	Glaxosmithkline	
			Loss Of Consciousness	Procardia Xl	C	Glaxosmithkline	
				Ambien	C		
				Pravacol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/26/04ISR Number: 4365523-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0512078A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300MG Per day	7	MON	Confusional State	Wellbutrin	PS	Glaxosmithkline	ORAL
40MG per day			Lymphoplasia	Prozac	SS		
			Memory Impairment				
			Suicidal Ideation				

Date:05/26/04ISR Number: 4365540-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0333110A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
150MG Per day	6	DAY	Coronary Artery Stenosis	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged			Myocardial Infarction				

Date:05/27/04ISR Number: 4366581-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511889A  
Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
1 YR			Grand Mal Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
			Libido Decreased	Paxil Cr	SS	Glaxosmithkline	
1 YR			Medication Error	Birth Control Pill	C		
			Mood Altered				

Date:05/27/04ISR Number: 4366582-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0512061A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
			Abortion Spontaneous	Wellbutrin	PS	Glaxosmithkline	ORAL
			Drug Exposure During				
			Pregnancy				

Date:05/27/04ISR Number: 4366585-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0323094A

Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 10 DAY				Zyban	PS	Glaxosmithkline	ORAL
250MG Twice				Erymax	C	Glaxosmithkline	ORAL
per day	8 DAY						
		Hypersomnia					
		Sleep Disorder		Cocodamol	C		ORAL
		Stress		Premarin	C		ORAL

Date:05/27/04ISR Number: 4366587-2Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0327731A

Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice				Zyban	PS	Glaxosmithkline	ORAL
per day	4 DAY						
		Skin Lesion					
				Thyroxine	C	Glaxosmithkline	

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Freedom Of Information (FOI) Report

Date:05/27/04ISR Number: 4366595-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0333228A

Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day	15 DAY	Abortion Spontaneous		Zyban	PS	Glaxosmithkline	ORAL
		Necrobiosis Placental Disorder					

Date:05/28/04ISR Number: 4367110-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0505967A

Age:16 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300MG See		Convulsion	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
dosage text	3 WK	Drooling Eye Rolling Fall Headache Medication Error Tremor	Professional				

Date:05/28/04ISR Number: 4367111-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0507194A

Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 300MG Per day		Medication Error	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day		Overdose	Professional	Wellbutrin	SS	Glaxosmithkline	ORAL
20MG Per day				Prilosec	C		ORAL
3.75MG Per day				Paxil Cr	C	Glaxosmithkline	ORAL

Nasonex

C

2SPR Per day

Date:05/28/04ISR Number: 4367112-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508672A

Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	300MG Per day 3 DAY	Dyspnoea	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Other	10MG per day	Rash	Professional	Ambien	C		ORAL
		Throat Tightness		Xanax	C		ORAL
1MG At night		Vomiting					

Date:05/28/04ISR Number: 4367113-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0510000A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day 10 WK	Aphasia	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
		Balance Disorder	Professional	Depakote	C		
		Coordination Abnormal					
		Dysarthria		Protonix	C		
		Irritability		Pravacol	C		
		Judgement Impaired					
		Road Traffic Accident					
		Sleep Disorder					
		Sluggishness					
		Tremor					

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Freedom Of Information (FOI) Report

Date:05/28/04ISR Number: 4367114-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0510367A  
Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Health Professional	Bupropion	PS	Glaxosmithkline	ORAL

Date:05/28/04ISR Number: 4367122-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0512178A  
Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous Multiple Sclerosis		Bupropion	PS	Glaxosmithkline	

Date:05/28/04ISR Number: 4367123-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0512399A  
Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice per day		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/28/04ISR Number: 4367128-6Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0327981A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG Three Initial or Prolonged times per day		Abdominal Pain Hepatic Enzyme Increased Pancreatic Abscess		Zyban	PS	Glaxosmithkline	ORAL

Date:05/28/04ISR Number: 4367145-6Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0043570A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 2 DAY Initial or Prolonged		Anaphylactic Shock	Health Professional	Zyban	PS	Glaxosmithkline	ORAL
		Cardiovascular Disorder Pharmaceutical Product Complaint					

Date:05/28/04ISR Number: 4367412-6Report Type:Direct Company Report #CTU 219743  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150 MG DAILY ORAL		Conjunctivitis Erythema Multiforme Myalgia Pharyngolaryngeal Pain Pyrexia Rhinorrhoea		Wellbutrin Xl 150 Mg Glaxosmithkline	PS	Glaxosmithkline	ORAL

Date:06/01/04ISR Number: 4367668-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0512619A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion Drug Interaction		Wellbutrin Sr Decongestant No Concurrent	PS SS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Medication C

Date:06/01/04ISR Number: 4367686-1Report Type:Expedited (15-DaCompany Report #FI-GLAXOSMITHKLINE-B0334085A  
 Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	15 DAY	Psoriasis		Zyban	PS	Glaxosmithkline	ORAL
				Nicotine Substitute	C		
				Buventol	C	Glaxosmithkline	
RESPIRATORY							
(INHALATION)							
				Beclomet	C	Glaxosmithkline	
RESPIRATORY							
(INHALATION)							

Date:06/01/04ISR Number: 4367687-3Report Type:Expedited (15-DaCompany Report #2004009670  
 Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Single	Erythema		Zyban	PS	Glaxosmithkline	ORAL
dose		Fatigue					
		Intentional Misuse					
		Suicide Attempt					

Date:06/01/04ISR Number: 4367869-0Report Type:Direct Company Report #CTU 219803  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	1 1 ORAL	Drug Ineffective		Vioxx 25 Mg Merck	PS	Merck	ORAL
Initial or Prolonged	1 1 ORAL	Fungal Infection		Wellbutrin 300 Mg			
		Stevens-Johnson Syndrome		Glaxo	SS	Glaxo	ORAL

Date:06/01/04ISR Number: 4369283-0Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 219884

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Life-Threatening	Agitation		Paxil	PS		
Hospitalization -	Diarrhoea		Prozac	SS		
Initial or Prolonged	Insomnia		Effexor	SS		
	Intentional Misuse		Celexa	SS		
	Muscle Contractions		Remeron	SS		
	Involuntary		Lexapro	SS		
	Restlessness		Serzone	SS		
	Stress		Wellbutrin	SS		
	Suicidal Ideation		Zoloft	SS		
	Suicide Attempt		Sleep Pills	C		

Date:06/02/04ISR Number: 4368391-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0512883A  
Age:61 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Cough		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 2 WK	Hallucination		Toprol Xl	C		
	Insomnia		Plavix	C		
	Sleep Talking		Amiloride	C		
			Norvasc	C		
			Avapro	C		
			Keppra	C		
			Lexapro	C		

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Freedom Of Information (FOI) Report

Vicodin C  
Ambien C

Date:06/02/04ISR Number: 4368392-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0512890A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day	7 DAY	Dyskinesia	Wellbutrin	PS	Glaxosmithkline	ORAL
			Hallucination	Vioxx	C		
			Insomnia	Effexor Xr	C		
			Paraesthesia				
			Tinnitus				

Date:06/02/04ISR Number: 4368393-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0512894A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Mental Disorder	Wellbutrin	PS	Glaxosmithkline	ORAL
				Depakote	C		

Date:06/02/04ISR Number: 4368394-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0512998A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Medication Error	Wellbutrin	PS	Glaxosmithkline	

Date:06/02/04ISR Number: 4371503-3Report Type:Expedited (15-DaCompany Report #TPA2004A00298  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other Required Intervention to 15 MG, 1 IN 1			Alanine Aminotransferase Increased Aspartate	Health Professional Company			
				Actos (Pioglitazone Hydrochloride) (15 Milligram, Tablets)	PS		ORAL

Prevent Permanent  
D, PER ORAL  
Impairment/Damage

Aminotransferase  
Increased  
Blood Alkaline  
Phosphatase Increased  
Liver Function Test  
Abnormal  
Medication Error

Representative

Wellbutrin Sr  
(Bupropion  
Hydrochloride) SS  
Cozaar(Losartan  
Potassium) SS  
Niaspan (Nicotinic  
Acid) SS  
Zoloft (Sertraline  
Hydrochloride) SS  
Birth Control Pills  
(All Other  
Therapeutic  
Products) C

Date:06/03/04ISR Number: 4369995-9Report Type:Expedited (15-DaCompany Report #A03200400600  
Age:48 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Per day Initial or Prolonged 5MG See Other dosage text 9 DAY	Suicide Attempt		Wellbutrin Sr  Ambien  Paroxetine	PS  SS	Glaxosmithkline	ORAL  ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

4 DAY Hydrochloride C Glaxosmithkline ORAL

Date:06/03/04ISR Number: 4370005-8Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0334041A  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Zyban	PS	Glaxosmithkline	ORAL
1 DAY		Delirium Tremens Hallucination Overdose Restlessness Tardive Dyskinesia					

Date:06/03/04ISR Number: 4370006-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0334091A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Myocardial Infarction		Zyban	PS	Glaxosmithkline	
UNKNOWN							

Date:06/03/04ISR Number: 4371820-7Report Type:Direct Company Report #CTU 220092  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Wellbutrin 33 Mg	PS		ORAL
ONCE DAILY		Hostility					
ORAL							

Date:06/03/04ISR Number: 4372073-6Report Type:Direct Company Report #CTU 220074  
 Age:9 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour		Wellbutrin 75 Mg			

37.5 MG 2	Aggression	Burpropion Hcl	PS	ORAL
DAILY ORAL	Communication Disorder			
	Confusional State	Trazadone	C	
	Crying			
	Mood Swings			
	Screaming			
	Social Avoidant Behaviour			

Date:06/04/04ISR Number: 4371192-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0509840A  
Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Insomnia	Health	Requip	PS	Glaxosmithkline	ORAL
3 DAY							
		Myoclonus	Professional	Wellbutrin	SS	Glaxosmithkline	ORAL
2 YR							
		Simple Partial Seizures		Celexa	C		ORAL
		Tremor					

Date:06/04/04ISR Number: 4371204-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0334011A  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cerebral Haemorrhage		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
Initial or Prolonged		Coordination Abnormal					
per day	14 DAY						
Other		Dizziness					
		Sensory Loss					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/04/04ISR Number: 4371206-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0334151A  
 Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day Other	2 MON	Neuralgia	Musculoskeletal Stiffness	Zyban	PS	Glaxosmithkline	ORAL

Date:06/07/04ISR Number: 4372161-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0513282A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Per day 6 MON	6 MON	Visual Acuity Reduced		Wellbutrin	PS	Glaxosmithkline	ORAL
				Advair Allergy Medication	C C	Glaxosmithkline	

Date:06/07/04ISR Number: 4372167-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0513477A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:06/07/04ISR Number: 4372168-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0513555A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Fatigue Liver Disorder Pyrexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	1TAB Per day	3 DAY	Abnormal Behaviour	Zyban	PS	Glaxosmithkline	ORAL
Other			Aggression	Inderal	C		
UNKNOWN			Dehydration				
			Depression				
			Incoherent				
			Insomnia				
			Mood Altered				
			Restlessness				
			Smoker				
			Tearfulness				
			Weight Decreased				

Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG per day		Abortion Spontaneous	Bupropion	PS	Glaxosmithkline	
Other			Drug Exposure During Pregnancy				
			Pregnancy				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/08/04ISR Number: 4373242-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0513586A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Medication Error	Wellbutrin	PS	Glaxosmithkline	
				Bupropion	SS	Glaxosmithkline	

Date:06/08/04ISR Number: 4373243-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0513587A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Medication Error	Wellbutrin	PS	Glaxosmithkline	
				Bupropion	SS	Glaxosmithkline	

Date:06/08/04ISR Number: 4373244-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0513589A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Medication Error	Wellbutrin	PS	Glaxosmithkline	
				Bupropion	SS	Glaxosmithkline	

Date:06/08/04ISR Number: 4373245-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0513590A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Medication Error	Wellbutrin	PS	Glaxosmithkline	
				Bupropion	SS	Glaxosmithkline	

Date:06/08/04ISR Number: 4373246-9Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0513592A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening			Coma	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
30TAB Unknown							

Hospitalization - Overdose  
Initial or Prolonged Suicide Attempt

Date:06/08/04ISR Number: 4373247-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0513593A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	
				Bupropion	SS	Glaxosmithkline	

Date:06/08/04ISR Number: 4373248-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0513594A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin	PS	Glaxosmithkline	
				Bupropion	SS	Glaxosmithkline	



Pregnancy  
Pregnancy  
Trisomy 21

Date:06/08/04ISR Number: 4374157-5Report Type:Direct Company Report #CTU 220286  
Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bursitis		Bupropion Sr	PS	Watson	
150 MG DAILY		Urticaria					

Date:06/09/04ISR Number: 4374652-9Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0334661A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Death		Zyban	PS	Glaxosmithkline	ORAL
Death							

Date:06/09/04ISR Number: 4374653-0Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0043865A  
Age:39 YR Gender:Male I/FU:F

Outcome	PT
Other	Angina Pectoris Chest Pain

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sense Of Oppression

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
100MG per day	22 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
			Ritalin	C		ORAL

Date:06/10/04ISR Number: 4375645-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0512065A  
Age:85 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	300MG Per day	3 WK	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Insomnia		Lasix	C	Glaxosmithkline	
		Loss Of Consciousness		Procardia Xl	C	Glaxosmithkline	
				Ambien	C		
				Pravacol	C		

Date:06/11/04ISR Number: 4376085-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506081B  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG per day	78 DAY	Health	Bupropion	PS	Glaxosmithkline	
		Pregnancy	Professional	Tobacco	SS		
		Sudden Infant Death Syndrome					

Date:06/11/04ISR Number: 4376088-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508423A  
Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day	8 DAY	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -		Bronchospasm	Professional	Spectazole Cream	C		
Initial or Prolonged		Dyspnoea					
Disability		Rash					

Other Urticaria

Date:06/11/04ISR Number: 4376097-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514128A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
3	WK		Abdominal Distension	Wellbutrin	PS	Glaxosmithkline	ORAL
			Flight Of Ideas	No Concurrent			
			Muscle Spasms	Medication	C		
			Suicidal Ideation				

Date:06/11/04ISR Number: 4376106-2Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0334822A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -			Depression	Bupropion			
Initial or Prolonged			Hyperventilation	Hydrochloride	PS	Glaxosmithkline	ORAL
300MG per day	23	DAY					
			Pyrexia				
			Rash				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/14/04ISR Number: 4377013-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514321A  
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 12.5G Single Initial or Prolonged dose		Coma		Wellbutrin	PS	Glaxosmithkline	ORAL
Disability 120MG Single dose		Overdose		Paxil	SS	Glaxosmithkline	
30MG Single dose				Ativan	SS		

Date:06/14/04ISR Number: 4377015-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514346A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 3 YR		Clavicle Fracture		Wellbutrin	PS	Glaxosmithkline	ORAL
		Grand Mal Convulsion Laceration		Accupril Lipitor	C C		

Date:06/14/04ISR Number: 4377016-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0307434A  
 Age:64 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG See dosage text 850MG Three times per day	1 MON	Blood Glucose Increased		Zyban	PS	Glaxosmithkline	ORAL
		Diabetes Mellitus		Glucophage	C		ORAL
		Diabetes Mellitus Inadequate Control		Diamicron Tahor	C C		ORAL ORAL

35MG Twice	Vastarel	C	ORAL
per day			
250MG Per day	Aspegic	C	ORAL
150MG Three	Mediator	C	ORAL
times per day			

Date:06/14/04ISR Number: 4377061-1Report Type:Expedited (15-DaCompany Report #US-ROCHE-370114  
 Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - UNKNOWN	Diarrhoea		Klonopin	PS	Roche	
Initial or Prolonged	Dizziness		Wellbutrin Xl	SS		ORAL
547 DAY	Drug Screen Positive		Wellbutrin Xl	SS		ORAL
	Muscle Spasms Overdose Vomiting		Lithium Salt	C		

Date:06/14/04ISR Number: 4378327-1Report Type:Direct Company Report #CTU 220700  
 Age:48 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Lip Pain Pruritus Rash Swelling Face Urticaria		Bupropion Hydrochloride	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/15/04ISR Number: 4377600-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514333A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Medication Error					
				Wellbutrin Sr	SS	Glaxosmithkline	
150MG Per day							

Date:06/16/04ISR Number: 4378533-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514663A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
		Medication Error					

Date:06/16/04ISR Number: 4378534-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514689A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Faecal Incontinence		Hydrochlorothiazide	C		
				Klonopin	C		
				Seroquel	C		
				Protonix	C		
				Bentyl	C		
				Singulair	C		

Date:06/16/04ISR Number: 4378535-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514694A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion Overdose		Zomig	SS		

Date:06/16/04ISR Number: 4378536-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514701A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Three times per day							

Date:06/16/04ISR Number: 4378537-3Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0514795A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Suicide Attempt		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:06/16/04ISR Number: 4378555-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0335047A

Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia		Zyban	PS	Glaxosmithkline	
UNKNOWN	150MG	See					
dosage text 27 DAY							
Dermatitis Allergic							
Joint Swelling							
Pruritus							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/16/04ISR Number: 4378561-0Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0335131A

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Cerebrovascular Accident		Zyban	PS	Glaxosmithkline	ORAL
6 WK						
Initial or Prolonged	Delirium		Atacand	C		
	Headache		Lipex	C		
	Ill-Defined Disorder					

Date:06/16/04ISR Number: 4378562-2Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0335216A

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Blood Pressure Increased		Zyban	PS	Glaxosmithkline	
UNKNOWN						
Initial or Prolonged	Dysstasia					
	Hallucination, Visual					
	Myocardial Infarction					
	Tremor					
	Vomiting					

Date:06/16/04ISR Number: 4378563-4Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0335239A

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Death	Brain Neoplasm		Zyban	PS	Glaxosmithkline	ORAL

Date:06/16/04ISR Number: 4380351-XReport Type:Direct Company Report #CTU 220794

Age:16 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Grand Mal Convulsion		Wellbutrin Xl 300 Mg	PS		ORAL
PO DAILY						
Hospitalization -	Rash		Seroquel	C		
Initial or Prolonged	Respiratory Arrest					
	Treatment Noncompliance					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other Required 10MG, 1IN 1 Intervention to D, ORAL Prevent Permanent 300MG, 1 IN 1 Impairment/Damage D, ORAL		Grand Mal Convulsion	Health Professional  Other	Reglan-10mg-Tablet (Metoclopramide Hcl)  Bupropion  Valsartan-Hct Amlodipine Simvastatin Diflunisal Doxepin Levothyroxine Omeprazole Venlafaxine Fluoxetine Hydrochloroquine Folic-Acid Vitamin B-12 Vitamin B-6	PS  SS  C C C C C C C C C C C C C C		ORAL  ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/17/04ISR Number: 4381141-4Report Type:Direct  
 Age:46 YR Gender:Male I/FU:I

Company Report #USP 56576

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Wellbutrin Xl	PS	Glaxosmithkline	
TABLET,							
EXTENDED							
RELEASE							

Wellbutrin Sr SS

Date:06/17/04ISR Number: 4381302-4Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #CTU 220949

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Rash		Bupropion	PS		ORAL
100 MG BID PO							
Initial or Prolonged Serum Sickness							

Date:06/17/04ISR Number: 4381370-XReport Type:Direct  
 Age:46 YR Gender:Male I/FU:I

Company Report #CTU 220990

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Confusional State		Buproprion 150 Mg	PS		ORAL
150 MG TWICE							
Intervention to Dizziness							
DAIL ORAL							
Prevent Permanent Headache							
Impairment/Damage Nausea							
Pharmaceutical Product							
Complaint							

Date:06/17/04ISR Number: 4381396-6Report Type:Direct  
 Age: Gender: I/FU:I

Company Report #CTU 220914

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Medication Error

Bupropion Sr 100 Mg		
Watson	PS	Watson
Bupropion Sr 150 Mg		
Watson	SS	Watson

Date:06/18/04ISR Number: 4380594-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514646A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error		Wellbutrin	PS	Glaxosmithkline	
450MG Per day				Bupropion	SS	Glaxosmithkline	
450MG Single							
dose							

Date:06/18/04ISR Number: 4380597-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514884A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Feeling Abnormal		Ultram	C		
300MG Per day		Feeling Jittery					
		Headache					
		Nausea					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/18/04ISR Number: 4380598-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLIN-A0515009A  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day	3 DAY	Aggression	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Anger	Zocor	C		
			Disorientation	Fosamax	C		
			Nervousness	Lotensin	C	Glaxosmithkline	
			Physical Assault	Celebrex	C		
				Nifedipine	C	Glaxosmithkline	
				Flonase	C	Glaxosmithkline	
				Patanol	C		

Date:06/18/04ISR Number: 4380620-3Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLIN-B0335716A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day		Aggression	Zyntabac	PS	Glaxosmithkline	ORAL
			Confusional State	Sertraline	C		
			Disorientation	Antidepressants	C		
			Irritability				

Date:06/18/04ISR Number: 4383496-3Report Type:Expedited (15-DaCompany Report #B0334697A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	5 WK		Abnormal Behaviour - Anhedonia Depression	Foreign Literature Health		Wellbutrin (Bupropion Hydrochloride)	PS
			Mania Thought Blocking	Professional			

Date:06/18/04ISR Number: 4383691-3Report Type:Expedited (15-DaCompany Report #001738  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Alopecia Amnesia Grand Mal Convulsion Incontinence	Health Professional Company Representative	Cenestin (Synthetic Conjugated Estrogens,A) Tablet, 1.25mg	PS		ORAL
1.00 TABLET,			Other				
QD,ORAL				Wellbutrin-Slow Release (Bupropion Hydrochloride)	SS		ORAL
300.00 MG,				Singulair (Montelukast Sodium)	C		
QD, ORAL				Levothyroxine (Levothyroxine)	C		
				Hydrochlorothiazide	C		
Date:06/21/04ISR Number: 4381499-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514657A							
Age: Gender: I/FU:I							
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	450MG per day	Drug Exposure During Pregnancy Vocal Cord Paralysis		Bupropion	PS	Glaxosmithkline	
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/21/04ISR Number: 4381500-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514662A

Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other				Bupropion	PS	Glaxosmithkline	
150MG per day							
		Pregnancy					
		Metrorrhagia					

Date:06/21/04ISR Number: 4381508-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0515139A

Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other				Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	4 WK						
		Asthenia					
		Chest Pain					
		Chills		Prednisone	C		
		Cough		Premarin	C		
		Disorientation		Medroxyprogesterone	C		
		Dizziness		Benadryl	C	Glaxosmithkline	
		Dysphagia		Maxair	C		
		Feeling Abnormal		Serevent	C	Glaxosmithkline	
		Gastric Disorder		Azmacort	C		
		Headache					
		Hyperhidrosis					
		Ill-Defined Disorder					
		Increased Tendency To Bruise					
		Mucous Membrane Disorder					
		Muscle Atrophy					
		Muscle Disorder					
		Muscle Spasms					
		Musculoskeletal Disorder					
		Nausea					
		Nervousness					
		Pain Of Skin					
		Pruritus					
		Pruritus Generalised					
		Pyrexia					
		Skin Disorder					
		Tonsillar Cyst					
		Tremor					

Urticaria  
Vasodilatation  
Wheezing

Date:06/21/04ISR Number: 4381522-9Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0335216A

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - UNKNOWN Initial or Prolonged	Blood Pressure Increased	Health	Zyban	PS	Glaxosmithkline	
	Dysstasia Hallucination, Visual Myocardial Infarction Tremor Vomiting	Professional				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/21/04ISR Number: 4382360-3Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 221114

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Anxiety		Bupropion Sr 150 Mg			
Disability	Drug Withdrawal Syndrome		Sr Watson Labs/Penn			
Required	Headache		Labs	PS	Watson Labs/Penn	
Intervention to	Major Depression				Labs	ORAL
150 MG BID						
Prevent Permanent	Panic Attack					
ORAL						
Impairment/Damage	Pharmaceutical Product		Bupropion Sr 150 Mg			
	Complaint		Sr Watson Labs/Penn			
	Self-Injurious Ideation		Labs	SS		
	Suicidal Ideation					
	Suicide Attempt					

Date:06/21/04ISR Number: 4382362-7Report Type:Direct  
Age:48 YR Gender:Male I/FU:I

Company Report #CTU 221116

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Disability	Anger		Budeprion Sr 100mg			
Required	Depression		Teva Pharm	PS	Teva Pharm	ORAL
2 TABLET 2X						
Intervention to	Drug Effect Decreased					
PER DAY ORAL						
Prevent Permanent	Irritability		Elavil-Amitriptylin-	C		
Impairment/Damage	Pharmaceutical Product					
	Complaint					

Date:06/21/04ISR Number: 4383969-3Report Type:Expedited (15-DaCompany Report #US-2004-026688  
Age:36 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Other	Arrhythmia	Consumer	Betaseron(Interferon			
	Blood Pressure Increased		Beta - 1b)			
	Cardiac Flutter		Injection, 250ug	PS		
SUBCUTANEOUS	EVERY					
	8 MIU,					
	Drug Interaction					
2D,						

SUBCUTANEOUS

Potentiation

Extrasystoles  
Liver Function Test  
Abnormal

Wellbutrin Xr (150  
Mg (Bupropion  
Hydrochloride) SS

ORAL

150 MG, 1

X/DAY, ORAL

Provigil (Modafinil) C

Date:06/21/04ISR Number: 4393783-0Report Type:Periodic

Company Report #KII-2002-0000442

Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose Multiple Drug Overdose	Health Professional Company	Oxycontin Tablets (Oxycodone Hydrochloride)	PS		
80 MG, TID			Representative	Amitriptyline (Amitriptyline) Prozac (Fluoxetine Hydrochloride) Ambien (Zolpidem Tartrate) Bupropion (Amfebutamone)	SS SS SS SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/21/04ISR Number: 4396249-7Report Type:Periodic  
 Age:44 YR Gender:Female I/FU:I

Company Report #USA-2003-0010973

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse Multiple Drug Overdose	Health Professional Other	Oxycontin Hydrochloride (Similar To 20-553) (Oxycodone Hydrochloride) Hydrocodone Bitartrate (Similar To Ind 59,175) (Hydrocodone Bitartrate) Acetaminophen (Acetaminophen) Diphenhydramine (Diphenhydramine) Bupropion (Amfebutamone) Temazepam (Temazepam) Venlafaxine (Venlafaxine) Caffeine (Caffeine)	PS     SS SS SS SS SS SS		

Date:06/21/04ISR Number: 4398890-4Report Type:Periodic  
 Age:47 YR Gender:Male I/FU:F

Company Report #USA-2003-0007098

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Multiple Drug Overdose Accidental	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553) Oxycodone Hydrochloride Hydrocodone Bitartrate (Similar To Ind 59,175) (Hydrocodone Bitartrate) Morphine Sulfate (Similar To Nda 19-516) (Morphine	PS     SS		

Sulfate)	SS
Dihydrocodeine/ Caffeine/ Acetaminophen (Similar To And 88-584)	SS
Diazepam (Diazepam)	SS
Temazepam (Temazepam)	SS
Oxazepam (Oxazepam)	SS
Bupropion (Amfebutamone)	SS

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/22/04ISR Number: 4382632-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514898A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Adverse Event		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2 YR						
		Agitation		Unknown Blood			
		Convulsion		Pressure Medication	SS		
		Disorientation					
		Headache					
		Hypertension					
		Insomnia					
		Logorrhoea					
		Medication Error					
		Nausea					

Date:06/22/04ISR Number: 4382633-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514998A

Age:18 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 DAY						
		Cerebrovascular Accident					
		Confusional State					
		Dysarthria					
		Headache					
		Hypoaesthesia					
		Movement Disorder					
		Speech Disorder					
		Transient Ischaemic					
		Attack					

Date:06/22/04ISR Number: 4382637-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0515236A

Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	6 WK						
		Fatigue		Bupropion	SS	Glaxosmithkline	
450MG per day							

MON  
Irritability Paxil SS Glaxosmithkline ORAL  
Medication Error No Concurrent  
Sexual Dysfunction Medication C

Date:06/22/04ISR Number: 4382654-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0335459A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	ORAL
10 DAY		Arthralgia					
		Constipation					
		Dizziness					
		Fatigue					
		Hallucination					
		Headache					
		Hypogeusia					
		Insomnia					
		Nausea					
		Nervousness					
		Pruritus					
		Rash					
		Tinnitus					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/23/04ISR Number: 4383643-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511755A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	4	DAY					
		Medication Error					
				Lexapro	C		
				Risperdal	C		

Date:06/23/04ISR Number: 4383658-5Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0328342A

Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cerebrovascular Accident		Zyban	PS	Glaxosmithkline	ORAL
150MG Three							
times per day							
				No Concurrent Medication	C		

Date:06/23/04ISR Number: 4383671-8Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0335720A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anhedonia Depression Social Problem Suicidal Ideation		Zyban	PS	Glaxosmithkline	ORAL

Date:06/23/04ISR Number: 4383674-3Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0335860A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depressed Level Of		Zyban	PS	Glaxosmithkline	ORAL
4 WK							

Consciousness  
Disturbance In Attention  
Memory Impairment  
Sleep Disorder  
Somnolence

Serepax

C

Date:06/24/04ISR Number: 4384405-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0513060A  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300MG Twice per day		Drug Abuser Medication Error Psychotic Disorder		Wellbutrin Seroquel	PS C	Glaxosmithkline	NASAL
600MG Twice per day				Trileptal	C		
600MG Twice per day				Neurontin	C		

Date:06/24/04ISR Number: 4384409-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0515707A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG per day		Abortion Spontaneous		Bupropion Celexa Remeron	PS C C	Glaxosmithkline	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/24/04ISR Number: 4384412-0Report Type:Expedited (15-DaCompany Report #CH-GLAXOSMITHKLINE-B0330728A  
Age:49 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Per day 18 DAY	Amnesia	Health	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged Disability	Demyelination Grand Mal Convulsion Incontinence Leukoencephalomyelitis Leukoencephalopathy	Professional				

Date:06/24/04ISR Number: 4384413-2Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0332456A  
Age:4 MON Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other	Abnormal Behaviour Accidental Exposure Acidosis Convulsion Crying Cyanosis Drug Toxicity Emotional Distress Extremity Contracture Eye Movement Disorder Haemoglobin Decreased Hypotonia Irritability Nervous System Disorder Pallor Respiratory Acidosis Restlessness Skin Discolouration Treatment Noncompliance	Health Professional	Zyban	PS	Glaxosmithkline	ORAL

Date:06/24/04ISR Number: 4384416-8Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0335535A  
Age:44 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Other	Rash	Zyban	PS	Glaxosmithkline	ORAL
1TAB See					
	Skin Exfoliation				
dosage text	20 DAY	Nicorette	C	Glaxosmithkline	
UNKNOWN					

Date:06/24/04ISR Number: 4384417-XReport Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0335693A  
 Age:23 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Convulsion		Zyban	PS	Glaxosmithkline	
150MG Twice						
Initial or Prolonged	Insomnia					
per day						
Other	Starvation					

Date:06/24/04ISR Number: 4388630-7Report Type:Expedited (15-DaCompany Report #B0335075A  
 Age:15 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Aggression
Initial or Prolonged	Anger
	Attention

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Deficit/Hyperactivity Disorder Condition Aggravated Drug Abuser	Report Source	Product	Role	Manufacturer	Route
		Feeling Abnormal Impulsive Behaviour Irritability Medication Error	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		NASAL
INTRANASAL		Restlessness Suicidal Ideation Theft Treatment Noncompliance		Salbutamol Sulphate Cannabis	C C		

Date:06/24/04ISR Number: 4388960-9Report Type:Expedited (15-DaCompany Report #2003-06064  
Age:26 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cholelithiasis Hepatitis	Study Health	Viread(Tenofovir Disoproxil Fumarate)	PS		ORAL
300 MG, ONCE A DAY, ORAL		Pancreatitis	Professional				
450 MG, TWICE A DAY, ORAL				Combivir(Zidovudine W/Lamivudine)	SS		ORAL
				Wellbutrin(Bupropion Hydrochloride)	SS		
				Ambien (Zolpidem Tartrate)	C		
				Effexor (Venlafaxine Hydrochloride)	C		
				Asacol (Mesalazine)	C		
				Clindagel (Clindamycin)	C		
				Elidel (Pimecrolimus)	C		
				Clotrim/Beta	C		
				Protopic(Tacrolimus)	C		
				Flonase(Fluticasone Propionate)	C		

Date:06/25/04ISR Number: 4385017-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0515239A  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice		Pulmonary Thrombosis	Wellbutrin	PS	Glaxosmithkline	ORAL
	per day	5 DAY	Road Traffic Accident				
1	YR			Tegretol	SS		
				Librium	SS		
1	YR			Nortriptyline	C		
1	YR			Depakote	C		
				Seroquel	C		
1	YR			Trazodone	C		
1	YR			Celexa	C		

Date:06/25/04ISR Number: 4385020-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0515738A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	8TAB per day		Overdose	Wellbutrin	PS	Glaxosmithkline	ORAL
			Suicidal Ideation				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/25/04ISR Number: 4385021-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0515739A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Coma		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day		Knee Operation WK		Zoloft	C		
				Alcohol	C		

Date:06/25/04ISR Number: 4385043-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0335857A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Unknown		Cerebrovascular Accident		Zyban	PS	Glaxosmithkline	ORAL

Date:06/25/04ISR Number: 4386443-3Report Type:Direct Company Report #CTU 221524

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 20 MG 3 X A		Agitation Anxiety		Prozac 20 Mg Eli-Lilly	PS	Eli-Lilly	ORAL
Initial or Prolonged DAY ORAL		Depression		Wellbutrin 150 Mg Glaxosmithkline	SS	Glaxosmithkline	ORAL
150 MG 2 X A DAY ORAL		Insomnia Restlessness Suicide Attempt					

Date:06/28/04ISR Number: 4386249-5Report Type:Direct Company Report #CTU 221662

Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Life-Threatening  
DAILY 2 WK  
Required  
Intervention to  
Prevent Permanent  
Impairment/Damage

Depression  
Mental Disorder

Wellbutrin PS

Date:06/28/04ISR Number: 4386252-5Report Type:Direct  
Age:16 YR Gender:Female I/FU:I

Company Report #CTU 221667

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG PO		Palpitations		Wellbutrin Sr	PS		ORAL
DAILY				Nicotine Gum	C		

Date:06/28/04ISR Number: 4386898-4Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 221717

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG 1 PER		Depression		Wellbutrin Xl 300 Mg	PS		
DAY		Pharmaceutical Product Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/28/04ISR Number: 4389102-6Report Type:Expedited (15-DaCompany Report #2004UW06035  
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Seroquel	PS		
Other		Abasia		Serzone	SS		
Required		Dysstasia		Wellbutrin	SS		
Intervention to		Fall		Motrin	C		
Prevent Permanent		Medication Error		Heparin	C		
Impairment/Damage		Musculoskeletal Stiffness		Ultram	C		
		Rhabdomyolysis		Prednisone	C		
				Cellcept	C		
				Sleeping Pill	C		

Date:06/29/04ISR Number: 4386599-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0515716A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other		Abortion Spontaneous					
150MG Per day							

Date:06/29/04ISR Number: 4386601-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0516112A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Death		Completed Suicide		Ativan	C		
150MG Per day				Thyroid Medication	C		
Life-Threatening		Gun Shot Wound					
Other		Murder					

Date:06/29/04ISR Number: 4386603-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0516246A  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Life-Threatening		Haematuria		Advair	C	Glaxosmithkline	
300MG Per day 1 WK		Pain					

Albuterol C Glaxosmithkline  
Singulair C

Date:06/29/04ISR Number: 4386624-9Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0336916A

Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fibromyalgia		Zyban	PS	Glaxosmithkline	
1TAB See							
dosage text	14	DAY	Suicidal Ideation				
			Thinking Abnormal	Morphine	C		

Date:06/30/04ISR Number: 4387784-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0516404A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Creatinine Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
		Blood Urea Increased					
		Hepatic Enzyme Increased					

Freedom Of Information (FOI) Report

Date:06/30/04ISR Number: 4387785-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0516415A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other	150MG Twice	Pregnancy	Drug Exposure During	Bupropion	PS	Glaxosmithkline	
	per day		Jaundice				
			Pulmonary Oedema				

Date:06/30/04ISR Number: 4387790-1Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0516554A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other	150MG Unknown	Vaginal Haemorrhage		Wellbutrin	PS	Glaxosmithkline	ORAL
	16 DAY			Sertraline	C		
UNKNOWN	25MG Unknown			Clonazepam	C		
UNKNOWN	1MG Unknown						

Date:06/30/04ISR Number: 4387797-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0334223A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Hospitalization -		Abasia		Zyban	PS	Glaxosmithkline	
Initial or Prolonged		Confusional State					
Other		Convulsion					
		Depression					
		Disability					
		Epilepsy					
		Impaired Driving Ability					
		Impaired Work Ability					
		Oral Infection					
		Suicidal Ideation					

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1000 MCG (500 Initial or Prolonged MCG, 2 IN 1 Other D), ORAL	Anaemia  Atrial Fibrillation  Chest Pain  Disease Recurrence Dizziness Electrocardiogram Qt Prolonged Gallbladder Disorder Hepatic Enzyme Increased Hyperhidrosis	Consumer	Tikosyn (Dofetilide)	PS		ORAL
			Peginterferon Alfa-2b (Peginterferon Alfa-2b)	SS		
			Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		
			Valdecoxib (Valdecoxib)	C		
			Metoprolol (Metoprolol)	C		
			Warfarin (Warfarin)	C		
			Potassium (Potassium)	C		
			Nizantidine (Nizantidine)	C		
			Antibiotics (Antibiotics)	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/30/04ISR Number: 4390538-8Report Type:Expedited (15-DaCompany Report #2004041004

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2400 MG (800 MG, 3 IN 1 D), ORAL		Drug Hypersensitivity Muscle Spasms  Nervous System Disorder  Pharmaceutical Product  Complaint Tremor Weight Increased	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
				Bupropion Hydrochloride (Bupropion Hydrochloride) Venlafaxine Hydrochloride (Venlafaxine Hydrochloride) Levothyroxine Sodium (Levothyroxine Sodium) Baclofen (Baclofen)	SS  C C C		

Date:06/30/04ISR Number: 4415545-8Report Type:Periodic

Company Report #2003122132

Age:13 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 12.5 MG Initial or Prolonged (DAILY), ORAL Other  75 MG (DAILY), UNKNOWN		Asthenia  Back Pain  Bipolar Ii Disorder Chest Pain Cough Decreased Appetite  Disturbance In Attention  Dizziness  Homicidal Ideation Hyperhidrosis	Consumer  Health  Professional	Zoloft (Sertraline)   Bupropion Hydrochloride (Bupropion Hydrochloride)  Paroxetine Hydrochloride	PS   SS		ORAL

Hypersensitivity  
Increased Appetite  
Influenza Like Illness  
Insomnia  
Lymphadenopathy  
Neck Pain  
Nightmare  
Pyrexia  
Rash  
Renal Pain  
Suicidal Ideation  
Swelling Face  
Vision Blurred

(Paroxetine  
Hydrochloride) C

Date:07/01/04ISR Number: 4388722-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0516229A  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Drug Exposure During Pregnancy Placental Insufficiency		Effexor	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/01/04ISR Number: 4388741-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0516614A

Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
		Akathisia					
per day							
		Hallucination		Seroquel	C		
		Tremor		Neurontin	C		
				Topamax	C		

Date:07/01/04ISR Number: 4388757-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0336854A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Generalised Oedema	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown							
		Nephrotic Syndrome	Professional	Nexen	SS	Glaxosmithkline	ORAL
		Renal Failure					
		Weight Increased					

Date:07/01/04ISR Number: 4388763-5Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0337371A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blindness Transient		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day 1 DAY							
				Betablocker	C		ORAL
				Zocor	C		ORAL
				Kardegic	C		ORAL
75MG Per day							
				Stilnox	C		ORAL

Date:07/01/04ISR Number: 4390067-1Report Type:Direct

Age: Gender: I/FU:I

Company Report #USP 56669

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Medication Error		Wellbutrin Sr	PS	Glaxosmithkline	
	TABLET 150 MG				Wellbutrin Xl	SS	Glaxosmithkline	
	TABLET 150 MG							

Date:07/01/04ISR Number: 4391700-0Report Type:Expedited (15-DaCompany Report #2003125358  
Age:35 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Multiple Drug Overdose Accidental	Health Professional Company Representative	Geodon (Ziprasidone) Bupropion Hydrochloride (Bupropion Hydrochloride) Clonazepam (Clonazepam)	PS    SS SS		

Date:07/01/04ISR Number: 4391982-5Report Type:Expedited (15-DaCompany Report #B0335849A  
Age:29 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Atrial Fibrillation Bundle Branch Block Left Delirium Disorientation Electrocardiogram Qt

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Corrected Interval Prolonged Electrocardiogram T Wave Abnormal	Report Source	Product	Role	Manufacturer	Route
		Grand Mal Convulsion Hallucination, Visual Lethargy Multiple Drug Overdose	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL		Sinus Tachycardia Supraventricular Tachycardia		Ziprasidone Hcl Unspecified Tablet (Ziprasidone Hcl)	SS		ORAL
ORAL				Semisodium Valproate Tablet (Divalproex Sodium)	SS		ORAL
ORAL				Fluoxetine Tablet (Fluoxetine)	SS		ORAL

Date:07/02/04ISR Number: 4389494-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0516613A  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100MG Three times per day 2		Drug Ineffective Psychiatric Symptom MON		Wellbutrin Valium	PS C	Glaxosmithkline	ORAL

Date:07/02/04ISR Number: 4389502-4Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0335441A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 15 DAY		Depression Dry Mouth Fear Hallucination, Auditory Insomnia	Health Professional	Zyntabac	PS	Glaxosmithkline	ORAL

Nervousness

Date:07/02/04ISR Number: 4392087-XReport Type:Expedited (15-DaCompany Report #A118566  
Age:29 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Atrial Fibrillation
Initial or Prolonged	Bundle Branch Block Left
Required	Delirium
Intervention to	Disorientation
Prevent Permanent	Electrocardiogram
Impairment/Damage	Abnormal
	Electrocardiogram Qrs
	Complex Prolonged
	Electrocardiogram Qt
	Corrected Interval
	Prolonged
	Electrocardiogram St
	Segment Abnormal
	Electrocardiogram St
	Segment Depression
	Electrocardiogram T Wave
	Abnormal
	Grand Mal Convulsion
	Hallucination, Visual

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Lethargy Multiple Drug Overdose Sinus Tachycardia					
80 MG		Supraventricular Tachycardia	Literature Health	Ziprasidone (Ziprasidone)	PS		
100 MG			Professional Company Representative	Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		
40 MG				Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	SS		
250 MG				Valproate Semisodium (Valproate Semisodium)	SS		

Date:07/02/04ISR Number: 4392196-5Report Type:Expedited (15-DaCompany Report #US-2004-026688  
Age:36 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Anxiety Arrhythmia Blood Pressure Increased	Consumer Health Professional	Betaseron (Interferon Beta-1b) Injection, 250 Ug	PS		
	SUBCUTANEOUS	8 MIU, EVERY 2D,	Cardiac Flutter					
	SUBCUTANEOUS		Drug Interaction					
	150 MG,		Liver Function Test Abnormal Palpitations		Wellbutrin Xr 150 Mg (Bupropion Hydrochloride)	SS		ORAL
	1X/DAY, ORAL							
	100 MG,				Provigil (Modafinil)	SS		ORAL
	1X/DAY, ORAL							

Neurontin  
(Gabapentin) C  
Baclofen C

Date:07/06/04ISR Number: 4390151-2Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0516866A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown		Alcohol Use Insomnia					

Date:07/06/04ISR Number: 4390152-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0516903A  
Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2	WK		Zyprexa	C		
				Lito	C	Glaxosmithkline	
				Depakote	C		
				Effexor	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/06/04ISR Number: 4390173-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0335459A  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression	Health	Zyban	PS	Glaxosmithkline	ORAL
15 DAY		Arthralgia	Professional				
		Constipation					
		Dizziness					
		Fatigue					
		Hallucination					
		Headache					
		Hypogeusia					
		Insomnia					
		Nausea					
		Nervousness					
		Pruritus					
		Rash					
		Tinnitus					

Date:07/06/04ISR Number: 4390206-2Report Type:Expedited (15-DaCompany Report #IL-GLAXOSMITHKLINE-B0337852A  
Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Energy Increased		Zyban	PS	Glaxosmithkline	ORAL
		Euphoric Mood		No Concurrent			
		Hypomania		Medication	C		
		Insomnia					

Date:07/06/04ISR Number: 4390207-4Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0043570A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anaphylactic Shock	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG Per day 2 DAY							
Initial or Prolonged		Cardiovascular Disorder	Professional				
		Pharmaceutical Product					
		Complaint					

Date:07/06/04ISR Number: 4390780-6Report Type:Direct  
Age:50 YR Gender:Male I/FU:I

Company Report #CTU 222148

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropion			
Other		Drug Ineffective		Bupropion100 Mg Teva	PS	Teva	ORAL
100 MG TID		Pharmaceutical Product					
ORAL		Complaint					
				Bupropion			
				Bupropion100 Mg Teva	SS	Teva	

Date:07/06/04ISR Number: 4394305-0Report Type:Expedited (15-DaCompany Report #04P-163-0264468-00  
Age:29 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Bundle Branch Block Left
Initial or Prolonged	Disorientation
	Electrocardiogram Qt
	Corrected Interval
	Prolonged
	Electrocardiogram T Wave
	Abnormal

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
250 MG, APPROXIMATELY 74 TABLETS, PER ORAL		Grand Mal Convulsion Hallucination, Visual Lethargy	Literature Health Professional	Divalproex Sodium (Depakote) (Divalproex Sodium) (Divalproex Sodium)	PS		ORAL
80 MG, APPROXIMATELY 74 TABLETS, PER ORAL		Multiple Drug Overdose Sinus Tachycardia Supraventricular Tachycardia		Ziprasidone	SS		ORAL
100 MG, APPROXIMATELY 157 TABLETS, PER ORAL				Bupropion	SS		ORAL
40 MG, APPROXIMATELY 180 TABLETS, PER ORAL				Fluoxetine	SS		ORAL

Date:07/07/04ISR Number: 4391419-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0517054A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Date:07/07/04ISR Number: 4391423-8Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0295775A  
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability	43 DAY	Aphasia	Consumer	Zyban	PS	Glaxosmithkline	ORAL
Other		Cerebral Arteritis Cerebral Haemorrhage Computerised Tomogram Abnormal Convulsion Electroencephalogram Abnormal Grand Mal Convulsion Headache Nausea Personality Change Vomiting					

Date:07/07/04ISR Number: 4391430-5Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0309137A  
 Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG See Initial or Prolonged dosage text	20 DAY	Arthralgia	Consumer	Zyban	PS	Glaxosmithkline	ORAL
		Feeling Cold General Physical Health Deterioration Myalgia Pyrexia Rash Pruritic					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/07/04ISR Number: 4391440-8Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0334297A  
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	6	DAY	Constipation	Zyban	PS	Glaxosmithkline	ORAL
			Hallucination				
RESPIRATORY (INHALATION)			Nausea	Spiriva	C		
			Smoker				
RESPIRATORY (INHALATION)				Ventolin	C	Glaxosmithkline	

Date:07/07/04ISR Number: 4391463-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0337415A  
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day			Diabetes Mellitus	Zyban	PS	Glaxosmithkline	ORAL
			Inadequate Control Hypoglycaemia	Insuline	C		

Date:07/07/04ISR Number: 4391465-2Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0337423A  
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day	5	DAY	Bradycardia Dizziness	Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
159 DAY			Hyperhidrosis	Tenormin	SS		ORAL
1.5G Per day	9	DAY	Palpitations	Amoxicillin	C	Glaxosmithkline	ORAL
			Syncope Tremor				

Date:07/07/04ISR Number: 4391467-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0337502A  
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Feeling Abnormal	Zyban	PS	Glaxosmithkline	
6 DAY			Suicidal Ideation	Risperidal	C		

Date:07/07/04ISR Number: 4391469-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0337576A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Heart Rate Increased	Zyban	PS	Glaxosmithkline	
2TAB per day			Hot Flush				
			Loss Of Consciousness				

Date:07/07/04ISR Number: 4391471-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0337791A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Ageusia	Zyban	PS	Glaxosmithkline	ORAL
12 DAY			Anosmia				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/07/04ISR Number: 4393442-4Report Type:Direct  
 Age:57 YR Gender:Female I/FU:I

Company Report #CTU 222340

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Asthenia		Ambien 10 Mg	PS		ORAL
10 MG HS PO		Insomnia		Wellbutrin 150 Mg	SS		ORAL
150 MG PO AM		Sedation					
AND PM		Somnolence					
		Tremor					

Date:07/07/04ISR Number: 4394274-3Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20040201456  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Drug Exposure During Pregnancy	Study Health	Topiramate- Blinded (Topiramate) Tablets	PS		
TRANSPLACENTAL	TRANSPLACENTA	Talipes	Professional				
L				Placebo (Placebo) Tablets	SS		
TRANSPLACENTAL	TRANSPLACENTA						
L				Wellbutrin (Bupropion Hydrochloride)			
TRANSPLACENTAL	TRNASPLACENTA			Unspecified	SS		
L				Prenatal Vitamin (Prenatal Vitamins)			
				Unspecified	C		
				Tylenol (Paracetamol)			
				Unspecified	C		

Date:07/08/04ISR Number: 4393008-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0517052A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG See		Medication Error		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
dosage text							

Date:07/08/04ISR Number: 4393016-5Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0331580A

Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Bupropion			
15 DAY		Anxiety		Hydrochloride	PS	Glaxosmithkline	ORAL
Condition Aggravated							
Depersonalisation							
Food Interaction							
Irritability							
Memory Impairment							

Date:07/08/04ISR Number: 4393020-7Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0335692A

Age:25 YR Gender:Female I/FU:F

Outcome	PT
Other	Arthralgia
	Arthritis
	Arthritis Reactive

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Joint Swelling Osteoarthritis Urinary Tract Infection	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	89 DAY			Zyban	PS	Glaxosmithkline	ORAL
200MG per day				Norethisterone	C		
UNKNOWN				Vioxx	C		

Date:07/08/04ISR Number: 4396008-5Report Type:Expedited (15-DaCompany Report #2004UW13429  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to Prevent Permanent Impairment/Damage		Convulsion Overdose	Health Professional	Zomig Wellbutrin - Slow Release	PS SS		

Date:07/08/04ISR Number: 4397279-1Report Type:Expedited (15-DaCompany Report #2004SE03643  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 25 MG DAILY Intervention to Prevent Permanent Impairment/Damage PO		Bradycardia	Foreign	Tenormin	PS		ORAL
PO Prevent Permanent Impairment/Damage PO		Dizziness	Health				
25 MG DAILY		Palpitations	Professional	Tenormin	SS		ORAL
		Syncope	Other				
		Tremor		Tenormin	SS		ORAL
150 MG DAILY				Zyntabac	SS		ORAL

Amoxicillin C

Date:07/09/04ISR Number: 4393710-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514346A  
 Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	3 YR		Clavicle Fracture	Wellbutrin	PS	Glaxosmithkline	ORAL
Other	40MG Per day		Face Injury	Accupril	C		ORAL
	10MG At night		Fall	Lipitor	C		ORAL
			Grand Mal Convulsion				
			Laceration				
			Rib Fracture				

Date:07/09/04ISR Number: 4393718-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0517370A  
 Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Medication Error	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Pharmaceutical Product	Wellbutrin Sr	SS	Glaxosmithkline	ORAL
			Complaint				

Date:07/09/04ISR Number: 4396206-0Report Type:Expedited (15-DaCompany Report #04P-163-0265030-00  
 Age:75 YR Gender:Male I/FU:I

Outcome  
 Death  
 Hospitalization -

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Initial or Prolonged Required Intervention to Prevent Permanent Dose Duration Impairment/Damage	PT	Report Source	Product	Role	Manufacturer	Route
SEE IMAGE	Abasia Angle Closure Glaucoma Asthenia Brain Damage	Consumer	Depakote Er 500mg (Depakote Er) (Divalproex Sodium) (Divalproex Sodium)	PS		ORAL
10 MG, 1 IN 1 D, PER ORAL	Coma Communication Disorder		Paroxetine Hydrochloride	SS		ORAL
1 IN 1 D, PER ORAL	Decreased Appetite					
1 IN 1 D, PER ORAL	Diabetes Mellitus Disturbance In Attention		Bupropion Hydrochloride	SS		ORAL
1 IN 1 D, PER ORAL	Dysphagia					
1 IN 1 D, PER ORAL	Dyspnoea Feeling Abnormal		Sertraline Hydrochloride	SS		ORAL
SEE IMAGE	Infected Cyst					
1 IN 1 D, PER ORAL	Lethargy		Mirtazapine	SS		ORAL
	Medication Error		Trilazem	SS		ORAL
	Overdose					
	Parkinson'S Disease Pneumonia Aspiration Sleep Disorder Somnolence		Panadeine Co Zocor Baby Asa	C C C		

Date:07/12/04ISR Number: 4394912-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514884A  
Age:29 YR Gender:Male I/FU:F

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day 14 DAY	Back Pain	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged 50MG Twice	Chest Pain	Professional	Ultram	SS		ORAL

per day 20 DAY  
Feeling Abnormal  
Feeling Jittery  
Grand Mal Convulsion  
Headache  
Nausea

Date:07/12/04ISR Number: 4394916-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0517442A  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anger		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice		Convulsion					
per day		Erectile Dysfunction		Ritalin	C		
		Fatigue		Claritin	C		
		Mania		Excedrin	C		
		Mood Altered					
		Tension					

Date:07/12/04ISR Number: 4394946-0Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0337695A  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erythema		Zyban	PS	Glaxosmithkline	ORAL
		Exanthem					
		Influenza Like Illness					
		Lupus-Like Syndrome					

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Freedom Of Information (FOI) Report

Date:07/12/04ISR Number: 4394947-2Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0337716A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Consciousness		Zyban	PS	Glaxosmithkline	ORAL

Date:07/12/04ISR Number: 4394948-4Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0337734A  
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea Haemorrhagic Rash Generalised		Zyban	PS	Glaxosmithkline	ORAL

Date:07/12/04ISR Number: 4394949-6Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0337740A  
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression Asthenia Dry Mouth Erythema		Zyban	PS	Glaxosmithkline	

Date:07/12/04ISR Number: 4394950-2Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0337744A  
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Crying Depressed Mood Excitability Irritability Psychotic Disorder		Zyban	PS	Glaxosmithkline	

Date:07/12/04ISR Number: 4394951-4Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0337779A  
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression Amnesia Loss Of Consciousness	Consumer	Zyban Alcohol	PS SS	Glaxosmithkline	

Date:07/12/04ISR Number: 4394952-6Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0337850A  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erythema Multiforme Exanthem Periorbital Oedema Rash Vesicular Urticaria Generalised		Zyban	PS	Glaxosmithkline	ORAL

Date:07/12/04ISR Number: 4394955-1Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0338473A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Consciousness		Zyban Alcohol	PS C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/12/04ISR Number: 4394957-5Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0043173A  
 Age:70 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1TAB Per day	17 DAY	Cerebral Atrophy	Zyban	PS	Glaxosmithkline	ORAL
UNKNOWN	300MG Per day	17 DAY	Disturbance In Attention	Allopurinol	C	Glaxosmithkline	
UNKNOWN	600MG Per day	17 DAY	Dysgraphia	Alpha-Lipon Acid 600	C		
UNKNOWN		17 DAY	Encephalopathy	Iscover	C		
UNKNOWN			Fall	Ace Inhibitor	C		
			Gait Disturbance				
			Pain				
			Parkinson'S Disease				
			Polyneuropathy				
			Vertigo				

Date:07/12/04ISR Number: 4395384-7Report Type:Direct Company Report #CTU 222528  
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG TWICE		Depression	Budeprion Sr 150 Mg	PS		ORAL
DAIL ORAL			Fatigue				
			Headache				
			Nausea				
			Pharmaceutical Product				
			Complaint				
			Somnolence				

Date:07/12/04ISR Number: 4395464-6Report Type:Direct Company Report #CTU 222517  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	1 TABLET		Abnormal Behaviour	Wellbutrin Sr 200 Mg	PS		ORAL

Required	Aggression			
TWICE DAIL				
Intervention to	Apathy			
ORAL				
Prevent Permanent	Bipolar I Disorder	Wellbutrin Xl 300 Mg	SS	ORAL
1 TABLET				
Impairment/Damage	Crying			
DAILY ORAL				
	Depression			
	Disinhibition			
	Emotional Disorder			
	Fear			
	Hallucination			
	Impaired Work Ability			
	Insomnia			
	Libido Increased			
	Mania			
	Suicidal Ideation			
	Thinking Abnormal			

Date:07/13/04ISR Number: 4396284-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0504585A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alcohol Poisoning	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
		Contusion	Professional				
per day	5 MON						
		Dizziness		Alcohol	SS		
		Fall		Albuterol	C	Glaxosmithkline	
		Syncope		Advair	C	Glaxosmithkline	
				Elavil	C	Glaxosmithkline	
				Tylenol	C	Glaxosmithkline	



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Freedom Of Information (FOI) Report

Date:07/13/04ISR Number: 4396303-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0517688A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
900MG Per day		Adverse Event					
		Medication Error					

Date:07/13/04ISR Number: 4396716-6Report Type:Direct Company Report #CTU 222615

Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropion	PS		ORAL
Other		Arthralgia					
100 MG, Q AM							
ORAL							

Date:07/13/04ISR Number: 4396825-1Report Type:Direct Company Report #CTU 222710

Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr			
Other		Feeling Abnormal		(Generic) 150 Mg One			
		Pharmaceutical Product		Bid	PS		
ONE BID		Complaint					
		Somnolence					

Date:07/14/04ISR Number: 4396978-5Report Type:Expedited (15-DaCompany Report #PHEH2004US07298

Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Tegretol	PS	Novartis Sector:	
Death		Pulmonary Thrombosis				Pharma	ORAL
365 DAY		Road Traffic Accident					
				Tegretol	SS	Novartis Sector:	
dose						Pharma	ORAL

increased

(unspecified)

150 mg, BID 7200 MIN

Wellbutrin - Slow Release	SS	ORAL
Librium "Hoffmann"	SS	
Nortriptyline	C	
Valproate Semisodium	C	
Quetiapine Fumarate	C	
Trazodone	C	
Citalopram Hydrobromide	C	

Date:07/14/04ISR Number: 4397309-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511391A  
 Age:40 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG per day 17 DAY	Abortion Spontaneous	Health	Wellbutrin	PS	Glaxosmithkline	
Initial or Prolonged 180MG per day	Drug Exposure During	Professional	Allegra	C		
Other	Pregnancy Pregnancy Vaginal Haemorrhage		Natalcare Plus	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/14/04ISR Number: 4397313-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0518091A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Medication Error					
150MG per day				Wellbutrin Sr	SS	Glaxosmithkline	

Date:07/14/04ISR Number: 4397332-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0337748A

Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
Other		Feeling Hot					
150MG Unknown		Pruritus					
		Urticaria					

Date:07/14/04ISR Number: 4398071-4Report Type:Direct Company Report #CTU 222731

Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropion 150 Mg	PS	Teva	ORAL
Other		Anxiety		Teva			
1 PER DAY		Depression					
ORAL		Drug Effect Decreased					
		Libido Decreased					
		Pharmaceutical Product					
		Complaint					

Date:07/15/04ISR Number: 4398511-0Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0518045A

Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
Other		Dizziness					
150MG Twice							

per day 1 WK  
Hallucination  
Loss Of Consciousness  
Road Traffic Accident

Date:07/15/04ISR Number: 4398517-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0518217A  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3 MON	Feeling Abnormal Suicidal Ideation		Oral Contraceptive	C		

Date:07/15/04ISR Number: 4398530-4Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0337502A  
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Feeling Abnormal	Health	Zyban	PS	Glaxosmithkline	
6 DAY		Suicidal Ideation	Professional	Risperidal	C		

Date:07/15/04ISR Number: 4398540-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0337749A  
Age:61 YR Gender:Female I/FU:F

Outcome  
Life-Threatening  
Hospitalization -

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Initial or Prolonged Disability

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	23 DAY	Cerebral Infarction		Zyban	PS	Glaxosmithkline	ORAL
3.75MG Per day		Dysphagia					
		Facial Paresis		Zopiclone	C		ORAL
		Speech Disorder					

Date:07/15/04ISR Number: 4398543-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0338492A  
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG See dosage text		Anorexia		Zyban	PS	Glaxosmithkline	ORAL
		Arthralgia					
		Asthenia					
		Blood Immunoglobulin E Increased					
		Cough					
		Dysphagia					
		Dyspnoea					
		Joint Swelling					
		Oesophageal Spasm					
		Serum Sickness					
		Urticaria Generalised					

Date:07/15/04ISR Number: 4403016-4Report Type:Expedited (15-DaCompany Report #S04-USA-04018-01  
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other Required 10 MG QD PO Intervention to		Alopecia	Consumer	Lexapro (Escitalopram)	PS		ORAL
		Arthralgia					
		Erectile Dysfunction		Lotrel	SS		

Prevent Permanent  
0.2 MG BID  
Impairment/Damage

Exostosis  
Fracture  
Hypertension  
Joint Dislocation  
Limb Injury  
Nausea  
Somnolence

Clonidine SS  
Amiloride  
W/Hydrochlorothiazid  
e SS  
Percocet SS  
Wellbutrin - Slow  
Release (Bupropion  
Hydrochloride) SS  
  
Trazodone C  
Nexium  
(Esomeprazole) C  
Carisoprodol C  
Bextra (Valdecoxib) C

150 MG

Date:07/16/04ISR Number: 4399514-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441718A  
Age:33 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice	Abortion Spontaneous	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day 1 YR Other	Drug Exposure During Pregnancy	Professional				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/16/04ISR Number: 4399519-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511866A  
 Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day	64 DAY	Atrial Fibrillation	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	20MG Per day		Palpitations	Paxil	C	Glaxosmithkline	ORAL
UNKNOWN				Synthroid	C	Glaxosmithkline	
UNKNOWN				Zocor	C		
UNKNOWN				Tricor	C		
UNKNOWN				Allegra	C		

Date:07/16/04ISR Number: 4399525-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0517685A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	450MG Per day		Amphetamines Positive	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged			Convulsion	No Concurrent Medication	C		

Date:07/16/04ISR Number: 4399536-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0518553A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	6000MG Single		Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -	dose		Intentional Misuse				
Initial or Prolonged			Suicide Attempt	No Concurrent Medication	C		
Other							

Date:07/16/04ISR Number: 4399551-8Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0338333A  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Dizziness					
per day	10 DAY	Palpitations		Eltroxin	C	Glaxosmithkline	ORAL
.2MG per day				Triphasil	C		ORAL

Date:07/16/04ISR Number: 4399560-9Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0338426A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Autoimmune Hepatitis		Zyban	PS	Glaxosmithkline	ORAL
		Malaise		Zoloft	C		

Date:07/16/04ISR Number: 4399562-2Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0338632A  
Age:43 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	C-Reactive Protein
Initial or Prolonged	Increased
	Discomfort
	Dysphagia
	Exanthem
	Face Oedema
	Hyperhidrosis
	Hypersensitivity
	Oedema Peripheral



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	3 WK	Palpitations Paraesthesia Pruritus Vertigo White Blood Cell Count Increased	Zyban	PS	Glaxosmithkline	ORAL

Date:07/16/04ISR Number: 4402890-5Report Type:Expedited (15-DaCompany Report #S04-USA-04077-01  
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose Other 10 MG QD PO 450 MG QD	PT	Grand Mal Convulsion Health Professional	Lexapro (Escitalopram) Wellbutrin Xl	PS SS		ORAL

Date:07/19/04ISR Number: 4401665-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0148065A  
Age:20 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	9 DAY	Abasia Aggression Amnesia Asthenia Blindness Blood Urine Present Disability Eyelid Function Disorder Grand Mal Convulsion Hypoaesthesia Insomnia Loss Of Consciousness Mouth Haemorrhage Muscle Twitching Renal Impairment	Wellbutrin Birth Control Pills	PS C	Glaxosmithkline	ORAL

Tongue Biting  
Tremor  
Vision Blurred

Date:07/19/04ISR Number: 4401668-6Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0510698A  
Age:22 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice Initial or Prolonged per day	Anxiety  Decreased Appetite  Depression Disability Impaired Work Ability		Zyban	PS	Glaxosmithkline	ORAL

Date:07/19/04ISR Number: 4401671-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0518588A  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death	Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/20/04ISR Number: 4402278-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0518769A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Paxil	PS	Glaxosmithkline	
		Injury		Wellbutrin	SS	Glaxosmithkline	
		Injury Asphyxiation					
		Mood Swings					
		Physical Assault					
		Sexual Assault Victim					

Date:07/20/04ISR Number: 4402285-4Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0330195A

Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Myocardial Infarction	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
Hospitalization -		Overdose	Professional				
per day							
Initial or Prolonged		Somnolence		Prozac	SS		ORAL
20TABS per							
Other		Sudden Death					
day							

Date:07/20/04ISR Number: 4402297-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0338283A

Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysphagia		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	1 DAY	Throat Tightness					

Date:07/20/04ISR Number: 4402298-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0338286A

Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Dizziness	Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	Headache				
	Nausea				
	Oropharyngeal Swelling				
	Swelling Face				

Date:07/20/04ISR Number: 4402308-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0339189A  
 Age:56 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Grand Mal Convulsion		Zyban	PS	Glaxosmithkline	ORAL
150MG See						
Initial or Prolonged	Metastases To Central					
dosage text 11 DAY	Nervous System		Nicopatch	C	Glaxosmithkline	
TRANSDERMAL 14MG Per day						
160MG Per day 1 YR			Kardegic	C		ORAL
200MG Per day			Monotildiem	C	Glaxosmithkline	ORAL
UNKNOWN			Solupred	C	Glaxosmithkline	
UNKNOWN			Valium	C		

Date:07/21/04ISR Number: 4403565-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0518217A  
 Age:24 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 3 MON						
	Feeling Abnormal		Oral Contraceptive	C		
	Suicidal Ideation					

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Freedom Of Information (FOI) Report

Date:07/21/04ISR Number: 4403578-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0336338A

Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour		Zyban	PS	Glaxosmithkline	
1	MON	Aggression		No Concurrent			
		Anxiety		Medication	C		
		Depression					
		Stress					

Date:07/21/04ISR Number: 4404607-7Report Type:Direct

Company Report #CTU 223298

Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pharmaceutical Product		Generic Paxil 10 Mg			
ONE DOSE PO		Complaint		Po Qam	PS		ORAL
QAM				Generic Wellbutrin			
PO QAM				Sr	SS		ORAL

Date:07/21/04ISR Number: 4404668-5Report Type:Direct

Company Report #CTU 223135

Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG 1X DAY		Abdominal Pain		Wellbutrin 150 Mg 1x			
		Confusional State		Day	PS		
		Constipation					
		Dyspnoea					
		Erythema					
		Flatulence					
		Hyperhidrosis					
		Pharmaceutical Product					
		Complaint					

Date:07/22/04ISR Number: 4404716-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0515876A  
Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	9 WK	Depression		Oxycodone	C		
		Drug Ineffective		Xanax	C		
		Feeling Abnormal					
		Nervousness					
		Psychomotor Hyperactivity					
		Suicidal Ideation					

Date:07/22/04ISR Number: 4404726-5Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0519183A  
Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG per day		Loss Of Consciousness					

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Freedom Of Information (FOI) Report

Date:07/22/04ISR Number: 4404727-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0519196A  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG per day	Abortion Spontaneous		Bupropion	PS	Glaxosmithkline	

Date:07/22/04ISR Number: 4404728-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0519214A  
Age:19 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day		Cerebral Haemorrhage		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Grand Mal Convulsion		No Concurrent Medication	C		

Date:07/22/04ISR Number: 4405183-5Report Type:Direct Company Report #CTU 223366  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG 1X DAY		Abdominal Pain Upper		Wellbutrin	PS		
		Confusional State					
		Constipation					
		Dyspnoea Exertional					
		Erythema					
		Flatulence					
		Hyperhidrosis					
		Pharmaceutical Product Complaint					

Date:07/22/04ISR Number: 4408411-5Report Type:Expedited (15-DaCompany Report #2004IC000340  
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alcoholism Convulsion	Consumer	Librium Bupropion	PS		

150 MG; TWICE		Pulmonary Embolism		Hydrochloride	SS		ORAL
A DAY; ORA		Road Traffic Accident					
ORAL	1	YR	Somnolence	Carbamazepine	SS		ORAL
				Nortriptyline	C		
				Valproate Semisodium	C		
				Quetiapine Fumarate	C		
				Citalopram			
				Hydrobromide	C		

Date:07/23/04ISR Number: 4406045-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504887A  
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Avandia	PS	Glaxosmithkline	ORAL
4MG Per day	5 DAY	Vomiting		Paxil	SS	Glaxosmithkline	ORAL
10MG							
Alternate							
days				Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day	2 WK			Glucovance	C		
				Aspirin	C	Glaxosmithkline	
				Nexium	C		
				Vioxx	C		
				Ferrous Sulfate	C	Glaxosmithkline	
				Lamisil	C	Glaxosmithkline	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/23/04ISR Number: 4406262-9Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US010889

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Glucose Decreased		Avandia	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Interaction		Wellbutrin	SS	Glaxosmithkline	ORAL
per day				Provigil	SS		ORAL
850MG Three				Glucophage	SS		ORAL
times per day							
INTRAMUSCULAR	156UNIT Per			Humalog	SS		
day							
INTRAMUSCULAR	50UNIT Twice			Humulin	SS		
per day							
40MG Per day				Celexa	SS		ORAL
				Neurontin	C		
				Vioxx	C		
				Prilosec	C		
				Lisinopril	C		
				Allegra	C		

Date:07/23/04ISR Number: 4406300-3Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #12286316

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Interaction		Avandia	PS	Glaxosmithkline	ORAL
150MG Twice		Hypoglycaemia		Wellbutrin	SS	Glaxosmithkline	ORAL
per day				Provigil	SS		ORAL
40MG Per day				Celexa	SS		ORAL

850MG Three	Glucophage	SS	ORAL
times per day			
SUBCUTANEOUS	Humalog Insulin	SS	
SUBCUTANEOUS	Regular Insulin	SS	
	Neurontin	C	
	Vioxx	C	
	Prilosec	C	
	Allegra	C	
	Lisinopril	C	

Date:07/23/04ISR Number: 4406575-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0320766A  
Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
1TAB Twice							
per day	13	DAY		Ramipril	C		
UNKNOWN				Aspirin	C	Glaxosmithkline	
UNKNOWN				Diltiazem Hydrochloride	C	Glaxosmithkline	
UNKNOWN				Atorvastatin	C		
UNKNOWN				Elantan La	C		
UNKNOWN	50MG	Per day					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/23/04ISR Number: 4406576-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0321121A  
 Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
1TAB Twice							
per day	13	DAY					
UNKNOWN				Ramipril	C		
UNKNOWN				Aspirin	C	Glaxosmithkline	
UNKNOWN				Diltiazem Hydrochloride	C	Glaxosmithkline	
UNKNOWN				Atorvastatin	C		
UNKNOWN				Elantan La	C		

Date:07/23/04ISR Number: 4406587-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0337415A  
 Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Diabetes Mellitus		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day							
Other		Inadequate Control Hypoglycaemia		Insulin Humalog Mix50	C		
SUBCUTANEOUS	4IU Per day			Humalog	C		
SUBCUTANEOUS	2IU Twice per						
day							
SUBCUTANEOUS	6IU At night			Lantus	C		
				Creon 25000	C		ORAL
				Inipomp	C		ORAL
1G per day							

Date:07/23/04ISR Number: 4406604-4Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0339775A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1TAB per day Initial or Prolonged		Amnesia Blood Pressure Decreased Cardiac Flutter Chest Pain Feeling Abnormal Headache Hiatus Hernia Insomnia		Zyban	PS	Glaxosmithkline	ORAL

Date:07/26/04ISR Number: 4407826-9Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0335716A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG Per day 8 DAY 200MG per day		Aggression Confusional State Difficulty In Walking Disorientation Drug Interaction Dysphoria Headache Illogical Thinking Irritability	Health Professional	Zyntabac Sertraline Tofranil	PS C C	Glaxosmithkline	ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/26/04ISR Number: 4407838-5Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0338941A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Suicide Attempt		Zyban	PS	Glaxosmithkline	ORAL

Date:07/26/04ISR Number: 4408230-XReport Type:Direct Company Report #CTU 223526

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety Drug Ineffective		Bupropion Hydrochloride	PS	Teva	ORAL
1 PILL TWICE		Heart Rate Increased					
DAIL ORAL		Hypoventilation Insomnia Irritability Pharmaceutical Product Complaint Restlessness					

Date:07/26/04ISR Number: 4410635-8Report Type:Expedited (15-DaCompany Report #2004046963

Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Loss Of Consciousness	Health Professional	Doxepin (Caps) (Doxepin)	PS		
UNKNOWN	UNKNOWN	Nausea Sinus Tachycardia Tremor		Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		
UNKNOWN	UNKNOWN			Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C		
				Lorazepam (Lorazepam)	C		

Benazepril  
Hydrochloride  
(Benazepril  
Hydrochloride) C  
Atenolol (Atenolol) C  
Vicodin (Hydrocodone  
Bitartrate,  
Paracetamol) C  
Methadone  
(Methadone) C

Date:07/27/04ISR Number: 4408850-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0519165A  
Age:40 YR Gender:Female I/FU:I

Outcome PT  
Death Agitation  
Anxiety  
Bedridden  
Completed Suicide  
Diarrhoea  
Dizziness  
Feeling Abnormal  
Headache

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Insomnia Irritability Lethargy	Report Source	Product	Role	Manufacturer	Route
		Nausea Nervousness Pyrexia Social Avoidant Behaviour Toothache Tremor Vomiting		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:07/27/04ISR Number: 4408856-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0519542A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Nephrolithiasis					
2	MON						

Date:07/27/04ISR Number: 4408875-7Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0338333A  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Anxiety					
per day	10 DAY	Dizziness		Eltroxin	C	Glaxosmithkline	ORAL
.2MG per day		Palpitations		Triphasil	C		ORAL

Date:07/27/04ISR Number: 4411246-0Report Type:Expedited (15-DaCompany Report #B0339624A  
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Paxil (Paroxetine Hydrochloride)	PS		
Required Intervention to PER DAY	4 WK	Alanine Aminotransferase Increased	Foreign Literature				

Prevent Permanent Impairment/Damage			Aspartate Aminotransferase Increased	Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	SS
150 MG / PER DAY	1	WK	Carotid Artery Occlusion			
			Carotid Artery Thrombosis			
			Cerebral Infarction		Trazodone (Formulation Unknown) (Trazodone)	SS
			Epstein-Barr Virus Infection			
PER DAY	3	WK	Ischaemic Stroke		Lithium Carbonate	C
			Lymphocyte Morphology Abnormal		Lorazepam	C
			Splenomegaly		Cyproheptadine	C
					Oral Contraceptive	C

Date:07/27/04ISR Number: 4413039-7Report Type:Expedited (15-DaCompany Report #S04-USA-04077-01  
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Electroencephalogram Abnormal	Health Professional	Lexapro (Escitalopram)	PS		ORAL
10 MG QD PO							
		Grand Mal Convulsion		Wellbutrin Xl	SS		
450 MG QD							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/28/04ISR Number: 4409939-4Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0043950A

Age:21 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	ORAL
UNKNOWN		Hypoaesthesia Oral		Haldol	C		
UNKNOWN		Muscle Spasms		Atosil	C	Glaxosmithkline	

Date:07/29/04ISR Number: 4410975-2Report Type:Expedited (15-DaCompany Report #2004-03238

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia		Wellbutrin Sr	PS	Glaxosmithkline	
150MG Per day 10 YR		Depressed Level Of Consciousness		Bupropion Hydrochloride	SS	Glaxosmithkline	ORAL
150MG Twice		Disturbance In Attention					
per day	DAY	Motor Dysfunction		Lithium	C	Glaxosmithkline	ORAL
10 YR		Tremor					

Date:07/29/04ISR Number: 4410984-3Report Type:Expedited (15-DaCompany Report #PT-GLAXOSMITHKLINE-B0339717A

Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Vision Blurred	Health	Zyban	PS	Glaxosmithkline	ORAL
300MG per day			Professional				

Date:07/29/04ISR Number: 4411973-5Report Type:Direct Company Report #CTU 223864

Age:15 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization -	Convulsion	Wellbutrin Sr 100 Mg	PS	ORAL
100 MG PO BID				
Initial or Prolonged	Intentional Misuse			
	Suicide Attempt			

Date:07/29/04ISR Number: 4411975-9Report Type:Direct Company Report #CTU 223865  
 Age:15 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Intentional Misuse		Wellbutrin Sr 150 Mg	PS		ORAL
150 MG PO BID	2 YR						
Initial or Prolonged		Suicide Attempt					

Date:07/29/04ISR Number: 4415067-4Report Type:Expedited (15-DaCompany Report #S04-USA-04386-01  
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Thrombocytopenia	Health Professional	Lexapro (Escitalopram)	PS		ORAL
20 MG QD PO				Topamax (Topiramate)	SS		
100 MG QAM				Topamax (Topiramate)	SS		
200 MG QHS				Wellbutrin (Bupropion Hydrochloride)	SS		
200 MG BID				Topamax (Topiramate)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/29/04  
 ISR Number: 4444293-3  
 Report Type:Periodic  
 Age:69 YR Gender:Female I/FU:I

Company Report #USA-2003-0011039

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Morphine Sulfate			
				(Morphine Sulfate)	PS		
				Tylenol With Codeine			
				(Codeine Phosphae, Paracetamol)	SS		
				Bupropion			
				(Amfebutamide)	SS		
				Diphenhydramine			
				(Diphenhydramine)	SS		

Date:07/30/04  
 ISR Number: 4412081-X  
 Report Type:Expedited (15-Da  
 Age: Gender:Male I/FU:F  
 Company Report #US-GLAXOSMITHKLINE-A0511243A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG See dosage text	Completed Suicide		Wellbutrin Sr	PS	Glaxosmithkline	
		Decreased Appetite					
		Depressed Mood					
		Depression					
		Disturbance In Attention					
		Feelings Of Worthlessness					
		Sleep Disorder					
		Social Avoidant Behaviour					
		Suicidal Ideation					
		Tremor					

Date:07/30/04  
 ISR Number: 4412109-7  
 Report Type:Expedited (15-Da  
 Age:43 YR Gender:Male I/FU:F  
 Company Report #GB-GLAXOSMITHKLINE-B0339907A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 11 DAY		Condition Aggravated		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged 25MG per day		Eye Pruritus		Atenolol	C		ORAL

10MG per day	Eye Swelling	Ramipril	C	ORAL
SUBCUTANEOUS	Ocular Hyperaemia	Mixtard	C	
40MG per day	Oedema Peripheral	Pravastatin	C	ORAL
75MG per day	Rash Pruritic	Aspirin	C	Glaxosmithkline ORAL
INTRADERMAL	Urticaria	Niquitin Cq	C	Glaxosmithkline
	21MG per day			

Date:07/30/04ISR Number: 4412112-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0339975A

Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG		Hallucination, Auditory		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged Variable dose	DAY	Hallucination, Visual					
		Muscle Spasms Muscle Twitching Paraesthesia Suicidal Ideation Tinnitus		St Johns Wort	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/30/04ISR Number: 4412119-XReport Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0340435A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Twice			Cerebrovascular Accident	Zyban	PS	Glaxosmithkline	
per day			Hemiparesis				

Date:07/30/04ISR Number: 4412130-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0341038A

Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
150MG Twice			Angioneurotic Oedema	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged			Bronchospasm				
per day			Inflammation				
			Lymph Node Pain				
			Lymphadenopathy				
			Oedema				
			Painful Respiration				
			Pyrexia				
			Rash Maculo-Papular				
			Rash Pruritic				

Date:07/30/04ISR Number: 4412822-1Report Type:Direct Company Report #CTU 223929

Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required							
150 MG QAM			Condition Aggravated	Wellbutrin Sr 150 Mg	PS		ORAL
Intervention to			Muscle Spasms				
ORAL							
Prevent Permanent			Tremor	Atenolol	C		
Impairment/Damage				Hydroxyzine	C		
				Ranitidine	C		
				Valproic Acid	C		
				Tylenol # 3	C		
				Meperidine	C		

Date:07/30/04ISR Number: 4413014-2Report Type:Direct  
 Age:48 YR Gender:Female I/FU:I

Company Report #CTU 223955

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Tolerance Decreased		Wellbutrin Sr	PS		ORAL
150 MG BID PO		Pharmaceutical Product		Klonopin	SS		ORAL
0.5 MG BID PO		Complaint Unevaluable Event					

Date:08/02/04ISR Number: 4413068-3Report Type:Direct  
 Age:21 YR Gender:Male I/FU:I

Company Report #CTU 224046

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG BID Initial or Prolonged ORAL		Convulsion		Bupropion 100 Mg	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/02/04ISR Number: 4413146-9Report Type:Direct  
Age: Gender: I/FU:I

Company Report #CTU 224061

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr Glaxo	PS	Glaxo	ORAL
ORAL		Medication Error		Wellbutrin Xl			
				Glaxo	SS	Glaxo	ORAL
ORAL							

Date:08/02/04ISR Number: 4413162-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0520283A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Hemiparesis					
1 WK		Palpitations		Unknown Blood Pressure Medication	C		

Date:08/03/04ISR Number: 4414030-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0520112A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Life-Threatening		Convulsion					
300MG Per day		Joint Dislocation		No Concurrent Medication	C		
Hospitalization - Initial or Prolonged Disability							
Other							

Date:08/03/04ISR Number: 4414032-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0520465A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other		Aggression	Health Professional	Anti-Psychotic			
		Drug Interaction					

Psychotic Disorder

Medication

SS

UNKNOWN

Date:08/03/04ISR Number: 4414054-XReport Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0341295A

Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Sleep Disorder		Zyban	PS	Glaxosmithkline	

Date:08/04/04ISR Number: 4415228-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0520725A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Cardiac Arrest		Wellbutrin	PS	Glaxosmithkline	ORAL
60TAB Unknown							
Hospitalization -		Convulsion		Cocaine	C		
Initial or Prolonged		Hepatic Enzyme Increased		Marijuana	C		
Disability		Overdose					
Other							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/04/04ISR Number: 4415238-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0340500A  
Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Rash Psoriaform	Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	8 DAY						

Date:08/05/04ISR Number: 4416362-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0520475A  
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Abortion Spontaneous	Bupropion	PS	Glaxosmithkline	
300MG per day							

Date:08/05/04ISR Number: 4416363-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0520519A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Twice			Medication Error				
per day	1 DAY			Prozac	C		

Date:08/05/04ISR Number: 4416367-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0520969A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Bladder Pain	Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice			Micturition Urgency				
per day	3 MON		Nocturia	Synthroid	C	Glaxosmithkline	
			Pollakiuria				
			Urinary Incontinence				

Date:08/06/04ISR Number: 4417253-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0517688A  
Age:41 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
900MG Per day 4 DAY						
Initial or Prolonged	Disorientation		Depakote	C		ORAL
1500MG per						
day	Medication Error					

Date:08/06/04ISR Number: 4417255-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0520283A  
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
	Hemiparesis		Unknown Blood			
	Migraine		Pressure Medication	C		
	Muscle Twitching		Diovan	C		
	Nightmare		Aciphex	C		
	Palpitations		Zyrtec	C	Glaxosmithkline	
	Paraesthesia		Singulair	C		
	Sensation Of Heaviness		Vioxx	C		
	Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/06/04ISR Number: 4417319-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0517688A

Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	900MG Per day 4 DAY	Agitation	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged	1500MG per day	Disorientation	Professional	Depakote	C		ORAL
		Medication Error					

Date:08/06/04ISR Number: 4417321-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0520283A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Hemiparesis	Professional	Unknown Blood			
		Migraine		Pressure Medication	C		
		Muscle Twitching		Diovan	C		
		Nightmare		Aciphex	C		
		Palpitations		Zyrtec	C	Glaxosmithkline	
		Paraesthesia		Singulair	C		
		Sensation Of Heaviness		Vioxx	C		
		Tremor					

Date:08/09/04ISR Number: 4418144-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0423404A

Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	25MG Per day 2 DAY	Agitation		Paxil Cr	PS	Glaxosmithkline	ORAL
150MG Twice		Anxiety		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
per day	3 DAY	Dizziness					
		Panic Attack		Xanax	C		
		Psychomotor Hyperactivity					

Date:08/09/04ISR Number: 4418199-XReport Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0490948A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
6 MON		Abnormal Dreams		Paxil	PS	Glaxosmithkline	ORAL
150MG Twice		Diarrhoea		Wellbutrin	SS	Glaxosmithkline	ORAL
per day	1 WK	Drug Withdrawal Syndrome					
		Influenza Like Illness		Retin-A	C		
		Migraine					
		Nausea					
		Sensory Disturbance					
		Throat Tightness					

Date:08/09/04ISR Number: 4418476-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510014A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75MG Per day		Drug Ineffective		Paxil Cr	PS	Glaxosmithkline	ORAL
450MG Per day				Wellbutrin	SS	Glaxosmithkline	ORAL
				Wellbutrin	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/09/04ISR Number: 4418488-9Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510426A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
12.5MG Per day	2 YR	Irritability Tremor		Paxil Cr	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY	Weight Increased		Wellbutrin	SS	Glaxosmithkline	ORAL
				Strattera	C		
				Estratest	C		

Date:08/09/04ISR Number: 4418602-5Report Type:Periodic  
Age:52 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0413109A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Four times per day	YR	Agitation Alopecia	Consumer	Paxil Neurontin	PS SS	Glaxosmithkline	ORAL
150MG Per day	5 WK	Anger Drug Withdrawal Syndrome		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
		Dysphonia Hypersensitivity Influenza Like Illness Insomnia Nausea Paraesthesia Performance Status Decreased Smoker Somnolence		Trazodone Klonopin	C C		

Date:08/09/04ISR Number: 4418723-7Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0490907A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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20MG Per day	3	MON	Drug Ineffective	Paxil	PS	Glaxosmithkline	ORAL
			Weight Increased	Wellbutrin Sr	SS	Glaxosmithkline	ORAL

Date:08/09/04ISR Number: 4418727-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0490970A  
 Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash Pruritic		Paxil	PS	Glaxosmithkline	
				Wellbutrin	SS	Glaxosmithkline	ORAL

300MG Per day

Date:08/09/04ISR Number: 4419115-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0493495A  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety		Paxil Cr	PS	Glaxosmithkline	ORAL
40MG Per day	4	YR		Wellbutrin	SS	Glaxosmithkline	ORAL
		Weight Decreased					
2	MON	Weight Increased		Albuterol	C	Glaxosmithkline	
				Wellbutrin	C	Glaxosmithkline	
				Flovent	C	Glaxosmithkline	
				Alphagan	C		
				Zyprexa	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/09/04ISR Number: 4419512-XReport Type:Periodic  
 Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497213A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 YR		Drug Ineffective		Paxil	PS	Glaxosmithkline	ORAL
1 YR		Insomnia		Wellbutrin	SS	Glaxosmithkline	ORAL
				Lithium	C	Glaxosmithkline	
				Other Medications	C		

Date:08/09/04ISR Number: 4419629-XReport Type:Periodic  
 Age:47 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0500586A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
30MG Per day		Urticaria		Paxil	PS	Glaxosmithkline	
150MG Twice				Wellbutrin Sr	SS	Glaxosmithkline	ORAL
per day				Unspecified Medication	SS		

Date:08/09/04ISR Number: 4419820-2Report Type:Periodic  
 Age:80 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0502389A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
10MG Per day		Agitation		Paxil	PS	Glaxosmithkline	ORAL
150MG Twice		Anxiety		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
per day	2 WK	Blood Pressure Increased					
300MG Per day	2 DAY	Confusional State		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Crying		Levoxyl	C	Glaxosmithkline	
		Decreased Appetite					
		Delusion					
		Depression					

Dizziness  
 Drug Withdrawal Syndrome  
 Gastrointestinal Pain  
 Headache  
 Hot Flush  
 Hyperchlorhydria  
 Insomnia  
 Nausea  
 Nervousness  
 Palpitations  
 Vomiting  
 Weight Increased

Date:08/09/04ISR Number: 4420418-0Report Type:Periodic  
 Age:53 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0499806A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Energy Increased		Paxil	PS	Glaxosmithkline	ORAL
25MG Per day		Insomnia		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
300MG Per day	5 DAY	Loss Of Libido		Xanax	C		
				Herbal Supplement	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/09/04ISR Number: 4420518-5Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503861A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
12.5MG Per		Haematuria		Paxil Cr	PS	Glaxosmithkline	ORAL
day	7 WK	Hypertension					
		Pollakiuria		Wellbutrin	SS	Glaxosmithkline	ORAL

Date:08/09/04ISR Number: 4420543-4Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504593A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
50MG Per day	8 YR	Anxiety		Paxil Cr	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Ineffective		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
per day		Paraesthesia					
		Weight Increased					

Date:08/09/04ISR Number: 4420553-7Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504905A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
12.5MG Per		Urticaria		Paxil Cr	PS	Glaxosmithkline	ORAL
day	4 DAY	Weight Decreased					
300MG Per day	11 DAY			Wellbutrin	SS	Glaxosmithkline	ORAL
				Oral Contraceptives	C		

Date:08/09/04ISR Number: 4420611-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506505A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Rash		Paroxetine	PS	Glaxosmithkline	ORAL
40MG Per day	2	MON			Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Twice								
per day	2	MON			Magnesium	C		
					Calcium	C		
					Albuterol	C	Glaxosmithkline	
					Zinc	C		

Date:08/09/04ISR Number: 4420671-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0507901A  
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Agitation		Paxil	PS	Glaxosmithkline	ORAL
20MG Per day	13	YR	Anger		Paxil Cr	SS	Glaxosmithkline	ORAL
12.5MG Per			Insomnia					
day	2	WK	Muscle Twitching		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Per day	1	MON	Negative Thoughts					

Date:08/09/04ISR Number: 4420672-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0507905A  
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Agitation		Paxil	PS	Glaxosmithkline	ORAL
30MG Per day	7	YR	Confusional State		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
150MG Per day	8	MON	Disturbance In Attention					
			Memory Impairment					
			Weight Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/09/04ISR Number: 4420755-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511060A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	4 DAY	Fatigue		Paxil	PS	Glaxosmithkline	
		Nausea		Wellbutrin	SS	Glaxosmithkline	

Date:08/09/04ISR Number: 4420757-3Report Type:Periodic  
Age:68 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0511097A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dissociation		Paxil	PS	Glaxosmithkline	ORAL
		Drug Ineffective		Wellbutrin	SS	Glaxosmithkline	ORAL
		Somnolence					

Date:08/09/04ISR Number: 4420763-9Report Type:Periodic  
Age:73 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511300A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	4 DAY	Balance Disorder		Paxil	PS	Glaxosmithkline	ORAL
		Dizziness		Wellbutrin	SS	Glaxosmithkline	ORAL
		Eructation		Lorazepam	C		
		Flatulence		Lipitor	C		
		Insomnia		Aciphex	C		
		Weight Increased		Ambien	C		

Date:08/09/04ISR Number: 4420799-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513053A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
10MG Per day		Constipation		Paxil	PS	Glaxosmithkline	ORAL
150MG Per day	24 DAY	Diarrhoea		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Drug Ineffective		Lipitor	C		

Hot Flush	Paxil	C	Glaxosmithkline
Insomnia	Tenuate	C	
Nightmare	Xanax	C	
Tension			

Date:08/09/04ISR Number: 4420993-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514128A  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Flight Of Ideas	Professional	No Concurrent			
WK		Suicidal Ideation		Medication	C		

Date:08/09/04ISR Number: 4420994-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0516903A  
Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Death	Health	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
Death		Overdose	Professional				
150MG Twice				Zyprexa	C		
per day	2 WK			Lito	C	Glaxosmithkline	
				Depakote	C		
				Effexor	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/09/04ISR Number: 4421000-1Report Type:Expedited (15-DaCompany Report #2004-03363

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged		Cardiac Disorder Dermatitis Exfoliative Medication Error Respiratory Disorder Skin Reaction		Bupropion	PS	Glaxosmithkline	ORAL

Date:08/09/04ISR Number: 4421007-4Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0336916A

Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1TAB See dosage text	15 DAY	Arthralgia Fibromyalgia Suicidal Ideation Thinking Abnormal	Consumer	Zyban Morphine	PS C	Glaxosmithkline	

Date:08/09/04ISR Number: 4421017-7Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0340435A

Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Cerebrovascular Accident Hemiparesis	Health Professional	Zyban	PS	Glaxosmithkline	

Date:08/09/04ISR Number: 4421751-9Report Type:Direct Company Report #CTU 224598

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
BID		Anxiety Apathy		Bupropion Sr 150mg Tab	PS		

Nausea  
Pharmaceutical Product  
Complaint

Climara Patch

C

Date:08/09/04ISR Number: 4423198-8Report Type:Direct  
Age:26 YR Gender:Female I/FU:I

Company Report #CTU 224502

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fatigue		Budeprion 150 Mg			
ONE PILL		Fluid Retention		Teva	PS	Teva	ORAL
TWICE A DA		Impatience					
ORAL		Irritability					
		Mood Swings					
		Pharmaceutical Product					
		Complaint					
		Weight Increased					

Date:08/09/04ISR Number: 4425658-2Report Type:Expedited (15-DaCompany Report #S04-USA-04077-01  
Age:43 YR Gender:Female I/FU:F

Outcome	PT
Other	Amnesia
	Coma
	Confusional State
	Eye Rolling

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Grand Mal Convulsion  
Tongue Biting

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
10 MG QD PO		Health Professional	Lexapro (Escitalopram)	PS		ORAL
450 MG QD			Wellbutrin Xl	SS		

Date:08/10/04ISR Number: 4422310-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0520977A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	450MG Per day 3	MON		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Convulsion		Doxepin	C		
		Liver Disorder		Clonazepam	C		
		Migraine		Chloral Hydrate	C		

Date:08/10/04ISR Number: 4422323-2Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0337087A  
Age:18 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Twice			Zyban	PS	Glaxosmithkline	
	per day	Laceration					
		Menarche		Alcohol	C		
		Postictal State					

Date:08/10/04ISR Number: 4422324-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0338405A  
Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	12 DAY			Zyban	PS	Glaxosmithkline	ORAL
		Orbital Oedema					

Initial or Prolonged Rash Papular  
 SUBCUTANEOUS 40UNIT Twice  
 Other  
 per day 14 YR

Mixtard C  
 Pravastatin C ORAL  
 Atenolol C ORAL  
 Aspirin C Glaxosmithkline ORAL  
 Ramipril C ORAL

Date:08/10/04ISR Number: 4422325-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0338704A  
 Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Enteritis		Zyban	PS	Glaxosmithkline	ORAL
1TAB Per day		Feeling Abnormal		Antidepressant	C		
		Nausea		Inhalers	C		
		Nervousness					
		Vomiting					

Date:08/10/04ISR Number: 4422328-1Report Type:Expedited (15-DaCompany Report #238104  
 Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN		Erythema		Zyban	PS	Glaxosmithkline	
Initial or Prolonged		Eye Swelling		Mixtard	SS		
SUBCUTANEOUS	80IU per day	Feeling Hot		Niquitin Cq	C	Glaxosmithkline	
TRANSDERMAL	21MG per day	Pruritus		Pravastatin	C		
UNKNOWN		Urticaria		Aspirin	C	Glaxosmithkline	
UNKNOWN							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

UNKNOWN

Atenolol C

UNKNOWN

Ramipril C

Date:08/10/04ISR Number: 4423252-0Report Type:Direct  
Age:24 YR Gender:Female I/FU:I

Company Report #CTU 224634

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Libido		Budeprion Teva	PS	Teva	ORAL
150 MG ONCE							
PER D ORAL		Nausea					
		Pharmaceutical Product Complaint					

Date:08/10/04ISR Number: 4424136-4Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 224679

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Bupropion Sr	PS		ORAL
150 MG PO BID							
		Pharmaceutical Product Complaint					

Date:08/10/04ISR Number: 4424170-4Report Type:Direct  
Age:36 YR Gender:Female I/FU:I

Company Report #CTU 224669

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fatigue		Budeprion Sr	PS		ORAL
150 MG PO BID							
		Insomnia					
		Irritability					
		Pharmaceutical Product Complaint					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea		Wellbutrin Sr 150			
		Pharmaceutical Product		Mg-Generic	PS		ORAL
150 MG PO BID		Complaint					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
WK		Flight Of Ideas	Professional	No Concurrent			
		Suicidal Ideation		Medication	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
		Fall					
		Tooth Fracture					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/11/04ISR Number: 4423295-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0521578A  
 Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 450MG Unknown		Contusion Convulsion		Wellbutrin Sr Wellbutrin Xl	PS SS	Glaxosmithkline Glaxosmithkline	ORAL ORAL
		Drug Ineffective Grand Mal Convulsion Insomnia Postictal State		Marijuana	C		

Date:08/11/04ISR Number: 4423299-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0521633A  
 Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day 3 WK	1 MON	Drug Ineffective Medication Error Pharmaceutical Product Complaint		Wellbutrin Bupropion Sr Zoloft	PS SS C	Glaxosmithkline Glaxosmithkline	ORAL

Date:08/11/04ISR Number: 4423303-3Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0336371A  
 Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG See dosage text	98 DAY	Activities Of Daily Living Impaired Affect Lability Anxiety Crying Depressed Mood Mental Disorder Nicotine Dependence Sleep Disorder		Zyban No Concurrent Medication	PS C	Glaxosmithkline	

Date:08/11/04ISR Number: 4423305-7Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0340435A  
Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Accident	Health	Zyban	PS	Glaxosmithkline	
150MG Twice		Hemiparesis	Professional				
per day							

Date:08/11/04ISR Number: 4423307-0Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0341295A  
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Zyban	PS	Glaxosmithkline	ORAL
5 WK		Dizziness					
		Fatigue					
		Sleep Disorder					

Date:08/11/04ISR Number: 4423315-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0341680A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	
33 DAY		Nightmare		Cipramil	C		
		Verbal Abuse					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/11/04ISR Number: 4423319-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0341933A  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 17 Initial or Prolonged	17 DAY	Agitation Anxiety Confusional State Trance		Zyban	PS	Glaxosmithkline	ORAL

Date:08/11/04ISR Number: 4425179-7Report Type:Direct Company Report #CTU 224756  
 Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
40 MG PO DAILY 150 MG PO Q DAY		Drug Interaction Mental Status Changes Tremor		Escitalopram Bupropion	PS SS		ORAL ORAL
				Benazepril Metronidazole Benicar Advair Morphine Ir	C C C C C		

Date:08/12/04ISR Number: 4424463-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0509451A  
 Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
135 DAY 32 DAY 100MG per day 50MG At night		Amnesia Contusion Convulsion Fall	Health Professional	Wellbutrin Trileptal Lamictal Benadryl	PS SS C C	Glaxosmithkline Glaxosmithkline Glaxosmithkline	ORAL ORAL ORAL

4TAB Per day  
Head Injury  
Medication Error  
Sedation  
Nicorette  
C  
Glaxosmithkline  
ORAL

Date:08/12/04ISR Number: 4424485-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0341269A  
Age:75 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anorexia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Decreased Interest					
per day	5	WK					
		Depression		Rabeprazole	C		ORAL
10MG per day		Dizziness					
		Fatigue					
		Tremor					

Date:08/12/04ISR Number: 4425302-4Report Type:Direct Company Report #CTU 224803  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tinnitus		Wellbutrin Xl	PS		
150 MG DAILY							
X 1 WK 300 MG							
DAILY X 3 WKS							

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Freedom Of Information (FOI) Report

Date:08/12/04ISR Number: 4426610-3Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #CTU 224843

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Wellbutrin Sr 150 Glaxosmithkline	PS	Glaxosmithkline	

Date:08/13/04ISR Number: 4425370-XReport Type:Expedited (15-DaCompany Report #2004-03222  
 Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression Ill-Defined Disorder		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
150MG Per day		Muscular Weakness		Paxil	C	Glaxosmithkline	ORAL
YR		Myalgia		Topamax	C		
YR		Pharmaceutical Product		Altace	C		
YR		Complaint		Zocor	C		
YR		Suicidal Ideation					

Date:08/13/04ISR Number: 4425377-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0521279A  
 Age:6 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination		Wellbutrin	PS	Glaxosmithkline	ORAL
37.5MG Per		Medication Error					
day	1 WK			Strattera	C		
				Clonidine	C		

Date:08/13/04ISR Number: 4425378-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0521306A  
 Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	75MG Four times per day Initial or Prolonged	Coma		Wellbutrin	PS	Glaxosmithkline	ORAL
		Death		No Concurrent Medication	C		

Date:08/13/04ISR Number: 4425379-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0521407A  
 Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Flushing Headache Medication Error		Wellbutrin Xl Ativan	PS C	Glaxosmithkline	ORAL ORAL

Date:08/13/04ISR Number: 4427245-9Report Type:Direct Company Report #CTU 224931  
 Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Grand Mal Convulsion Skin Laceration		Wellbutrin	PS		



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Freedom Of Information (FOI) Report

Date:08/13/04ISR Number: 4427264-2Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 224940

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Congenital Anomaly	Coloboma		Wellbutrin	PS		
	Congenital Eye Disorder		Prozac	SS		
	Cryptophthalmos					
	Drug Exposure During					
	Pregnancy					

Date:08/13/04ISR Number: 4448179-XReport Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #T04-USA-03512-01

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Anxiety	Health	Namenda (Memantine)	PS		ORAL
20 MG BID PO						
Initial or Prolonged	Aphasia	Professional	Wellbutrin			
	Convulsion		(Bupropion			
	Fall		Hydrochloride)	SS		ORAL
150 MG BID PO						
			Exelon (Rivastigmine			
			Tartrate)	C		
			Prempro	C		
			Paxil (Paroxetine			
			Hydrochloride)	C		

Date:08/16/04ISR Number: 4426806-0Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0521802A  
Age:28 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Abdominal Pain		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice						
per day	Rectal Haemorrhage					

Date:08/16/04ISR Number: 4426807-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0521812A  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Medication Error		Lexapro	SS		
		Somnolence		Clonazepam	SS		

Date:08/16/04ISR Number: 4426816-3Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0334041A

Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Zyban	PS	Glaxosmithkline	ORAL
Other		Delirium Tremens					
1 DAY		Hallucination					
		Overdose					
		Restlessness					
		Tardive Dyskinesia					

Date:08/16/04ISR Number: 4426826-6Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0341325A

Age:49 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Chest Pain
Initial or Prolonged	Coronary Artery Surgery
	Triple Vessel Bypass

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Graft

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day			Zyban	PS	Glaxosmithkline	
50MG per day			Atenolol	C		ORAL
400MG per day			Trental	C		ORAL
81MG per day			Ecotrin	C	Glaxosmithkline	ORAL
4MG per day			Coversyl Plus	C		ORAL

Date:08/16/04ISR Number: 4426838-2Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0342217A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Ankyloglossia Congenital Anorexia Dizziness Flatulence Middle Insomnia Reading Disorder Sleep Disorder Somnolence Thought Blocking Tremor Urticaria Visual Acuity Reduced Visual Disturbance		Zyban	PS	Glaxosmithkline	

Date:08/16/04ISR Number: 4426839-4Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0342260A  
Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1TAB Variable		Feeling Abnormal		Zyban	PS	Glaxosmithkline	

dose  
Mental Disorder  
Personality Change  
Psychotic Disorder

Date:08/17/04ISR Number: 4427736-0Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0318010A  
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1TAB Twice Initial or Prolonged per day	5 DAY	Aggression Agitation	Health Professional	Zyban	PS	Glaxosmithkline	ORAL
Other		Blood Pressure Increased Depression Dizziness Feeling Abnormal Headache Pharmaceutical Product Complaint					

Date:08/17/04ISR Number: 4427747-5Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0341325A  
Age:49 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Chest Pain Coronary Artery Surgery

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Freedom Of Information (FOI) Report

		Triple Vessel Bypass Graft	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Health Professional	Zyban	PS	Glaxosmithkline	
150MG Twice per day				Atenolol	C		ORAL
50MG per day				Trental	C		ORAL
400MG per day				Ecotrin	C	Glaxosmithkline	ORAL
81MG per day				Coversyl Plus	C		ORAL
4MG per day							

Date:08/17/04ISR Number: 4427758-XReport Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0342487A  
 Age: Gender:Male I/FU:F

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Cardiac Disorder		Zyban	PS	Glaxosmithkline	ORAL
Dose							
Hospitalization - Initial or Prolonged							

Date:08/17/04ISR Number: 4429450-4Report Type:Direct Company Report #CTU 225143  
 Age:53 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Condition Aggravated Depression		Wellbutron Sr (Generic)	PS		ORAL
Dose							
Other		Drug Effect Decreased					
120 MG 2 PO				Trazadone	C		
BID		Pharmaceutical Product Complaint		Xanax	C		

Date:08/17/04ISR Number: 4430036-6Report Type:Expedited (15-DaCompany Report #S04-USA-04732-01  
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abnormal Behaviour Agitation	Consumer	Lexapro (Escitalopram)	PS		ORAL
20 MG QD PO		Belligerence Confusional State		Lexapro (Escitalopram)	SS		ORAL
10 MG QD PO		Delusion Insomnia		Lexapro (Escitalopram)	SS		ORAL
10 MG QD PO		Mania Oral Intake Reduced Paranoia		Wellbutrin (Bupropion Hydrochloride)	SS		
300 MG QD		Speech Disorder		Wellbutrin (Bupropion Hydrochloride)	SS		
150 MG QD				Ambien (Zolpidem Tartrate)	C		
				Trazodone	C		
				Lipitor (Atorvastatin)	C		
				Achiphex (Rabeprazole Sodium)	C		
				Aspirin	C		

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Freedom Of Information (FOI) Report

Date:08/18/04ISR Number: 4428823-3Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0339775A  
 Age:58 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 1TAB per day Initial or Prolonged	Amnesia	Health	Zyban	PS	Glaxosmithkline	ORAL
	Blood Pressure Decreased Cardiac Flutter Chest Pain Feeling Abnormal Headache Hiatus Hernia Insomnia	Professional				

Date:08/18/04ISR Number: 4430053-6Report Type:Direct Company Report #CTU 225159  
 Age:65 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 1 TABLET BY MOUTH TWICE DAILY	Depression Movement Disorder Muscular Weakness Pharmaceutical Product Complaint Suicidal Ideation		Bupropion Sr 150 Mg Ta Wat	PS	Wat	

Date:08/18/04ISR Number: 4431278-6Report Type:Expedited (15-DaCompany Report #B0299104A  
 Age:25 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Myoclonic Epilepsy	Foreign Literature Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Disability	Activities Of Daily Living Impaired Convulsion Drug Interaction Drug Withdrawal Syndrome	Health Professional	Efexor Er (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		ORAL
75 MG 1X PER 11DAY	Flight Of Ideas					
UNKNOWN DOSE TWO TIMES PER DAY	Impaired Driving Ability Irritability Memory Impairment Petit Mal Epilepsy Status Epilepticus		Zyban (Amfebutamone Hydrochloride, )	SS		ORAL

Outcome	PT
Other	Anxiety Drug Hypersensitivity Drug Ineffective Muscle Spasms Pharmaceutical Product



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Dose	Duration	Complaint Post Procedural Complication Tremor	Report Source	Product	Role	Manufacturer	Route
2400 MG (800 MG, 3 IN 1 D), ORAL		Weight Decreased Weight Increased	Consumer Health  Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
				Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		
				Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Baclofen (Baclofen)	C		

Date:08/19/04ISR Number: 4429532-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514884A  
Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day	14 DAY	Back Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged 50MG Twice		Chest Pain		Ultram	SS		ORAL
per day	20 DAY	Dizziness Feeling Abnormal Feeling Jittery Grand Mal Convulsion Headache Nausea					

Date:08/19/04ISR Number: 4429536-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0521578A  
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 450MG Per day		Blood Pressure Diastolic Increased		Wellbutrin Sr Wellbutrin Xl	PS SS	Glaxosmithkline Glaxosmithkline	ORAL ORAL
		Contusion Convulsion Drug Ineffective Grand Mal Convulsion Insomnia Postictal State		Marijuana	C		

Date:08/19/04ISR Number: 4429544-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0522437A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG Per day	4 WK	Convulsion Glossodynia		Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	ORAL

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Date:08/19/04ISR Number: 4430640-5Report Type:Direct  
Age:40 YR Gender:Male I/FU:I

Company Report #CTU 225302

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
TID		Depression		Bupropion Hcl 150 Mg	PS		

Date:08/20/04ISR Number: 4430240-7Report Type:Expedited (15-DaCompany Report #2004-03363  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged		Cardiac Disorder Dermatitis Exfoliative Medication Error Polytraumatism Respiratory Disorder Skin Reaction	Other	Bupropion	PS	Glaxosmithkline	ORAL

Date:08/20/04ISR Number: 4430249-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0338405A  
Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 12 DAY Initial or Prolonged		Erythema	Health	Zyban	PS	Glaxosmithkline	ORAL
SUBCUTANEOUS 40UNIT		Orbital Oedema	Professional	Mixtard	SS		
Other per day	14 YR	Rash Papular					
		Skin Degenerative Disorder		Pravastatin	C		ORAL
				Atenolol	C		ORAL
2 YR		Urticaria		Aspirin	C	Glaxosmithkline	ORAL
2 YR				Ramipril	C		ORAL
2 YR				Niquitin Cq	C	Glaxosmithkline	
TRANSDERMAL	21MG per day						

Date:08/20/04ISR Number: 4430255-9Report Type:Expedited (15-DaCompany Report #238104  
Age:43 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - UNKNOWN	Erythema	Consumer	Zyban	PS	Glaxosmithkline	
Initial or Prolonged SUBCUTANEOUS 80IU per day	Eye Swelling		Mixtard	SS		
Feeling Hot			Niquitin Cq	C	Glaxosmithkline	
TRANSDERMAL 21MG per day UNKNOWN	Pruritus		Pravastatin	C		
UNKNOWN	Urticaria		Aspirin	C	Glaxosmithkline	
UNKNOWN			Atenolol	C		
UNKNOWN			Ramipril	C		
UNKNOWN						

Date:08/20/04ISR Number: 4430262-6Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0342245A  
Age:49 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Per day 27 DAY Initial or Prolonged	Psoriasis		Zyban	PS	Glaxosmithkline	ORAL

Date:08/20/04ISR Number: 4430265-1Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0342289A  
Age:29 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Twice	Confusional State		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day 3 WK	Fall					
UNKNOWN 10MG per day	Lower Limb Fracture		Stilnoct	C		
UNKNOWN 30MG per day	Pelvic Fracture		Cipralext	C		

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UNKNOWN	100MG per day	Imigran	C	Glaxosmithkline
UNKNOWN	25MG per day	Atarax	C	
UNKNOWN	5MG per day	Stesolid	C	
UNKNOWN	5MG per day	Nitrazepam	C	
UNKNOWN		Desolett	C	
UNKNOWN	50MG per day	Diclofenac	C	
UNKNOWN		Propylene Glycol	C	

Date:08/20/04ISR Number: 4432489-6Report Type:Expedited (15-DaCompany Report #A-US2004-07003  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SEE IMAGE		Abdominal Pain Upper Anorexia	Study Health	Tracleer (Bosentan) Tablet	PS		ORAL
Other ORAL		Antibody Test Positive	Professional				
		Autoimmune Disorder Constipation	Distributor	Lovastatin (Lovastatin)	SS		
20 MG, QD		Hepatitis International Normalised Ratio Increased Nephrolithiasis		Wellbutrin (Amfebutamone Hydrochloride) Coumadin (Warfarin Sodium) Prinivil (Lisinopril) Aspirine (Acetylsalicylic Acid) Milk Of Magnesia (Magnesium Hydroxide)	SS   C C  C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged SEE IMAGE	Antibody Test Positive Hepatitis	Study Health	Tracleer(Bosentan) Tablet	PS		ORAL
Other 20 MG, QD	International Normalised Ratio Increased  Nephrolithiasis	Professional Distributor	Lovastatin (Lovastatin)  Wellbutrin(Amfebutam one Hydrochloride0 Coumadin (Warfarin Sodium) Prinivil (Lisinopril) Aspirine (Acetylsalicylic Acid0 Milk Of Magnesia (Magnesium Hydroxide)	SS  SS C C C C		

Outcome  
Hospitalization -  
Initial or Prolonged  
  
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Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Pain Anorexia	Study Health	Tracleer (Bosentan) Tablet	PS		ORAL
SEE IMAGE		Antibody Test Positive Constipation	Professional Distributor	Lovastatin (Lovastatin)	SS		
20 MG, QD		Hepatitis International Normalised Ratio Increased Nephrolithiasis		Wellbutrin (Amfebutamone Hydrochloride) Coumadin (Warfarin Sodium) Prinivil (Lisinopril) Aspirine (Acetylsalicylic Acid) Milk Of Magnesia (Magnesium Hydroxide)	SS C C C C		

Date:08/23/04ISR Number: 4430847-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0515876A  
Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Depression	Professional	Oxycodone	C		
300MG Per day	9 WK	Drug Ineffective Feeling Abnormal Nervousness Psychomotor Hyperactivity Suicidal Ideation		Xanax	C		

Date:08/23/04ISR Number: 4430848-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0520969A  
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other 100MG Twice		Bladder Pain	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3	MON	Micturition Urgency	Professional			
			Nocturia	Synthroid	C	Glaxosmithkline	
			Pollakiuria				
			Urinary Incontinence				

Date:08/23/04ISR Number: 4430850-7Report Type:Expedited (15-DaCompany Report #S04--USA-04732-01  
Age:61 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day	5 YR	Abnormal Behaviour		Wellbutrin	PS	Glaxosmithkline	ORAL
	10MG Per day	2 MON	Agitation		Lexapro	SS		ORAL
			Belligerence		Ambien	C		ORAL
			Confusional State		Trazodone	C		
UNKNOWN	25MG Per day		Delusion		Lipitor	C		
UNKNOWN			Mania		Aciphex	C		
UNKNOWN			Paranoia		Aspirin	C	Glaxosmithkline	
UNKNOWN								



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Date:08/23/04ISR Number: 4430858-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0522920A  
 Age:10 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2550MG Single Initial or Prolonged dose	1 DAY			Wellbutrin	PS	Glaxosmithkline	ORAL
		Grand Mal Convulsion Overdose Visual Disturbance					

Date:08/23/04ISR Number: 4430873-8Report Type:Expedited (15-DaCompany Report #FI-GLAXOSMITHKLINE-B0342423A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1TAB Variable Initial or Prolonged dose	9 DAY			Zyban	PS	Glaxosmithkline	ORAL
47.5MG Per day	5 YR	Eczema Vesicular Eye Oedema Hypersensitivity		Seloken Zoc	C		ORAL
20MG Per day	10 YR	Liver Function Test Abnormal Pharyngolaryngeal Pain Pyrexia Urticaria		Linatil	C		ORAL

Date:08/23/04ISR Number: 4447597-3Report Type:Periodic Company Report #PHEH2004US06923  
 Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 MG, QD, ORAL		Drug Interaction Palpitations Throat Tightness	Consumer	Zelnorm (Tegaserod) Tablet	PS		ORAL
				Wellbutrin - Slow			

Release (Bupropion  
Hydrochloride) SS

200 MG, 6QD

Date:08/24/04ISR Number: 4431406-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514321A  
Age:25 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 12.5G Single Initial or Prolonged dose	Coma		Wellbutrin	PS	Glaxosmithkline	ORAL
Disability 120MG Single Other dose	Overdose		Paxil	SS	Glaxosmithkline	
30MG Single dose			Ativan	SS		
			Vitamins Minerals	C C		

Date:08/24/04ISR Number: 4431421-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0522921A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Bulimia Nervosa Oesophageal Varices Haemorrhage		Wellbutrin	PS	Glaxosmithkline	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/24/04ISR Number: 4431422-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0522931A  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Twice			Blood Glucose Increased	Wellbutrin	PS	Glaxosmithkline	ORAL
			Convulsion				
per day	1	WK	Fall	Ludiomil	C		
			Upper Limb Fracture	Unknown Medication	C		

Date:08/24/04ISR Number: 4431424-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0523061A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Twice			Medication Error	Wellbutrin	PS	Glaxosmithkline	ORAL
per day							

Date:08/24/04ISR Number: 4431437-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0342585A  
Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Per day	12	DAY	Drug Interaction	Zyban	PS	Glaxosmithkline	ORAL
			Haemorrhage	Marvelon	SS		ORAL
180MG Per day				Fexofenadine	C		ORAL

Date:08/24/04ISR Number: 4433260-1Report Type:Expedited (15-DaCompany Report #A0393333A  
Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death							
			Blood Ph Decreased	Wellbutrin Sr			
			Blood Pressure Systolic	Tablet-Controlled			
			Increased	Release (Bupropion			
			Literature				
			Health				
			Professional				

SEE DOSAGE                      Cardio-Respiratory Arrest                      Hydrochloride)                      PS

TEXT                      Coma

- Completed Suicide
- Electrolyte Imbalance
- Grand Mal Convulsion
- Heart Rate Increased
- Hepatic Enzyme Increased
- Hyperthermia
- Overdose
- Renal Failure

Date:08/24/04    ISR Number: 4433764-1    Report Type:Direct                      Company Report #CTU 225531  
 Age:              Gender:Female              I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Wellbutrin Sr 150mg			
		Feeling Abnormal		Bid	PS		ORAL
150MG 1 BID		Nausea					
PO		Pharmaceutical Product					
		Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/25/04ISR Number: 4432378-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0522795A

Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	2	YR	Depressed Mood	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day	2	MON	Ear Pain	Bupropion	SS	Glaxosmithkline	ORAL
			Headache				
			Neck Pain				
			Photophobia	Albuterol	C	Glaxosmithkline	
			Suicidal Ideation	Lexapro	C		
				Synthroid	C	Glaxosmithkline	
				Vioxx	C		
				Protonix	C		

Date:08/25/04ISR Number: 4432397-0Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0341359A

Age:22 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	10	DAY	Delirium	Zyban	PS	Glaxosmithkline	
			Dizziness				
			Epilepsy	No Concurrent Medications	C		
			Foaming At Mouth				
			Grand Mal Convulsion				
			Insomnia				

Date:08/25/04ISR Number: 4432411-2Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0342628A

Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Unknown			Balance Disorder	Zyban	PS	Glaxosmithkline	ORAL
			Diplopia	Fludex	C		
			Dizziness	Neurobion	C		

5MG per day	Gait Disturbance	Norvasc	C
30MG Three	Loss Of Consciousness	Isordil	C
times per day	Syncope		
100MG per day		Carbasalate Calcium	C
75MG Three		Dipyridamole	C
times per day			
20MG per day		Lipitor	C
200MG per day		Selokeen	C
40MG Twice		Famotidine	C
per day			

Date:08/26/04ISR Number: 4433850-6Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0425393A  
Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - per day		Dizziness					
Initial or Prolonged		Pharmaceutical Product		Paxil	C	Glaxosmithkline	ORAL
20MG Per day				Insulin	C		
Other		Complaint		Lorazepam	C		
INTRAVENOUS				Micardis	C	Glaxosmithkline	
				Cephalexin	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/26/04ISR Number: 4433857-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0523500A

Age:24 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Bupropion	PS	Glaxosmithkline	ORAL
300MG Per day		Drug Exposure During Pregnancy					

Date:08/26/04ISR Number: 4433865-8Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0320766A

Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
1TAB Twice							
per day	13 DAY			Ramipril	C		
UNKNOWN	1UNIT Twice						
per day				Aspirin	C	Glaxosmithkline	
UNKNOWN	1UNIT Per day			Diltiazem Hydrochloride	C	Glaxosmithkline	
UNKNOWN	1UNIT Per day			Atorvastatin	C		
UNKNOWN	1UNIT At						
night				Elantan La	C		
UNKNOWN	50MG In the						
morning							

Date:08/26/04ISR Number: 4433875-0Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0342716A

Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 14 DAY Initial or Prolonged	Cold Sweat Dizziness Nausea Pruritus Rash Sensation Of Foreign Body	Zyban	PS	Glaxosmithkline	ORAL
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Date:08/26/04ISR Number: 4433880-4Report Type:Expedited (15-DaCompany Report #IE-GLAXOSMITHKLINE-B0342939A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alcohol Use Convulsion Death		Zyban Alcohol	PS SS	Glaxosmithkline	ORAL

Date:08/27/04ISR Number: 4435225-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0523621A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Disability Other		Intentional Misuse Suicide Attempt		Wellbutrin	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/27/04ISR Number: 4435253-7Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0343579A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice per day	Death	Stevens-Johnson Syndrome	Zyban	PS	Glaxosmithkline	

Date:08/27/04ISR Number: 4435258-6Report Type:Expedited (15-DaCompany Report #138579

Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day 1 WK	Aggression	Apathy Personality Change	Zyban	PS	Glaxosmithkline	ORAL

Date:08/27/04ISR Number: 4435317-8Report Type:Direct Company Report #CTU 225774

Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Pruritus		Wellbutrin Sr 100 Mg Tablet	PS		

Date:08/27/04ISR Number: 4443621-2Report Type:Expedited (15-DaCompany Report #B0300505A

Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aggression Cardio-Respiratory Arrest Completed Suicide	Literature Health Professional	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL		Convulsion Depressed Level Of Consciousness Disorientation		Imipramine (Imipramine) Desipramine (Desipramine)	SS SS		

Drug Screen Positive  
Dysarthria  
Intentional Misuse  
Lethargy  
Postictal State  
Sinus Tachycardia

Clonazepam C  
Thiamine C

Date:08/30/04ISR Number: 4437037-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508671A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG See dosage text	7 DAY	Burning Sensation Dizziness Hot Flush Hyperhidrosis Hypersensitivity Nausea Pain Urticaria Vomiting	Health Professional	Zyban	PS	Glaxosmithkline	ORAL



Hospitalization -	Confusional State	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 1 WK					
Initial or Prolonged	Incontinence	Glyburide	C		
Other	Rash	Synthroid	C	Glaxosmithkline	
	Respiratory Distress	Zyloprim	C	Glaxosmithkline	
	Swollen Tongue	Ambien	C		
	Urinary Tract Infection	Vicodin	C		
		Lasix	C	Glaxosmithkline	
		Gabitril	C		

Date:08/30/04ISR Number: 4438714-XReport Type:Direct Company Report #CTU 225881  
 Age:9 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Abnormal Behaviour		Wellbutrin 150 Mg	PS		
DAILY	Aggression		Strattera	C		
	Feelings Of Worthlessness					
	Headache					
	Suicidal Ideation					

Date:08/31/04ISR Number: 4437949-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0523784A  
 Age:31 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Brain Neoplasm		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 3 WK	Grand Mal Convulsion		No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4437951-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0523797A  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Mania		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK		Suicidal Ideation	Zocor	C		

Date:08/31/04ISR Number: 4437952-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0523830A  
 Age:79 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
18 MON			Drug Ineffective Euphoric Mood Insomnia Psychomotor Hyperactivity	Clonidine	C		

Date:08/31/04ISR Number: 4437960-9Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0343398A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice			Multi-Organ Failure				
per day				Thyroxine Hormone Replacement Therapy Amlodipine	C C C	Glaxosmithkline	

Date:08/31/04ISR Number: 4437985-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498029A  
 Age:8 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Convulsion Wellbutrin PS Glaxosmithkline ORAL  
35 DAY

Date:08/31/04ISR Number: 4437986-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0503199A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	7 WK						

Date:08/31/04ISR Number: 4437987-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504891A  
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dry Mouth		Wellbutrin Sr	SS	Glaxosmithkline	
		Hepatic Enzyme Increased		Klonopin	C		
		Insomnia		Soy Supplement	C		
		Memory Impairment		Vitamins	C		
		Pharmaceutical Product					
		Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4437988-9Report Type:Periodic  
Age:29 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0506097A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	108 DAY	Grand Mal Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other		Tongue Biting					

Date:08/31/04ISR Number: 4437989-0Report Type:Periodic  
Age:63 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0506289A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	91 DAY	Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4437990-7Report Type:Periodic  
Age:49 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0506577A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	5 YR	Oedema Peripheral		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Pruritus		Ativan	C		ORAL
	.5MG Four times per day			Unknown Medication	C		ORAL
	25MG Per day						

Date:08/31/04ISR Number: 4437991-9Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0506578A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day 2 WK	Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	300MG Per day	Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged	MON	Convulsion		Insulin	C		
Other	5MG per day			Altace	C		
	150MCG per			Synthroid	C	Glaxosmithkline	
	day						
	25MG per day			Atenolol	C		
	180MG per day			Diltiazem Xr	C	Glaxosmithkline	
	60MG per day			Imdur	C		
				Asa	C	Glaxosmithkline	

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	71 DAY	Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
	50MG As	Fatigue		Ultram	C		ORAL
	required	Grand Mal Convulsion					
		Hypoaesthesia		Fosamax	C		
		Incontinence		Tranxene	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4437994-4Report Type:Periodic  
Age:29 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0508732A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	22 DAY	Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Hypersensitivity Throat Tightness		Unknown Medication	C		

Date:08/31/04ISR Number: 4437995-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509209A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3 WK	Gastrooesophageal Reflux Disease		Wellbutrin Xl Prevacid	PS C	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4437996-8Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509214A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	5 MON	Constipation Dyspepsia Paraesthesia		Wellbutrin Xl Ticar	PS C	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4437997-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509220A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 WK	Tinnitus		Wellbutrin Xl Effexor	PS C	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4437998-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0509223A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Myalgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	4	WK		Zocor	C		

Date:08/31/04ISR Number: 4437999-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0509237A  
Age:13 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Musculoskeletal Stiffness		Wellbutrin	PS	Glaxosmithkline	ORAL
Disability							
300MG Per day	2	WK					
Other							

Date:08/31/04ISR Number: 4438000-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0509240A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Euphoric Mood		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438001-XReport Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509452A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Herpes Simplex		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Nasal Oedema Oedema Mouth		Paxil	C	Glaxosmithkline	

Date:08/31/04ISR Number: 4438002-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509459A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 300MG Per day 2 WK Initial or Prolonged		Convulsion Facial Bones Fracture Fall		Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438003-3Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509464A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hyperglycaemia White Blood Cell Count Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438004-5Report Type:Periodic  
 Age:37 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0509465A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day 1 MON		Disturbance In Attention		Wellbutrin	PS	Glaxosmithkline	ORAL
				Oral Contraceptives	C		

Date:08/31/04ISR Number: 4438005-7Report Type:Periodic  
Age:49 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0509466A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2	MON		Valium	C		ORAL
10MGM At							
night							

Date:08/31/04ISR Number: 4438006-9Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509473A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3	WK		No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438007-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509548A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Delusion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day	7	WK					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438008-2Report Type:Periodic  
Age:73 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509634A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	1	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Dry Mouth							
Nausea							
Nervousness							

Date:08/31/04ISR Number: 4438009-4Report Type:Periodic  
Age:54 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0509648A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
450MG Per day	7	MON		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -							
Grand Mal Convulsion							
Initial or Prolonged				Celexa	C		
				Lamictal	C	Glaxosmithkline	
				Zyprexa	C		
				Buspar	C		

Date:08/31/04ISR Number: 4438010-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509649A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	10	DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
Abdominal Pain							
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438011-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509654A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Headache							
Somnolence							

Date:08/31/04ISR Number: 4438012-4Report Type:Periodic  
Age:41 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509657A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Blood Triglycerides	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	11	MON	Increased	Zoloft	C		
				Flintstone Vitamins	C		

Date:08/31/04ISR Number: 4438013-6Report Type:Periodic  
Age:54 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509812A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Tinnitus	Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	6	WK		Flomax	C		

Date:08/31/04ISR Number: 4438014-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509814A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Blood Glucose Increased	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438015-XReport Type:Periodic  
 Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509818A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1	WK	Drug Ineffective	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Stool Analysis Abnormal	No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438016-1Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0509820A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Erythema Nodosum	Wellbutrin Xl	PS	Glaxosmithkline	

Date:08/31/04ISR Number: 4438017-3Report Type:Periodic  
 Age:13 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509828A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3	WK	Tremor	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Zoloft	C		

Date:08/31/04ISR Number: 4438018-5Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0509850A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day			Visual Disturbance	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Wellbutrin Sr	C	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438019-7Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509855A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression		Wellbutrin	PS	Glaxosmithkline	ORAL
2	MON	Anger Suicidal Ideation		Paxil	C	Glaxosmithkline	

Date:08/31/04ISR Number: 4438020-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0509918A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG	Unknown	Pharmaceutical Product Complaint					

Date:08/31/04ISR Number: 4438021-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0509919A  
 Age: Gender:I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Spasms		Wellbutrin Xl	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438022-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509925A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cough		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438023-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509936A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression		Wellbutrin	PS	Glaxosmithkline	ORAL
2 MON		Anger Suicidal Ideation		Antidepressant (Unspecified)	C		

Date:08/31/04ISR Number: 4438024-0Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509991A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decreased Appetite		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 4 DAY		Energy Increased		Trazodone Birth Control Pills	SS C		

Date:08/31/04ISR Number: 4438025-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0509992A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438026-4Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0509993A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 WK		Panic Reaction		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Zoloft	C		

Date:08/31/04ISR Number: 4438027-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0509996A  
 Age:39 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 MON	Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Synthroid	C	Glaxosmithkline	
				Effexor	C		

Date:08/31/04ISR Number: 4438028-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510006A  
 Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
450MG Unknown		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Gastrooesophageal Reflux Disease		No Concurrent Medication	C		
		Headache					
		Nausea					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438029-XReport Type:Periodic  
Age:46 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0510020A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
55 DAY				Paxil Cr	SS	Glaxosmithkline	ORAL
75MG Per day				Lamictal	C	Glaxosmithkline	ORAL
200MG per day	76 DAY			Trileptal	C		ORAL
300MG Twice							
per day				Vistaril	C		ORAL

Date:08/31/04ISR Number: 4438030-6Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510021A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3 WK	Migraine		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Sensation Of Pressure		Clarinet	C		
		Vision Blurred		Fosamax	C		

Date:08/31/04ISR Number: 4438031-8Report Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510024A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3 MON	Skin Discolouration		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Prevacid	C		

Date:08/31/04ISR Number: 4438032-XReport Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510032A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Per day 1 DAY      Dissociation      Wellbutrin Xl      PS      Glaxosmithkline      ORAL  
Tequin      C

Date:08/31/04ISR Number: 4438033-1Report Type:Periodic      Company Report #US-GLAXOSMITHKLINE-A0510209A  
Age:      Gender:Female      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Contusion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 1	MON			No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438034-3Report Type:Periodic      Company Report #US-GLAXOSMITHKLINE-A0510212A  
Age:      Gender:Male      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 3	MON			No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438035-5Report Type:Periodic      Company Report #US-GLAXOSMITHKLINE-A0510214A  
Age:83 YR      Gender:Male      I/FU:I

Outcome      PT  
Balance Disorder  
Constipation  
Dizziness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Fatigue

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Lexapro	C		

Date:08/31/04ISR Number: 4438036-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510235A  
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 MON	Pruritus		Naproxen	C		
		Rash Erythematous		Allegra	C		

Date:08/31/04ISR Number: 4438037-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510240A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Anxiety		Effexor	SS		
		Drug Withdrawal Syndrome		Alprazolam	C		
		Headache					
		Mood Altered					

Date:08/31/04ISR Number: 4438038-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510336A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR	Therapeutic Response					
		Decreased					

Date:08/31/04ISR Number: 4438039-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510340A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sexual Dysfunction		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438040-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510399A  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 MON	Vomiting		No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438041-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510408A  
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day	5 DAY			Trazodone	C		
				Trazodone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438042-2Report Type:Periodic  
Age:16 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0510413A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day	1 WK		Wellbutrin	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438043-4Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510419A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG Per day	2 WK		Wellbutrin	PS	Glaxosmithkline	ORAL
				Zoloft	C		
				Ambien	C		
				Xanax	C		
				Seroquel	C		

Date:08/31/04ISR Number: 4438044-6Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510424A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG Per day	5 MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Zyrtec	C	Glaxosmithkline	

Date:08/31/04ISR Number: 4438045-8Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510434A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Lipitor	C		

225MG Per day	Nausea		Effexor	C		
			Wellbutrin Sr	C	Glaxosmithkline	ORAL
YR						
Date:08/31/04ISR Number: 4438046-XReport Type:Periodic			Company Report #US-GLAXOSMITHKLINE-A0510503A			
Age:	Gender:Female	I/FU:F				
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	Cough					
	Insomnia					
	Urinary Incontinence					
Date:08/31/04ISR Number: 4438047-1Report Type:Periodic			Company Report #US-GLAXOSMITHKLINE-A0510513A			
Age:	Gender:	I/FU:I				
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	Tendonitis					
300MG Per day						
Date:08/31/04ISR Number: 4438048-3Report Type:Periodic			Company Report #US-GLAXOSMITHKLINE-A0510517A			
Age:	Gender:Female	I/FU:I				
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	Alopecia					
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Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438049-5Report Type:Periodic  
Age:53 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510523A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	4 DAY	Dizziness	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Plavix	C		
				Lipitor	C		
				Aspirin	C	Glaxosmithkline	
				Platil	C		
				Bextra	C		
				Terazosin	C		
				Aceon	C		
				Indapamide	C		
				Acetaminophen	C	Glaxosmithkline	
				Propoxyphene	C		

Date:08/31/04ISR Number: 4438050-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510527A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	4 WK	Dizziness	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438051-3Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0510530A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Vomiting	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438052-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510531A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

450MG Per day Drug Ineffective Wellbutrin Xl PS Glaxosmithkline ORAL

Date:08/31/04ISR Number: 4438053-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510534A  
 Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day 1 WK	Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Localised Oedema		Coumadin	C	Glaxosmithkline	
				Lasix	C	Glaxosmithkline	
				Beta Blocker	C		
				Diovan	C		

Date:08/31/04ISR Number: 4438054-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510537A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438055-0Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510539A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Heart Rate Increased		Strattera	C		
		Irritability		Trileptal	C		
		Medication Error					
		Stomach Discomfort					

Date:08/31/04ISR Number: 4438056-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0510540A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438057-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510686A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	
Other				Ambien	C		

Date:08/31/04ISR Number: 4438058-6Report Type:Periodic  
Age:12 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510729A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	Haemorrhage Subcutaneous		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Advair	C	Glaxosmithkline	
				Flonase	C	Glaxosmithkline	
				Zyrtec	C	Glaxosmithkline	

Date:08/31/04ISR Number: 4438059-8Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510739A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY						

Date:08/31/04ISR Number: 4438060-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510746A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mania		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Nicotine Patches	C	Glaxosmithkline	

Date:08/31/04ISR Number: 4438061-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510752A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR			Wellbutrin Sr	SS	Glaxosmithkline	ORAL
				Ambien	C		
				Synthroid	C	Glaxosmithkline	
				Clonazepam	C		

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Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438062-8Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510757A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day				Lexapro	C		

Date:08/31/04ISR Number: 4438063-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510758A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438064-1Report Type:Periodic  
Age:55 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510761A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day				Zoloft	C		

Date:08/31/04ISR Number: 4438065-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510763A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:08/31/04ISR Number: 4438066-5Report Type:Periodic  
Age:41 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0510764A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day	170 DAY	Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged 2.5MG Per day		Grand Mal Convulsion		Cenestin	SS		
		Incontinence		Singulair	C		
				Levo-Thyroxin	C	Glaxosmithkline	
.025MG Per day				Hydrochlorothiazide	C		
12.5MG Per day							

Date:08/31/04ISR Number: 4438068-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510860A  
Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day	15 DAY	Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Initial or Prolonged Other		Facial Bones Fracture Fall Tremor					

Date:08/31/04ISR Number: 4438069-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510881A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	7 MON	Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438070-7Report Type:Periodic  
Age:51 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0510904A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	8 DAY	Bacterial Infection	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Ativan	C		
				Dicyclomine	C		

Date:08/31/04ISR Number: 4438071-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510916A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Insomnia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438072-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510917A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
Other							

Date:08/31/04ISR Number: 4438073-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510929A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	1 MON	Rash	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438074-4Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510936A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Per day 1 DAY	Insomnia	Wellbutrin	PS	Glaxosmithkline	ORAL
		Celexa	C		

Date:08/31/04ISR Number: 4438075-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510937A  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 1 WK				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438076-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510940A  
 Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 11 DAY		Crying		Zoloft	C		
		Dizziness		Cyclessa	C		
		Dry Mouth		Singulair	C		
		Nausea					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438077-XReport Type:Periodic  
 Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510947A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK			Lexapro	C		
		Disorientation					
		Insomnia					

Date:08/31/04ISR Number: 4438078-1Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510948A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Alcohol	SS		
		Convulsion		No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438079-3Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511013A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Abdominal Discomfort					
		Crying					
		Mood Altered					

Date:08/31/04ISR Number: 4438080-XReport Type:Periodic  
 Age:16 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511025A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
1 YR		Alopecia					

Date:08/31/04ISR Number: 4438081-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511062A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
5	DAY	Feeling Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Irritability Panic Reaction		Zoloft	C		

Date:08/31/04ISR Number: 4438082-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0511066A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2	WK	Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Fatigue		Zoloft	C		

Date:08/31/04ISR Number: 4438083-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0511086A  
 Age:86 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	23 DAY	Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hypertension		Baby Aspirin	C	Glaxosmithkline	
		Insomnia		Ditropan Xl	C		
		Nervousness		Ambien	C		
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438084-7Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511100A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	14 DAY	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Crying		Xanax	C		
		Dizziness					
		Headache					

Date:08/31/04ISR Number: 4438085-9Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511106A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Zoloft	C		
				Multiple Medications	C		

Date:08/31/04ISR Number: 4438086-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511113A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 WK	Sinus Congestion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Vision Blurred		No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438087-2Report Type:Periodic  
Age:51 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511120A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	1 YR	Depressed Mood		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Erectile Dysfunction		Trazodone	C		
		Semen Volume Decreased		Viracept	C		
				Combivir	C	Glaxosmithkline	

Pravachol C  
Zetia C

Date:08/31/04ISR Number: 4438088-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511233A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
1 WK		Urticaria					

Date:08/31/04ISR Number: 4438089-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511266A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438090-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511303A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - 10 DAY Initial or Prolonged				Cipro Alcohol	C C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438092-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511403A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying Emotional Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438093-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511410A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438094-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511463A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	29 DAY			Lorazepam	C		

Date:08/31/04ISR Number: 4438095-1Report Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511471A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK	Nervousness		Norco	C		
				Soma	C		

Date:08/31/04ISR Number: 4438096-3Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0511473A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

300MG Per day

Fatigue

Wellbutrin XL PS Glaxosmithkline ORAL

Date:08/31/04ISR Number: 4438097-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0511588A  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK						

Date:08/31/04ISR Number: 4438098-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0511590A  
 Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Withdrawal Syndrome		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	30 WK	Headache		Ambien	C		
		Nausea		Klonopin	C		

Date:08/31/04ISR Number: 4438099-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0511598A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Feeling Jittery		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	7 DAY	Insomnia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438100-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511599A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Haemoglobin Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438101-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511606A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	6 WK	Insomnia Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438102-6Report Type:Periodic  
Age:29 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511611A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1 DAY	Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438103-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511617A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1 WK	Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Adderall	C		
				Neurontin	C		

Date:08/31/04ISR Number: 4438104-XReport Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511620A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3 WK	Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Neck Pain Radiotherapy		No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438105-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0511626A  
 Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Vomiting		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438106-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0511720A  
 Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Myoclonus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438107-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0511741A  
 Age: Gender: I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression Irritability		Wellbutrin Xl	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438108-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511763A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin Xl No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438109-9Report Type:Periodic  
 Age:22 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511876A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	450MG Per day	Cold Sweat  Dizziness Hyperhidrosis Nausea Vision Blurred Vomiting		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438110-5Report Type:Periodic  
 Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511898A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mouth Ulceration		Wellbutrin Xl Prozac Lipitor Norvasc Vitamins Water Pill Wellbutrin Sr	PS C C C C C C	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438111-7Report Type:Periodic  
 Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511906A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

300MG Per day Drug Ineffective Wellbutrin Xl PS Glaxosmithkline ORAL

No Concurrent Medication C

Date:08/31/04ISR Number: 4438112-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0511907A  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice		Oedema Peripheral					
per day	2 YR			Bupropion Hydrochloride	SS	Glaxosmithkline	

Date:08/31/04ISR Number: 4438113-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0511912A  
 Age:78 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 DAY			Amitriptyline	C		
				Perphenazine	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438114-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511917A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Muscle Twitching		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438115-4Report Type:Periodic  
Age:76 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512053A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3 WK	Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Capoten	C	Glaxosmithkline	
				Lipitor	C		
				Hctz	C		

Date:08/31/04ISR Number: 4438116-6Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512054A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	1 MON	Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Prednisone	C		
				Urso	C		
				Salagen	C		
				Cyclobenzaprine	C		
				Femhrt	C		

Date:08/31/04ISR Number: 4438117-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512060A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	17 DAY	Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Ineffective					
		Insomnia					

Date:08/31/04ISR Number: 4438118-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512062A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438119-1Report Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512082A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	6 DAY	Headache		Lantus	C		
		Nausea		Humalog Insulin	C		
				Lanoxin	C	Glaxosmithkline	
				Potassium	C		
				Lipitor	C		
				Bumex	C		
				Multi-Vitamins	C		
				Aspirin	C	Glaxosmithkline	
				Zaroxolyn	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438120-8Report Type:Periodic  
Age:18 MON Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512085A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438121-XReport Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0512087A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK	Dizziness Headache Myalgia Neck Pain Tension		No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438122-1Report Type:Periodic  
Age:41 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512228A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	8 DAY	Insomnia		Pain Medication	SS		

Date:08/31/04ISR Number: 4438123-3Report Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512234A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
5 WK				Lexapro	C		

Date:08/31/04ISR Number: 4438124-5Report Type:Periodic  
Age:14 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0512239A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lethargy		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438125-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512249A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK			Insulin	C		
				Blood Pressure Medication	C		
				Thyroid Medication	C		
				Milk Of Magnesium	C	Glaxosmithkline	
				Stool Softener	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438126-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512340A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Muscle Twitching		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438127-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512341A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438128-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512343A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oropharyngeal Swelling		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438129-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512388A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	4 MON	Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438130-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512396A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1 MON	Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
				Lexapro	C		

Date:08/31/04ISR Number: 4438131-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512400A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
0	DAY						

Date:08/31/04ISR Number: 4438132-4Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512403A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4 MON			Ativan	C		
				Vivelle	C		
				Singulair	C		
				Wellbutrin Sr	C	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438133-6Report Type:Periodic  
Age:25 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512404A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vomiting		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK			No Concurrent			



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Medication C

Date:08/31/04ISR Number: 4438134-8Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512414A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3	WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Candidiasis					
		Dry Mouth		No Concurrent Medication	C		
		Oral Discomfort					

Date:08/31/04ISR Number: 4438135-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512418A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	4	DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dizziness					
		Hypertension		Dhea	C		
		Palpitations		Vitamins	C		
				Antioxidant	C		
				Calcium	C		
				Garlic Capsules	C		
				Ginger Root	C		

Date:08/31/04ISR Number: 4438136-1Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0512451A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Insomnia					

Date:08/31/04ISR Number: 4438137-3Report Type:Periodic  
Age:34 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0512534A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

300MG Per day 14 DAY Rash Generalised

Wellbutrin Xl PS Glaxosmithkline ORAL

INTRAMUSCULAR

Depo Medrol C

Zyrtec C Glaxosmithkline ORAL

Date:08/31/04ISR Number: 4438138-5Report Type:Periodic  
Age:22 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0512603A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Consciousness		Wellbutrin	PS	Glaxosmithkline	ORAL
248 DAY		Muscle Rigidity Petit Mal Epilepsy		Oral Contraceptives	C		

Date:08/31/04ISR Number: 4438139-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512645A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Nightmare Thinking Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438140-3Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512868A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1	MON		No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438141-5Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512874A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1	MON		Lexapro	C		
				Fosamax	C		
				Nasonex	C		
				Astelina	C		
				Advair	C	Glaxosmithkline	
				Combivent	C		
				Klonopin	C		
				Protonix	C		
				Hydroxyzine	C		
				Dapsone	C		
				Wellbutrin Sr	C	Glaxosmithkline	

Date:08/31/04ISR Number: 4438142-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512878A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice				Zocor	C		
per day				Aspirin	C	Glaxosmithkline	

Date:08/31/04ISR Number: 4438143-9Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512893A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 DAY	Feeling Abnormal		Advil	C	Glaxosmithkline	
		Mood Swings		Klonopin	C		
		Negative Thoughts					

Date:08/31/04ISR Number: 4438144-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512960A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Xanax	SS		ORAL

Date:08/31/04ISR Number: 4438145-2Report Type:Periodic  
Age:37 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0512964A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Generalised		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	14 DAY			Depo Medrol	C		
80MG Per day				Zyrtec	C	Glaxosmithkline	ORAL
10MG Per day							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438146-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513012A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 WK		Oedema Peripheral		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Pain		Diovan	C		
				Procardia	C	Glaxosmithkline	
				Nexium	C		
				Atenolol	C		
				Flonase	C	Glaxosmithkline	

Date:08/31/04ISR Number: 4438148-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513023A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Insomnia		Vitamin E	C		
		Nausea		Vitamin B	C		
				Vitamin C	C	Glaxosmithkline	

Date:08/31/04ISR Number: 4438149-XReport Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513026A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	18 DAY	Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Synthroid	C	Glaxosmithkline	
				Birth Control	C		

Date:08/31/04ISR Number: 4438150-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513057A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1 DAY	Flushing		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Lopid	C	
Synthroid	C	Glaxosmithkline
Premarin	C	
Zyrtec	C	Glaxosmithkline
Lipitor	C	
Nexium	C	
Accolate	C	

Date:08/31/04ISR Number: 4438151-8Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513226A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Therapeutic Response Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438152-XReport Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513267A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Screen False Positive		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438153-1Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513270A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	14 DAY	Tremor		Xanax	C		
				Percocet	C		

Date:08/31/04ISR Number: 4438154-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513285A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:08/31/04ISR Number: 4438155-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513289A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nervousness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Lescol	C		
				Foradil	C		
				Doxazosin	C		
				Albuterol	C	Glaxosmithkline	
				Zantac	C	Glaxosmithkline	
				Ativan	C		
				Aspirin	C	Glaxosmithkline	
				Nasacort	C		

Date:08/31/04ISR Number: 4438156-7Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513290A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Myalgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	17 DAY						

Date:08/31/04ISR Number: 4438157-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513295A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	5	WK		Sotalol	C		
			Dizziness				
			Nausea				
			Stomach Discomfort				

Date:08/31/04ISR Number: 4438158-0Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0513404A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Tinnitus				

Date:08/31/04ISR Number: 4438159-2Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0513437A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3	MON		Allegra	C		
			Dysphagia				
			Swollen Tongue				



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Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438160-9Report Type:Periodic  
Age:14 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513438A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	2	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438161-0Report Type:Periodic  
Age:59 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513443A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	7	DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438162-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513466A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day				Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438163-4Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513478A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3	WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Zoloft	C		

Date:08/31/04ISR Number: 4438164-6Report Type:Periodic  
Age:43 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0513480A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	7 MON	Asthenia		No Concurrent Medication	C		
		Depression					
		Disturbance In Attention					
		Fatigue					

Date:08/31/04ISR Number: 4438165-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0513481A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Menstruation Irregular		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK			Papaverine	C		

Date:08/31/04ISR Number: 4438166-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0513483A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Myocardial Infarction		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438167-1Report Type:Periodic  
Age:27 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513486A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 DAY	Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Heart Rate Increased		Atenolol	C		
				Xenadrine	C		

Date:08/31/04ISR Number: 4438168-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513490A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1 MON	Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Lexapro	C		

Date:08/31/04ISR Number: 4438169-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513618A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Nervous System Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438170-1Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513644A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	5 DAY	Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Ineffective		Klonopin	C		
		Headache					
		Nausea					

Date:08/31/04ISR Number: 4438171-3Report Type:Periodic  
Age:10 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513648A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:08/31/04ISR Number: 4438172-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0513653A  
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Conjunctivitis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 1 YR		Constipation Ill-Defined Disorder		No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438173-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0513656A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Stool Analysis Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438174-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513658A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3 DAY			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Vision Blurred					
		Weight Increased					

Date:08/31/04ISR Number: 4438175-0Report Type:Periodic  
Age:31 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513775A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice				Wellbutrin	PS	Glaxosmithkline	ORAL
		Anger					
per day	2 MON			No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438176-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513779A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	6 MON			Wellbutrin	PS	Glaxosmithkline	ORAL
		Arthralgia					
		Myalgia					

Date:08/31/04ISR Number: 4438177-4Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0513783A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 MON			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Tinnitus					
25MG Per day				Hctz	C		
80MG Per day				Zocor	C		

Vitamins C  
Aciphex C

20MG Twice

per day

Date:08/31/04ISR Number: 4438178-6Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513821A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Cough		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 11 DAY			Spirolactone	C		
			Estradiol	C		
			Crestor	C		

Date:08/31/04ISR Number: 4438179-8Report Type:Periodic  
Age:10 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513835A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Irritability		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day			Strattera	C		
	Tremor					

Date:08/31/04ISR Number: 4438180-4Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513957A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 5 WK			Celebrex	C		

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Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438181-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513977A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 YR		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438182-8Report Type:Periodic  
 Age:54 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0513978A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
UNKNOWN		Drug Ineffective Vision Blurred		No Concurrent Medication	C		
10MG In the morning				Clonazepam	C		
				Zestril	C		ORAL

Date:08/31/04ISR Number: 4438183-XReport Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0513986A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Psychotropic Drugs	C		

Date:08/31/04ISR Number: 4438184-1Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514094A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown		Sexual Dysfunction		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438185-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514111A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Pharmaceutical Product Complaint Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438186-5Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514140A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK	Diarrhoea		Lorazepam	C		

Date:08/31/04ISR Number: 4438187-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514149A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438188-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514151A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG Per day	Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Effexor	C		
				Prednisone	C		

Date:08/31/04ISR Number: 4438189-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514278A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Menstruation Irregular		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438190-7Report Type:Periodic  
Age:74 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514312A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day 1 MON	Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Constipation		Tigan	C	Glaxosmithkline	
		Nausea		Compazine	C	Glaxosmithkline	
		Weight Decreased		Strattera	C		
				Vitamins	C		

Date:08/31/04ISR Number: 4438191-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514322A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	
				Gabitril	C		

Date:08/31/04ISR Number: 4438192-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514323A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438193-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514328A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Asthenia Musculoskeletal Stiffness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438194-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514330A  
 Age:54 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	14 DAY	Gastric Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Nausea		Levoxyl	C	Glaxosmithkline	
		Paraesthesia		Tenormin	C		
				Lorazepam	C		
				Amitriptyline	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438195-6Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514345A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 WK			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dizziness					
		Drug Ineffective		Zoloft	C		
		Feeling Abnormal					
		Nausea					

Date:08/31/04ISR Number: 4438196-8Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514507A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
450MG Per day				Wellbutrin	PS	Glaxosmithkline	ORAL
		Anorexia					
		Confusional State		No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438197-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514510A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
				Wellbutrin	PS	Glaxosmithkline	ORAL
		Salivary Hypersecretion					

Date:08/31/04ISR Number: 4438198-1Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0514518A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -	7 DAY			Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged				Restoril	C		
Other		Drug Ineffective					
		Suicidal Ideation					

Date:08/31/04ISR Number: 4438199-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514615A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amenorrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
6	MON						

Date:08/31/04ISR Number: 4438200-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514626A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other				Lexapro	C		
				Unspecified Medications	C		

Date:08/31/04ISR Number: 4438201-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514670A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG	Per day						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438202-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514674A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Retching		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:08/31/04ISR Number: 4438203-2Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514679A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 1 WK							
Heart Rate Increased							
Hyperhidrosis							
Migraine							
Nausea							

Date:08/31/04ISR Number: 4438204-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514853A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Repetitive Speech		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 30 DAY							
				Alcohol	C		

Date:08/31/04ISR Number: 4438205-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514869A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Jittery		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438206-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514870A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	7 MON	Hair Texture Abnormal		Multiple Medications	C		

Date:08/31/04ISR Number: 4438207-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514874A  
Age: Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia		Wellbutrin Xl	PS	Glaxosmithkline	

Date:08/31/04ISR Number: 4438208-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514878A  
Age:39 YR Gender:Female I/FU:I

Outcome	PT
	Anxiety
	Blood Pressure Decreased
	Disorientation
	Dizziness
	Heart Rate Increased
	Migraine
	Nausea
	Nervousness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Tinnitus Tremor					
Dose	Duration		Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 MON			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Estrogen	C		
				Aleve Cold + Sinus	C		

Date:08/31/04ISR Number: 4438209-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514885A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	6 WK	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Educational Problem		No Concurrent			
		Irritability		Medication	C		

Date:08/31/04ISR Number: 4438210-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514895A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Headache		Ibuprofen	C	Glaxosmithkline	
				Actos	C		
				Zaroxolyn	C		
				Aldactone	C		
				Butalbital/Apap/Caff			
				eine	C		
				Coumadin	C	Glaxosmithkline	
				Hydrocodone	C		
				Docusate	C		
				Glucophage	C		
				Prevacid	C		
				Diovan	C		
				Lanoxin	C	Glaxosmithkline	
				Betapace	C	Glaxosmithkline	
				Alprazolam	C		
				Albuterol Inhaler	C	Glaxosmithkline	
				Advair	C	Glaxosmithkline	

Phazyme	C	Glaxosmithkline
Afrin	C	Glaxosmithkline
Ocean Nasal	C	Glaxosmithkline
Flonase	C	Glaxosmithkline
Lasix	C	Glaxosmithkline
Potassium	C	

Date:08/31/04ISR Number: 4438211-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514908A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3	WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438212-3Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0515014A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Skin Test Negative		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438213-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515025A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438214-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515026A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438215-9Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515134A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	10 DAY	Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Impatience		Buspar	C		

Date:08/31/04ISR Number: 4438216-0Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515154A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Twice per day	6 WK	Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
		Psychomotor Hyperactivity					

Respiratory Sighs

Nicotine Transdermal

Patch

C

Glaxosmithkline

Date:08/31/04ISR Number: 4438217-2Report Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0515163A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	300MG Per day 10	MON	Amnesia	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged			Grand Mal Convulsion				
Other							

Date:08/31/04ISR Number: 4438218-4Report Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0515225A

Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	300MG Per day 8	MON	Menopause	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged				Prozac	C		
Other				Oral Contraceptives	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438219-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515309A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438220-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515362A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscular Weakness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:08/31/04ISR Number: 4438221-4Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515367A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 2 WK							
				Diovan	C		
				Metformin	C		
				Blood Pressure Medication	C		ORAL

Date:08/31/04ISR Number: 4438222-6Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515383A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ageusia	Consumer	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150TAB Per day							
	1 MON	Dysphagia					
		Nausea Vomiting		Pravachol	C		

Date:08/31/04ISR Number: 4438223-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515385A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash Erythematous		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Rash Pruritic		No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438224-XReport Type:Periodic  
Age:43 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0515395A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day	3 MON	Depressed Level Of Consciousness		Lexapro	SS		ORAL
10MG Per day	442 DAY	Eye Rolling					
		Grand Mal Convulsion					
		Postictal State					
		Tongue Biting					
		Tonic Clonic Movements					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438225-1Report Type:Periodic  
Age:61 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0515536A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3 DAY	Agitation Drug Ineffective Headache Nausea Panic Attack		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438226-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515573A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1 MON	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438227-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515610A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 WK	Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438228-7Report Type:Periodic  
Age:70 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515730A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG per day	8 DAY	Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Glucophage	C		

Date:08/31/04ISR Number: 4438229-9Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515731A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 HR	Nausea Pain		No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438230-5Report Type:Periodic  
Age:25 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0515753A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menorrhagia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	18 DAY	Menstrual Disorder		Ortho Tricyclen	C		
1TAB Per day	5 YR			Nortriptyline	C		
50MG At night							

Date:08/31/04ISR Number: 4438231-7Report Type:Periodic  
Age:80 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0515873A

Outcome	PT
	Dermatitis Exfoliative
	Dry Skin
	Erythema
	Oedema
	Pain

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Agitation

Wellbutrin Xl

PS

Glaxosmithkline

ORAL

Date:08/31/04ISR Number: 4438235-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515911A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		Ocular Hyperaemia		Wellbutrin Xl	PS	Glaxosmithkline	

Date:08/31/04ISR Number: 4438236-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515916A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3 WK	Disturbance In Attention Irritability		Wellbutrin Xl No Concurrent Medication	PS C	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438237-8Report Type:Periodic  
 Age:49 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0516121A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	1	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Hydrochlorothiazide	C		
				Atenolol	C		
				Albuterol	C	Glaxosmithkline	
				Advair	C	Glaxosmithkline	

Date:08/31/04ISR Number: 4438238-XReport Type:Periodic  
 Age:62 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516132A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3	WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Zocor	C		
				Prevacid	C		
				Plavix	C		
				Klonopin	C		
				Antihypertensive	C		

Date:08/31/04ISR Number: 4438239-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516141A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438240-8Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0516143A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Other Angioneurotic Oedema Swollen Tongue Wellbutrin Xl PS Glaxosmithkline ORAL

Date:08/31/04ISR Number: 4438241-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0516243A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3	MON	Breast Tenderness	Pravachol	C		

Date:08/31/04ISR Number: 4438242-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0516248A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cognitive Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	6	MON		Methotrexate	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438243-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516251A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	Amnesia Cognitive Disorder Dysphasia Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438244-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516252A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
				Depakote Er	C		

Date:08/31/04ISR Number: 4438245-7Report Type:Periodic  
Age:45 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0516258A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Hospitalization - 300MG In the Initial or Prolonged morning Other	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
				Dilantin	C		

Date:08/31/04ISR Number: 4438246-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0516384A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	2 WK	Menorrhagia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438248-2Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0516423A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG	Per day						

Date:08/31/04ISR Number: 4438249-4Report Type:Periodic  
Age:70 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516428A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3	DAY	Confusional State		Synthroid	C	Glaxosmithkline	
				Zyrtec	C	Glaxosmithkline	
				Zestril	C		

Date:08/31/04ISR Number: 4438250-0Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516436A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG	Twice						
per day	1 WK			Lexapro	C		
				Nexium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438251-2Report Type:Periodic  
 Age:59 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516440A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 MON	Aggression Agitation Anxiety Energy Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Glucophage	C		
				Glucovance	C		
				Estrogen Replacement	C		
				Xanax	C		
				Proton Pump Inhibitor	C		

Date:08/31/04ISR Number: 4438252-4Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516452A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Screen False Positive		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438253-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516460A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	12 MON	Weight Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Blood Pressure Medication	C		

Date:08/31/04ISR Number: 4438254-8Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0516549A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Menorrhagia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438255-XReport Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0516553A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Unknown	1 YR	Dyspepsia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Irritable Bowel Syndrome					

Date:08/31/04ISR Number: 4438256-1Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516599A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3 WK	Dry Mouth		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dyspepsia					

Date:08/31/04ISR Number: 4438257-3Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0516601A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dysgeusia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438258-5Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516611A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	300MG Per day	3 MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Oral Contraceptives	C		

Date:08/31/04ISR Number: 4438259-7Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516621A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	300MG Per day	10 WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Ambien	C		
				Xanax	C		
				Seroquel	C		
				Combivent	C		
				Qvar	C	Glaxosmithkline	
				Xopenex	C		

Date:08/31/04ISR Number: 4438260-3Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516630A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	300MG Per day	3 MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Amaryl	C		
				Synthroid	C	Glaxosmithkline	
				Avapro	C		

Date:08/31/04ISR Number: 4438261-5Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516631A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Per day 3 WK Feeling Jittery Wellbutrin Xl PS Glaxosmithkline ORAL  
Zoloft C

Date:08/31/04ISR Number: 4438262-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0516754A  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
				Lithium	C	Glaxosmithkline	
				Klonopin	C		

Date:08/31/04ISR Number: 4438263-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0516760A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Muscular Weakness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Paraesthesia					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438264-0Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0516766A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	4 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Aggression					
		Depression					
		Nausea					
		Vomiting					

Date:08/31/04ISR Number: 4438265-2Report Type:Periodic  
 Age: Gender:I I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516777A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Alopecia					

Date:08/31/04ISR Number: 4438266-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516778A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	8 WK		Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Ineffective					

Date:08/31/04ISR Number: 4438267-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516911A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Insomnia					
		Medication Error		Trazodone	C		
				Prozac	C		

Date:08/31/04ISR Number: 4438268-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516990A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Coordination Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438269-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0517063A  
 Age:71 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
450MG Twice per day	3 YR	Drug Ineffective Sexual Dysfunction Sleep Disorder		Wellbutrin Xl  Lamictal Provigil Viagra	PS  C C C	Glaxosmithkline  Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438270-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0517090A  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nausea Vomiting		Wellbutrin Xl No Concurrent Medication	PS C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438271-8Report Type:Periodic  
 Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLIN-A0517091A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Dry Mouth		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dysphagia		Neurontin	C		
		Migraine		Skelaxin	C		
		Throat Tightness		Elavil	C	Glaxosmithkline	
				Xanax	C		
				Toprol Xl	C		
				Vicodin	C		
				Advil	C	Glaxosmithkline	
				Tylenol Es	C	Glaxosmithkline	
				Ambien	C		
				Maxalt	C		
				Proventil	C	Glaxosmithkline	

Date:08/31/04ISR Number: 4438272-XReport Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLIN-A0517246A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day 1	MON	Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Drug Ineffective					

Date:08/31/04ISR Number: 4438273-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLIN-A0517248A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day 6	WK	Hyperhidrosis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Effexor	SS		

Date:08/31/04ISR Number: 4438274-3Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLIN-A0517253A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Accident Hyperhidrosis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438275-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0517255A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
4 MON		Memory Impairment		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Neurontin	C		

Date:08/31/04ISR Number: 4438276-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0517259A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Sunburn		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438277-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517263A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Claustrophobia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Insomnia		Zelnorm	C		
		Irritability					
		Mood Altered					

Date:08/31/04ISR Number: 4438278-0Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517273A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Crying		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Allegra	C		

Date:08/31/04ISR Number: 4438279-2Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517389A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
3 MON		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438280-9Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517405A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Unknown 7 DAY		Feeling Hot		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hot Flush					

Date:08/31/04ISR Number: 4438281-0Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517421A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 WK	Tension					

Date:08/31/04ISR Number: 4438282-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0517432A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:08/31/04ISR Number: 4438283-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0517439A  
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Arthritis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	5 WK	Joint Swelling		Yasmin	C		
		Oedema Peripheral					
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438284-6Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0517441A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dysphonia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2	MON					

Date:08/31/04ISR Number: 4438285-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517445A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Face Oedema		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6	MON					
		Rash Pustular		Trileptal	C		
				Nexium	C		

Date:08/31/04ISR Number: 4438286-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517667A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	
UNKNOWN							
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438287-1Report Type:Periodic  
Age:72 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517669A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Somnolence	Consumer	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	10	MON					
				Aricept	C		
				Nexium	C		
				Lasix	C	Glaxosmithkline	
				Albuterol	C	Glaxosmithkline	
				Zyprexa	C		

Date:08/31/04ISR Number: 4438288-3Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517675A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Irritability		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	10 DAY			Hydrochlorothiazide	C		
				Zyrtec	C	Glaxosmithkline	
				Lipitor	C		
				Plavix	C		
				Nexium	C		

Date:08/31/04ISR Number: 4438289-5Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517681A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Local Swelling	Consumer	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	7 WK			Dhea	SS		
				Fosamax	C		
				Soma	C		
				Ultracet	C		
				Combi-Patch	C		
				Ginseng	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438290-1Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517683A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	24 DAY	Abdominal Distension Faeces Discoloured Flatulence Stomach Discomfort		Wellbutrin Xl Tegretol	PS C	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438291-3Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517689A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 MON	Tinnitus		Wellbutrin Xl Synthroid	PS C	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438292-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517692A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Prostatic Specific Antigen Increased		Wellbutrin Xl No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438293-7Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517693A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3 WK	Headache Thinking Abnormal Weight Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438294-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0517698A  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Alcohol Use		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	90 DAY	Personality Change		Prozac	C		
				Alcohol	C		

Date:08/31/04ISR Number: 4438295-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0517849A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 MON			No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438296-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0517857A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Leukopenia		Wellbutrin	PS	Glaxosmithkline	ORAL
6 MON							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438298-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517878A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Discomfort		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438299-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518021A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438300-1Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0518022A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abnormal Dreams Nightmare		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438301-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518024A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Unknown		Hiccups		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438302-5Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518073A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3 WK	Agitation	Consumer	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Anger		Humalog	C		
		Anorexia		Evista	C		

Anxiety  
Crying  
Dizziness  
Dysgeusia  
Emotional Disorder  
Fungal Infection  
Hyperhidrosis  
Insomnia  
Myalgia  
Nausea  
Tremor  
Weight Decreased

Vitamins

C

Date:08/31/04ISR Number: 4438303-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518082A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Irritability		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	450MG Per day			No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438304-9Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0518095A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mouth Ulceration		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438305-0Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518099A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cough		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 MON			Insulin	C		

Date:08/31/04ISR Number: 4438306-2Report Type:Periodic  
Age:14 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518100A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 DAY	Nasal Congestion		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
		Rhinorrhoea		Allergy Shots	C		
		Sneezing		Proventil	C	Glaxosmithkline	
				Singulair	C		

Date:08/31/04ISR Number: 4438307-4Report Type:Periodic  
Age:15 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518179A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Papular		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438308-6Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518210A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Accidental Overdose		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Feeling Abnormal		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
		Nausea		Iron Supplement	C		
		Tinnitus		Calcium Supplement	C		
				Yasmin	C		
				Vitamin D Supplement	C		

Date:08/31/04ISR Number: 4438309-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0518227A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	
4 MON		Fatigue		No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438310-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0518228A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oedema Peripheral		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6 MON			Hytrin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438311-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518231A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Agitation Drug Ineffective Heart Rate Increased Insomnia Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438312-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518236A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Single Initial or Prolonged dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438313-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518246A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day 2 MON		Weight Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438314-1Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518361A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG See dosage text		Dry Mouth Weight Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438315-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518423A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438316-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518518A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day				Luvox	C		
				Anafranil	C		
				Xanax	C		

Date:08/31/04ISR Number: 4438317-7Report Type:Periodic  
Age:23 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518522A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Deafness		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				No Concurrent Medication	C		
2 WK							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438318-9Report Type:Periodic  
Age:35 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518537A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown	1	MON	Blood Pressure Increased	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Heart Rate Increased	No Concurrent Medication	C		
			Hypervigilance				

Date:08/31/04ISR Number: 4438319-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518540A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice			Metrorrhagia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
per day	7	MON		Provera	C		
				Atenolol	C		

Date:08/31/04ISR Number: 4438320-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518547A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day			Lethargy	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
200MG per day			Nervousness	Wellbutrin Sr	SS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438321-9Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518549A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day			Dysmenorrhoea	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438322-0Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0518612A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day							

300MG Per day

Date:08/31/04ISR Number: 4438323-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518621A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Rash

Date:08/31/04ISR Number: 4438324-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518630A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

300MG Per day 4 WK

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438325-6Report Type:Periodic  
Age:43 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0518642A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus	Health	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4 WK		Professional	No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438326-8Report Type:Periodic  
Age:52 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518712A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Excessive Masturbation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	MON	Libido Increased					

Date:08/31/04ISR Number: 4438327-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518753A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bruxism Dysgeusia Salivary Hypersecretion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438328-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518785A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pollakiuria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day				Klonopin	C		

Date:08/31/04ISR Number: 4438329-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518791A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Feeling Jittery		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	7 DAY	Heart Rate Increased		Prozac	C		
		Insomnia					

Date:08/31/04ISR Number: 4438330-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0518798A  
 Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 MON			Adderall	C		

Date:08/31/04ISR Number: 4438331-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0518819A  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Asthenia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Drug Ineffective		Unknown Medication	C		
		Headache		Effexor	C		
		Nausea		Vistaril	C		
		Sensation Of Heaviness		Vitamins	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438332-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518984A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Zoloft	C		
				Ortho Tricyclen	C		

Date:08/31/04ISR Number: 4438333-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519027A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:08/31/04ISR Number: 4438334-7Report Type:Periodic  
Age:55 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0519035A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK	Crying		Estrogen	C		
		Depressive Symptom		Cartia	C	Glaxosmithkline	
		Stool Analysis Abnormal					
		Vision Blurred					

Date:08/31/04ISR Number: 4438335-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519053A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK	Vitreous Floaters		Zoloft	C		
				Ambien	C		

Date:08/31/04ISR Number: 4438336-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0519064A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Affect Lability		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:08/31/04ISR Number: 4438337-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0519065A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Screen Positive		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				Trazodone	C		
				Levothyroxine	C	Glaxosmithkline	

Date:08/31/04ISR Number: 4438338-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0519070A  
Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 3 WK		Palpitations		Wellbutrin	SS	Glaxosmithkline	ORAL
200MG Per day 1 WK		Throat Tightness		No Concurrent			
		Visual Disturbance		Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438339-6Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #S04-USA-04077-01

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day	4	MON	Professional	Lexapro	SS		ORAL
10MG Per day							

Date:08/31/04ISR Number: 4438340-2Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519210A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3	WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Headache					
		Insomnia		Inderal	C		
		Lethargy		Topamax	C		

Date:08/31/04ISR Number: 4438341-4Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519216A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Abdominal Pain Upper					
				Methadone	C	Glaxosmithkline	
				Percocet	C		

Date:08/31/04ISR Number: 4438342-6Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519248A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
1	MON						
		Tremor		Duragesic	C		
				Xanax	C		
				Prednisone	C		
				Protonix	C		

Lexapro	C
Klonopin	C
Altace	C
Lipitor	C
Clindamycin	C
Levaquin	C
Neurontin	C
Percocet	C

Date:08/31/04ISR Number: 4438343-8Report Type:Periodic  
 Age:44 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0519390A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypertension		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	9	MON					

Date:08/31/04ISR Number: 4438344-XReport Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519391A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438345-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519405A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438346-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519410A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MGML Per day	3 WK	Adverse Event		Wellbutrin Xl  No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438347-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519535A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
600MG Per day		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
75MG Per day		Insomnia		Wellbutrin	C	Glaxosmithkline	ORAL
		Muscle Twitching Overdose					

Date:08/31/04ISR Number: 4438348-7Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0519541A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Headache		Wellbutrin Xl No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438349-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519550A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Palpitations					

Date:08/31/04ISR Number: 4438350-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519695A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Memory Impairment		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438351-7Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519701A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Panic Reaction		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Xanax	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438352-9Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519704A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Wellbutrin Sr	SS	Glaxosmithkline	ORAL
				Actonel	C		
				Nexium	C		
				Oral Contraceptives	C		
				Ativan	C		

Date:08/31/04ISR Number: 4438353-0Report Type:Periodic  
Age:59 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0519706A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	12	DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Lipitor	C		
				Aspirin	C	Glaxosmithkline	
				Vitamins	C		

Date:08/31/04ISR Number: 4438354-2Report Type:Periodic  
Age:45 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519707A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3	WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438355-4Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519723A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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150MG Per day	1	MON	Dyspnoea	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Heart Rate Increased	Lexapro	C		
			Migraine				
			Nausea				
			Pollakiuria				
			Tremor				

Date:08/31/04ISR Number: 4438356-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0519728A  
 Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Blood Pressure Increased	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Birth Control	C		ORAL

Date:08/31/04ISR Number: 4438357-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0519735A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				Seroquel	C		
Initial or Prolonged							
Other							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438358-XReport Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519736A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	4 DAY	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Thinking Abnormal		No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438359-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519855A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	2 MON	Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dyspnoea					

Date:08/31/04ISR Number: 4438360-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519869A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Cough		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Rhinitis					

Date:08/31/04ISR Number: 4438361-XReport Type:Periodic  
Age:37 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0519875A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	19 DAY	Erythema		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Pruritus		Dhea	C		
		Rash		Herbal Medicine	C		

Date:08/31/04ISR Number: 4438362-1Report Type:Periodic  
Age:44 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0519880A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3	WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Ambien	C		
				Novocaine	C		

Date:08/31/04ISR Number: 4438363-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0519882A  
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300MG Per day	3	DAY		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438364-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0519896A  
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	8	DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Ultram	SS		
				Norvasc	C		
				Zyrtec	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438365-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519900A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mouth Ulceration		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2	WK						

Date:08/31/04ISR Number: 4438366-9Report Type:Periodic  
 Age:50 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519908A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
4	DAY	Tinnitus		Ultram	C		

Date:08/31/04ISR Number: 4438367-0Report Type:Periodic  
 Age:19 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519923A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG	Per day			No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438368-2Report Type:Periodic  
 Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520077A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG	Per day	Depression Headache Hot Flush Nervousness Vision Blurred		No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438369-4Report Type:Periodic  
Age:17 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0520093A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 1 DAY		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Aura Grand Mal Convulsion Headache		Remeron	C		

Date:08/31/04ISR Number: 4438370-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520094A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nipple Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438371-2Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520107A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 450MG Per day 2 WK		Vaginal Haemorrhage		Wellbutrin	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438372-4Report Type:Periodic  
Age:26 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0520111A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	300MG Per day	MON	Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
				Risperdal	C		
				Prozac	C		

Date:08/31/04ISR Number: 4438373-6Report Type:Periodic  
Age:34 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520281A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2	MON	Amnesia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Emotional Disorder	Prozac	C		
			Personality Change				

Date:08/31/04ISR Number: 4438374-8Report Type:Periodic  
Age:33 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0520285A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
105 DAY			Blood Pressure Diastolic	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Increased	Hyzaar	C		ORAL

Date:08/31/04ISR Number: 4438375-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520286A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Dizziness	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Ill-Defined Disorder	Toprol	C		

Date:08/31/04ISR Number: 4438376-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520288A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL
4 MON				Toprol	C		
Initial or Prolonged							

Date:08/31/04ISR Number: 4438377-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0520300A  
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3 DAY	Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438378-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0520310A  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	21 WK	Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Alopecia Effluvium		No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438379-7Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520312A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	8 WK						

Date:08/31/04ISR Number: 4438380-3Report Type:Periodic  
Age:28 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0520319A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN	150MG	Unknown 23 DAY					
Initial or Prolonged		Overdose		Diphenhydramine	C		
				Benzodiazepine	C		
				Cannabis	C		
				Alcohol	C		
				Sleeping Pill	C		

Date:08/31/04ISR Number: 4438381-5Report Type:Periodic  
Age:13 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520321A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Behaviour		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438382-7Report Type:Periodic  
Age:56 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520373A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Overdose		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
600MG Single							
dose		Tremor		Concerta	C		

Ritalin C  
Klonopin C

Date:08/31/04ISR Number: 4438383-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520375A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
450MG Per day	3	MON	Irritability	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Stool Analysis Abnormal	Unknown Medication	C		

Date:08/31/04ISR Number: 4438384-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520387A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day			Sleep Walking	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438385-2Report Type:Periodic  
Age:46 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0520485A

Outcome  
PT  
Fatigue  
Migraine  
Muscle Twitching  
Nightmare

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Palpitations Paraesthesia Sensation Of Heaviness Tremor	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3 WK			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Diovan	C		
				Aciphex	C		
				Zyrtec	C	Glaxosmithkline	
				Singulair	C		
				Zomig	C		
				Vioxx	C		

Date:08/31/04ISR Number: 4438387-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0520508A  
 Age: Gender: I/FU:I

Outcome Dose	Duration	PT Vomiting	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438388-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0520509A  
 Age: Gender: I/FU:I

Outcome Dose	Duration	PT Blood Glucose Increased	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438389-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0520521A  
 Age: Gender:Female I/FU:F

Outcome Dose Other	Duration	PT Convulsion	Report Source	Product	Role	Manufacturer	Route
300MG Per day				Wellbutrin	PS	Glaxosmithkline	ORAL
				Alcohol	C		

Date:08/31/04ISR Number: 4438390-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0520523A  
 Age: Gender:Female I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:08/31/04ISR Number: 4438391-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0520524A  
Age:17 YR Gender:Female I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	DAY			No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438392-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0520541A  
Age: Gender:Female I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Myalgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438393-1Report Type:Periodic  
Age:41 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0520546A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2 MON			No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438394-3Report Type:Periodic  
Age:17 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0521113A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				No Concurrent Medication	C		
450MG Per day							

Date:08/31/04ISR Number: 4438395-5Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0521115A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438396-7Report Type:Periodic  
Age:17 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0521117A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Furuncle		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Stevens-Johnson Syndrome		No Concurrent Medication	C		
300MG Per day							

Date:08/31/04ISR Number: 4438397-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0521119A  
Age:13 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438398-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0521834A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Geodon	C		
				Temazepam	C		

Date:08/31/04ISR Number: 4438399-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0522803A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4438421-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493781A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Loss Poor	Consumer	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438422-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494320A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash	Health Professional	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/31/04ISR Number: 4438423-7Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0500121A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus	Health Professional	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Unspecified Drugs	C		

Date:08/31/04ISR Number: 4438424-9Report Type:Periodic  
Age:30 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0507659A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Grand Mal Convulsion		Advair	C	Glaxosmithkline	
28 DAY		Tongue Biting		Beconase	C	Glaxosmithkline	
				Singulair	C		
				Ultram	C		
				Maxalt	C		
				Vioxx	C		
				Allegra D	C		

Date:08/31/04ISR Number: 4438425-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0507906A  
Age:15 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Osteoporosis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
250MG Per day				Depakote Er	C		ORAL
1500MG Per							
day				Celexa	C		ORAL
20MG Per day							

Date:08/31/04ISR Number: 4438426-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510522A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bruxism		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				No Concurrent Medication	C		

Date:08/31/04ISR Number: 4438427-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0513782A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pollakiuria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/01/04ISR Number: 4439062-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0524069A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Suicidal Ideation		Wellbutrin No Concurrent Medication	PS  C	Glaxosmithkline	

Date:09/01/04ISR Number: 4439066-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0335857A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Accident	Health  Professional	Zyban	PS	Glaxosmithkline	ORAL
150MG	Unknown						

Date:09/01/04ISR Number: 4439067-3Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0339775A

Age:58 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150MG See Initial or Prolonged dosage text	6 DAY	Amnesia  Blood Pressure Decreased  Cardiac Flutter  Chest Pain  Feeling Abnormal  Headache  Hiatus Hernia  Insomnia	Health  Professional	Zyban    Lipitor  Lopid   Pariet   Endep   Premarin	PS    C  C   C   C	Glaxosmithkline	ORAL    ORAL  ORAL   ORAL   ORAL
40G Per day							
60G Twice per day							
20G Twice per day							
10G As required							

Date:09/01/04ISR Number: 4439078-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0343937A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
Disability		Cerebral Infarction					
150MG Unknown		Hemiplegia		Levothyrox	C	Glaxosmithkline	
				Tenormine	C		
				Mopral	C		

Date:09/01/04ISR Number: 4439721-3Report Type:Direct Company Report #CTU 226057  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr 100 Mg			
Death		Completed Suicide		Gsk	PS	Gsk	
		Treatment Noncompliance					
1 TABLET							
TWICE A DA							

Date:09/01/04ISR Number: 4440010-1Report Type:Expedited (15-DaCompany Report #A-US2004-07003  
Age:41 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain Upper
Initial or Prolonged	Antibody Test Positive
Other	Constipation
	Hepatitis
	International Normalised
	Ratio Increased

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Freedom Of Information (FOI) Report

Nephrolithiasis

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
125 MG, BID, ORAL : 62.5 MG, BID, ORAL		Study Health Professional Distributor	Tracleer (Bosentan) Tablet	PS		ORAL
20 MG, QD			Lovastatin(Lovastatin)	SS		
			Wellbutrin(Amfebutamone Hydrochloride)	SS		
			Coumadin(Warfarin Sodium)	C		
			Prinivil(Lisinopril)	C		
			Aspirine(Acetylsalicylic Acid)	C		
			Milk Of Magnesia (Magnesium Hydroxide)	C		

Date:09/01/04ISR Number: 4440362-2Report Type:Direct  
Age:36 YR Gender:Female I/FU:I

Company Report #CTU 226055

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other SEE IMAGE PER FDA ONSITE REP SEE IMAGE PER JW FDA ONSITE REP		Balance Disorder Drug Ineffective Muscular Weakness Paraesthesia Vision Blurred		Zoloft Down To 50mg Unknown	PS		
				Wellbutrin Xl 150 Mgs Un Known	SS		

Date:09/02/04ISR Number: 4440061-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0335857A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Accident	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown			Professional				

Date:09/02/04ISR Number: 4440062-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0337371A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blindness Transient	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG Per day 1 DAY			Professional	Betablocker	C		ORAL
				Zocor	C		ORAL
				Kardegic	C		ORAL
75MG Per day				Stilnox	C		ORAL

Date:09/02/04ISR Number: 4440073-3Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0344111A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination		Zyban	PS	Glaxosmithkline	
5 DAY		Sensation Of Pressure					
		Tinnitus					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/02/04ISR Number: 4444581-0Report Type:Expedited (15-DaCompany Report #2004059295

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zyrtec-D 12 Hour			
Other		Chronic Obstructive Pulmonary Disease		(Cetirizine, Pseudoephedrine)	PS		
		Dental Operation		Pramipexole			
		Hypersensitivity		(Pramipexole)	SS		
		Polyneuropathy		Clonazepam			
		Sinus Disorder		(Clonazepam)	SS		
				Mirtazapne			
				(Mirtazapine)	SS		
				Amitriptyline			
				(Amitriptyline)	SS		
				Bupropion			
				Hydrochloride			
				(Bupropion			
				Hydrochloride)	SS		
				Montelukast Sodium			
				(Montelukast Sodium)	C		
				Combivent			
				(Ipratropium			
				Bromide, Salbutamol			
				Sulfate)	C		
				Mirtazapine			
				(Mirtazapine)	C		
				Esomeprazole			
				(Esomeprazole)	C		
				Salbutamol			
				(Salbutamol)	C		

Date:09/02/04ISR Number: 4446647-8Report Type:Expedited (15-DaCompany Report #2004-119599-NL

Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction		Estrogen Progestron	PS		ORAL
DF		Haemorrhage		Bupropion			
		Seasonal Allergy		Hydrochloride	SS		ORAL
150 MG	13 DAY			Fexofenadine	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fatigue	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
20 DAY		Hemiparesis	Professional	Diovan	C		ORAL
		Migraine		Aciphex	C		ORAL
		Muscle Twitching		Zyrtec	C	Glaxosmithkline	ORAL
		Nightmare		Singulair	C		ORAL
		Palpitations		Vioxx	C		ORAL
		Paraesthesia		Lexapro	C		ORAL
.5TAB Per day		Sensation Of Heaviness					
		Tremor					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/03/04ISR Number: 4440821-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0524499A  
 Age:19 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	100MG Per day	2 WK	Agitation	Wellbutrin	PS	Glaxosmithkline	ORAL
			Headache	No Concurrent Medication	C		
			Psychiatric Symptom				
			Suicidal Ideation				

Date:09/03/04ISR Number: 4440823-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0524503A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day	1 DAY	Homicidal Ideation	Wellbutrin	PS	Glaxosmithkline	ORAL
				Zoloft	C		

Date:09/03/04ISR Number: 4440826-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0524516A  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	1 MON		Amnesia	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged			Convulsion	Alcohol	C		
			Overdose				

Date:09/03/04ISR Number: 4440839-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0342217A  
 Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Anorexia	Zyban	PS	Glaxosmithkline	
			Dizziness	Fluvax	C	Glaxosmithkline	
INTRAMUSCULAR			Flatulence				
			Middle Insomnia				
			Sleep Disorder				

Somnolence  
Speech Disorder  
Thought Blocking  
Tremor  
Urticaria  
Visual Acuity Reduced  
Visual Disturbance

Date:09/03/04ISR Number: 4440844-3Report Type:Expedited (15-DaCompany Report #DK-GLAXOSMITHKLINE-B0343749A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	ORAL
		Ill-Defined Disorder		Alcohol	C		
		Laceration					
		Wound					

Date:09/03/04ISR Number: 4440847-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0344156A  
Age:56 YR Gender:Male I/FU:I

Outcome	PT
Other	Anxiety
	Drug Dependence
	Euphoric Mood

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Excitability Formication Malaise	Report Source	Product	Role	Manufacturer	Route
150MG See dosage text		Medication Error		Zyban	PS	Glaxosmithkline	ORAL
		Pallor					
		Transaminases Increased		Lexomil	C		

Date:09/03/04ISR Number: 4440848-0Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0344204A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 1TAB Unknown Initial or Prolonged		Adverse Event	Health	Zyban	PS	Glaxosmithkline	ORAL
		Aggression	Professional	Valium	SS		
		Amnesia		Warfarin	SS	Glaxosmithkline	
		Mental Disorder		Tritace	SS		
		Overdose		Panadeine Forte	SS	Glaxosmithkline	
		Suicide Attempt					

Date:09/07/04ISR Number: 4441953-5Report Type:Expedited (15-DaCompany Report #2004-03848  
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day 3 DAY 25MG Per day		Dizziness		Bupropion	PS	Glaxosmithkline	ORAL
		Loss Of Consciousness		Atenolol	C		
		Syncope					
		Vomiting					

Date:09/07/04ISR Number: 4441955-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0524551A  
Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Intentional Misuse Wellbutrin PS Glaxosmithkline ORAL  
1 MON  
Overdose  
Suicide Attempt

Date:09/07/04ISR Number: 4441989-4Report Type:Expedited (15-DaCompany Report #CH-GLAXOSMITHKLINE-B0344090A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State		Zyban	PS	Glaxosmithkline	ORAL
300MG Per day		Crying		Coaprovel	SS		ORAL
		Depressed Mood					
		Depression					
		Drug Withdrawal Syndrome					
		Fatigue					
		Feeling Abnormal					

Date:09/07/04ISR Number: 4442955-5Report Type:Direct Company Report #CTU 226470  
Age: Gender:Female I/FU:I

Outcome	PT
Other	Contusion
	Dehydration
	Disturbance In Attention
	Dizziness
	Dysgeusia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache Hot Flush Impaired Work Ability					
150MG	ONCE	Influenza Like Illness Nausea		Wellbutrin Xl Glaxosmithkline	PS	Glaxosmithkline	ORAL
DAILY	ORAL	Rash  Tremor Vomiting					

Date:09/07/04ISR Number: 4443023-9Report Type:Expedited (15-DaCompany Report #A-US2004-07003  
Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	SEE IMAGE	Antibody Test Positive Hepatitis	Study Health	Tracleer (Bosentan ) Tablet	PS		
Other		International Normalised Ratio Increased Nephrolithiasis	Professional Distributor	Wellbutrin (Amfebutamone Hydrochloride) Coumadin (Warfarin Sodium) Prinivil (Lisinopril) Aspirine (Acetylsalicylic Acid) Milk Of Magnesia (Magnesium Hydroxide)	SS  C C  C C		

Date:09/07/04ISR Number: 4444753-5Report Type:Direct Company Report #USP 56827  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
TABLET,  EXTENDED		Medication Error		Wellbutrin Sr	PS	Glaxosmithkline	

RELEASE

Wellbutrin Xl SS Glaxosmithkline

TABLET ,

EXTENDED

RELEASE

Date:09/07/04ISR Number: 4447128-8Report Type:Direct

Company Report #USP 56822

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Medication Error	Wellbutrin Sr	PS	Glaxosmithkline	

TABLET ,

EXTENDED

RELEASE

Wellbutrin Xl SS Glaxosmithkline

TABLET,

EXTENDED

RELEASE

Date:09/07/04ISR Number: 4455142-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0524551A

Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Health	Wellbutrin	PS	Glaxosmithkline	ORAL
1 MON			Professional				
			Suicide Attempt				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/08/04ISR Number: 4443650-9Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0524692A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day Initial or Prolonged		Dysphonia Tobacco Abuse Tremor		Zyban	PS	Glaxosmithkline	ORAL

Date:09/08/04ISR Number: 4443654-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0524757A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG Per day 3 MON		Agitation Confusional State Disorientation Fall Heart Rate Increased Hypoaesthesia Palpitations Panic Attack Syncope		Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:09/08/04ISR Number: 4443655-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0524762A  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 200MG Twice per day	4 YR	Nystagmus		Wellbutrin Seroquel Zinc Sulfate Levothyroxine Ursodiol Propranolol Ranitidine	PS C C C C C C	Glaxosmithkline Glaxosmithkline	ORAL

Date:09/08/04ISR Number: 4443659-5Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0336823A  
Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Agitation					
per day	61 DAY	Asthenia		No Concurrent			
		Decreased Interest		Medication	C		
		Depressed Mood					
		Suicidal Ideation					

Date:09/08/04ISR Number: 4443665-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0343830A  
Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Chest Pain		Zyban	PS	Glaxosmithkline	ORAL
150MG per day 3	DAY	Hyperhidrosis		Vioxx	C		ORAL
Initial or Prolonged		Nausea					
12.5MG Per		Pain In Extremity		Co-Codamol	C		ORAL
day				Priadel	C	Glaxosmithkline	ORAL
400MG per day 57	WK						



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/08/04ISR Number: 4447167-7Report Type:Direct  
Age:50 YR Gender:Female I/FU:I

Company Report #CTU 226644

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Migraine		Bupriopion	PS		ORAL
150 MG SA BID							
		Pharmaceutical Product					
PO		Complaint					

Date:09/09/04ISR Number: 4444865-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0514663A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error	Health	Wellbutrin	PS	Glaxosmithkline	
		Pharmaceutical Product	Professional				
		Complaint					

Date:09/09/04ISR Number: 4444867-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0517054A  
Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Anxiety	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
2 YR							
		Completed Suicide					
		Dissociation					
		Hostility					
		Weight Increased					

Date:09/09/04ISR Number: 4444900-5Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0344183A  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Psoriasis		Bupropion			
Initial or Prolonged				Hydrochloride	PS	Glaxosmithkline	ORAL
150MG Per day 7 DAY							

.1MG per day

Date:09/09/04ISR Number: 4444902-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0344253A  
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia		Zyban	PS	Glaxosmithkline	ORAL
150MG per day	2	MON					
		Blister					
		Oedema Peripheral					
		Rash					

Date:09/09/04ISR Number: 4444904-2Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0344335A  
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema		Zyban	PS	Glaxosmithkline	ORAL
300MG Per day	5	WK					
		Chills					
		Oedema Mouth					
		Swollen Tongue					
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/09/04ISR Number: 4445382-XReport Type:Expedited (15-DaCompany Report #US-MERCK-0408USA02318  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abdominal Pain Upper		Mevacor	PS	Merck & Co., Inc	ORAL
Initial or Prolonged		Alanine Aminotransferase		Tracleer	SS		ORAL
Other		Increased		Tracleer	SS		ORAL
UNKNOWN		Anorexia		Wellbutrin	SS		
UNKNOWN		Aspartate		Aspirine	C		
UNKNOWN		Aminotransferase		Prinivil	C		
UNKNOWN		Increased		Magnesia [Milk Of]	C		
UNKNOWN		Blood Alkaline		Coumadin	C		
UNKNOWN		Phosphatase Increased					
		Blood Bilirubin Increased					
		Constipation					
		Hepatitis					
		International Normalised					
		Ratio Increased					

Date:09/10/04ISR Number: 4446022-6Report Type:Expedited (15-DaCompany Report #BPC-SR-04-034  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Remeron	C		ORAL

Date:09/10/04ISR Number: 4446032-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0340981A  
 Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Alcoholic	Health	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Injury	Professional				
per day	3 DAY						

1 DAY

Alcohol

SS

Depo-Provera

C

INTRAMUSCULAR

Date:09/10/04ISR Number: 4446045-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0344251A

Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depressed Mood		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	11 DAY	Feeling Abnormal Insomnia					

Date:09/10/04ISR Number: 4448384-2Report Type:Direct Company Report #CTU 226841

Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Convulsion		Wellbutrin Sr 150 Mg			
Intervention to				Bid	PS		
ORAL BID	1 YR						
Prevent Permanent				Strattera 40 Mg	SS		ORAL
ORAL QD	1 MON						
Impairment/Damage							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/13/04  
 Age:57 YR  
 Gender:Female  
 I/FU:I

Report Type:Direct  
 Company Report #CTU 226935

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ONE PO BID		Drug Hypersensitivity		Wellbutrin Sr 100 Mg	PS		ORAL
		Drug Tolerance Decreased Gastrointestinal Disorder Pharmaceutical Product Complaint					

Date:09/13/04  
 Age:35 YR  
 Gender:Female  
 I/FU:I

Report Type:Expedited (15-Da  
 Company Report #HQWYE752701SEP04

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
SEE IMAGE/OVER 4-5 DAYS		Drug Interaction Drug Level Increased Electrocardiogram Qrs Complex Prolonged	Literature	Effexor (Venlafaxine Hydrochloride, Unspec)	PS		
INTRA VENOUS	400 MG ,	Medical Device Complication		Avelox (Moxifloxacin, )	SS		
INTRA VENOUS				Bupropion (Amfebutamone, )	SS		
SEE IMAGE/OVER 48 HOURS				Carvedilol (Carvedilol, )	SS		
3.125 MG 2X PER 1 DAY				Vancomycin (Vancomycin) Aspirin	C		

(Acetylsalicylic  
 Acid) C  
 Furosemide C  
 (Furosemide) C  
 Spironolactone C  
 (Spironolactone) C  
 Milrinone C  
 (Milrinone) C  
 Clonazepam C  
 (Clonazepam) C  
 Calcium (Calcium) C  
 Multivitamins  
 (Ascorbic  
 Acid/Ergocalciferol/  
 Folic  
 Acid/Nicotinamide/Pa C

Date:09/15/04ISR Number: 4449918-4Report Type:Expedited (15-DaCompany Report #BPC-SR-04-034  
 Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Chest Discomfort	Health	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
17 DAY		Dyspnoea	Professional	Remeron	C		ORAL
30MG Per day		Myocardial Infarction Pain In Jaw					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/15/04ISR Number: 4449925-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0525433A  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	5 YR	Condition Aggravated Suicidal Ideation	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
				Paxil	C	Glaxosmithkline	
				Elavil	C	Glaxosmithkline	
				Klonopin	C		

Date:09/15/04ISR Number: 4449935-4Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0343937A  
 Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 150MG Unknown Hospitalization - Initial or Prolonged Disability	WK	Aphasia Bradyphrenia Cerebral Infarction Cerebrovascular Accident Coma Confusional State Disturbance In Attention Headache Hemiparesis Hemiplegia Hypertension Hypertensive Crisis Hypertriglyceridaemia	Health Professional	Zyban Levothyrox Tenormine Mopral Female Hormones	PS C C C C	Glaxosmithkline Glaxosmithkline	ORAL ORAL

Date:09/15/04ISR Number: 4449937-8Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0344111A  
 Age:26 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	5 DAY	Hallucination Hallucination, Auditory	Health Professional	Zyban	PS	Glaxosmithkline	ORAL

500MG Twice                      Hallucination, Visual                      Epilim                      C  
per day                              Sensation Of Pressure

Date:09/15/04ISR Number: 4449938-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0344204A  
Age:                      Gender:Male                      I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose                      Duration						
Hospitalization - 1TAB Unknown	Adverse Event	Consumer	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged	Aggression		Valium	SS		
	Amnesia		Warfarin	SS	Glaxosmithkline	
	Mental Disorder		Tritace	SS		
	Overdose		Panadeine Forte	SS	Glaxosmithkline	
	Suicide Attempt					

Date:09/15/04ISR Number: 4449941-XReport Type:Expedited (15-DaCompany Report #PT-GLAXOSMITHKLINE-B0344722A  
Age:40 YR                      Gender:Male                      I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose                      Duration						
Hospitalization - Initial or Prolonged	Angioneurotic Oedema		Zyban	PS	Glaxosmithkline	ORAL
	Pruritus					
	Pyrexia					
	Rash Generalised					
	Skin Lesion					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/15/04ISR Number: 4449943-3Report Type:Expedited (15-DaCompany Report #IL-GLAXOSMITHKLINE-B0344959A

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Chest Pain		Zyban	PS	Glaxosmithkline	
UNKNOWN	2 WK					
Initial or Prolonged	Feeling Abnormal		Aspirin	C	Glaxosmithkline	
UNKNOWN			Simvastatin	C		
UNKNOWN			Tritace	C		
UNKNOWN			Vitamin B12 + B6	C		

Date:09/15/04ISR Number: 4451843-XReport Type:Direct

Company Report #CTU 227110

Age:31 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Arthralgia		Effexor 75 Mg	PS		ORAL
75 MG DAILY						
ORAL	Diarrhoea					
	Headache		Wellbutrin 100 Mg	SS		ORAL
200 MG DAILY						
ORAL	Nausea					
	Night Sweats					
	Pyrexia					
	Vomiting					

Date:09/16/04ISR Number: 4451321-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0525619A

Age:26 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2 MON		Strattera	C		

Date:09/16/04ISR Number: 4451322-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0525720A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin XL	PS	Glaxosmithkline	ORAL
300MG Per day	4 DAY	Mouth Injury					
		Muscle Twitching					
		Vomiting					

Date:09/16/04ISR Number: 4451339-5Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0345167A  
Age:17 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Creatinine		Zyban	PS	Glaxosmithkline	ORAL
50TAB per day		Increased					
Initial or Prolonged		Convulsion					
		Gait Disturbance					
		Hallucination					
		Memory Impairment					
		Mydriasis					
		Myoglobin Blood Increased					
		Overdose					
		Tachycardia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/16/04ISR Number: 4451340-1Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0345169A  
 Age:26 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 80TAB per day Initial or Prolonged		Agitation Confusional State Convulsion Dysarthria Hallucination Hypertension Overdose Somnolence Tachycardia		Zyban	PS	Glaxosmithkline	ORAL

Date:09/16/04ISR Number: 4453046-1Report Type:Direct Company Report #CTU 227348  
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage 150 MG TWICE DAILY ORAL		Fatigue Heart Rate Increased Hypoventilation Insomnia Pharmaceutical Product Complaint		Bupropion Sr Equiv. Of 150 Mg Wellbutrin Sr - Teva Laboratories	PS	Teva Laboratories	ORAL

Date:09/16/04ISR Number: 4454308-4Report Type:Direct Company Report #CTU 227395  
 Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abnormal Behaviour Agitation Completed Suicide Hostility Insomnia Irritability		Welbutrin	PS		

Date:09/17/04ISR Number: 4454421-1Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 227416

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin Sr 150	PS		ORAL
150 MG TID BY							
MOUTH							

Date:09/17/04ISR Number: 4454536-8Report Type:Direct  
Age:54 YR Gender:Female I/FU:I

Company Report #CTU 227489

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying		Bupropion Hcl			
Other		Feeling Abnormal		Sustained-Release			
		Hallucination, Visual		Tablets 100 Mg	PS	Mfd By Watson	
		Mood Altered				(Imprint "Wpi, 858")	
300 MG DAILY		Pharmaceutical Product					
AS 1 TID	1	Complaint		Allegra	C		
		Suicidal Ideation		Albuterol	C		
		Tearfulness		Advair	C		
		Thinking Abnormal		Zyrtec	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/17/04ISR Number: 4458216-4Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-JNJFOC-20040304012

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Thrombocytopenia	Health Professional	Topamax (Topiramate) Unspecified	PS		
300 MG, IN 1							
DAY							
				Wellbutrin (Bupropion Hydrochloride)	SS		
200 MG, 2 IN							
1 DAY							
				Lexapro (All Other Therapeutics Products)	SS		
20 MG, IN 1							
DAY							

Date:09/20/04ISR Number: 4453953-XReport Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0525756A  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Aggression		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown 44 DAY							
		Drug Ineffective Ill-Defined Disorder		Rivotril (Clonazepam)	C		

Date:09/20/04ISR Number: 4453954-1Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0525758A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Aggression Alcoholism		Zyban	PS	Glaxosmithkline	ORAL

Date:09/20/04ISR Number: 4453971-1Report Type:Expedited (15-DaCompany Report #PT-GLAXOSMITHKLINE-B0344722A  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 15 DAY		Angioneurotic Oedema	Health	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Bronchospasm Oedema Peripheral Pyrexia Skin Lesion Urticaria	Professional				

Date:09/20/04ISR Number: 4453985-1Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0345610A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN	2TAB per day	Bradyphrenia		Zyban	PS	Glaxosmithkline	
		Dizziness Syncope Vision Blurred		No Concurrent Medication	C		

Date:09/20/04ISR Number: 4454746-XReport Type:Direct Company Report #CTU 227557  
Age:27 YR Gender:Female I/FU:I

Outcome	PT
Other	Condition Aggravated Depression

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Effect Decreased Medication Error Pharmaceutical Product	Report Source	Product	Role	Manufacturer	Route
150 MG 2 TIMES ORAL		Complaint Somnolence		Budeprion 150 Mg Teva	PS	Teva	ORAL

Date:09/20/04ISR Number: 4454769-0Report Type:Expedited (15-DaCompany Report #2004UW19220  
Age: Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 10 MG DAILY Initial or Prolonged PO			Convulsion	Foreign Health Professional Other	Crestor  Wellbutrin	PS  SS		ORAL

Date:09/21/04ISR Number: 4454868-3Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0526395A  
Age:51 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice  per day			Glaucoma		Zyban	PS	Glaxosmithkline	ORAL

Date:09/21/04ISR Number: 4457315-0Report Type:Expedited (15-DaCompany Report #B0344156A  
Age:46 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other  150 MG/SEE			Anhedonia Anxiety Asthenia	Foreign Health Professional	Zyban-Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

DOSAGE  
 TEXT/ORAL  
 Drug Dependence  
 Euphoric Mood  
 Excitability  
 Formication  
 Intentional Misuse  
 Malaise  
 Medication Error  
 Morbid Thoughts  
 Pallor  
 Transaminases Increased

Bromazepam C

Date:09/21/04ISR Number: 4457395-2Report Type:Expedited (15-DaCompany Report #B0312762A  
 Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged 150 MG/ ORAL		Agitation Anxiety Insomnia  Overdose Suicidal Ideation Suicide Attempt	Foreign	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/21/04ISR Number: 4457474-XReport Type:Expedited (15-DaCompany Report #A0525434A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Balance Disorder Cerebrovascular Accident Dysphemia	Consumer	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/PER DAY/ORAL		Hallucination, Visual		Lansoprazole Oestradiol Thyroxine Sodium Hydrochlorothiazide	C C C C		

Date:09/21/04ISR Number: 4457477-5Report Type:Expedited (15-DaCompany Report #A0525250A

Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Crying Impaired Driving Ability Vision Blurred	Health Professional	Wellbutrin Xl Tablets-Extended Release (Bupropion Hydrochloride)	PS		ORAL
PER DAY/ORAL				Aripiprazole Oxcarbazepine	C C		

Date:09/21/04ISR Number: 4457486-6Report Type:Expedited (15-DaCompany Report #A0519334A

Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Depression Muscular Weakness Myalgia Pharmaceutical Product	Consumer	Bupropion Hydrochloride Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ PER DAY/ ORAL		Complaint					

Suicidal Ideation

Paroxetine	C
Hydrochloride	C
Topiramate	C
Ramipril	C
Simvastatin	C

Date:09/22/04ISR Number: 4455959-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0526418A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Foreign Body Trauma		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:09/22/04ISR Number: 4457195-3Report Type:Expedited (15-DaCompany Report #S04-USA-04018-01

Age:42 YR Gender:Male I/FU:F

Outcome	PT
Other	Alopecia
Required	Blood Pressure
Intervention to	Fluctuation
Prevent Permanent	Erectile Dysfunction
Impairment/Damage	Exostosis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Fracture Hypertension Joint Dislocation	Report Source	Product	Role	Manufacturer	Route
10 MG QD PO		Nausea Somnolence	Consumer	Lexapro (Escitalopram)	PS		ORAL
0.2 MG BID				Lotrel Clonidine	SS SS		
				Amiloride W/Hydrochlorothiazid e	SS		
				Percocet	SS		
				Percocet	SS		
150 MG				Wellbutrin - Slow Release (Bupropion Hydrochloride)	SS		
				Trazodone	C		
				Nexium (Esomeprazole)	C		
				Bextra (Valdecoxib)	C		
				Carisoprodol	C		

Date:09/23/04ISR Number: 4456996-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0526711A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Feeling Abnormal Overdose		Wellbutrin Xanax	PS SS	Glaxosmithkline	ORAL

Date:09/23/04ISR Number: 4456997-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0526785A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion Depressed Level Of Consciousness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:09/23/04ISR Number: 4457001-7Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0318010A  
Age:49 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 1TAB Twice	Aggression	Health	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day 5 DAY	Agitation	Professional				
Other	Blood Pressure Increased Depression Dizziness Feeling Abnormal Headache Pharmaceutical Product Complaint					

Date:09/23/04ISR Number: 4457004-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0333110A  
Age:52 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Arteritis
Initial or Prolonged	Blood Pressure Systolic Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Coronary Artery Stenosis Myocardial Infarction Pain In Extremity	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	6 DAY						

Date:09/23/04ISR Number: 4457025-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0345694A  
Age:26 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
Other		Asthma					
150MG per day	3 DAY			Salbutamol	C	Glaxosmithkline	
100MCG per		Wheezing					
day							
				Seretide	C	Glaxosmithkline	
250MCG Per							
day							

Date:09/23/04ISR Number: 4457029-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0346004A  
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
Hospitalization -		Diabetes Mellitus					
150MG Per day	7 DAY						
Initial or Prolonged		Inadequate Control		Crestor	C		ORAL
		Myocardial Infarction		Diamicron	C		ORAL
205MG Three							
times per day							

Date:09/23/04ISR Number: 4457794-9Report Type:Direct Company Report #USP 56844  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Medication Error	Wellbutrin Sr	PS	Glaxosmith Kline
TABLET, EXTENDED RELEASE			
	Bupropion	SS	Watson
TABLET, EXTENDED RELEASE			
	Budeprion Sr	SS	Teva
TABLET, EXTENDED RELEASE			
	Bupropion	SS	Eon Labs
TABLET, EXTENDED RELEASE			
	Wellbutrin Xl	SS	Glaxosmithkline

Date:09/23/04ISR Number: 4457797-4Report Type:Direct Company Report #USP 56843  
Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error		Bupropion (Generic For Zyban)	PS	Watson/Ndc # 0591-0867-60	
TABLET				Bupropion (Generic For Wellbutrin Sr)	SS	Watson / Ndc # 0591-0839-60	
TABLET							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/23/04ISR Number: 4457965-1Report Type:Direct  
Age: Gender: I/FU:I

Company Report #USP 56873

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Bupropion Xr	PS		
				Bupropion Sr	SS		

Date:09/23/04ISR Number: 4459838-7Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20040903575  
Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aggression	Literature	Acetaminophen			
Other		Agitation	Health	(Acetaminophen)			
ORAL		Atrioventricular Block	Professional	Unspecified	PS		ORAL
ORAL		Blood Creatine Phosphokinase Increased Blood Creatine		Bupropion (Long-Actng) (Bupropion )	SS		ORAL
ORAL		Phosphokinase Mb Increased Blood Glucose Increased		Aspirin (Acetylsalicylic Acid)	SS		ORAL
ORAL		Bradycardia		Caffeine (Caffeine)	SS		ORAL
ORAL		Cardiac Arrest Convulsion		Fluoxetine (Fluoxetine)	SS		ORAL
ORAL		Electrocardiogram Qrs Complex Prolonged		Valproic Acid (Valproic Acid)	SS		ORAL
ORAL		Incoherent Multiple Drug Overdose		Sertraline (Sertraline)	SS		ORAL
		Muscle Rigidity Po2 Increased Pyrexia Ventricular Arrhythmia Ventricular Extrasystoles					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest Coma Completed Suicide	Literature Health Professional	Acetaminophen (Acetaminophen) Unspecified	PS		ORAL
ORAL		Convulsion Haematemesis Heart Rate Increased		Aspirin (Acetylsalicylic Acid)	SS		ORAL
ORAL		Multiple Drug Overdose Respiratory Rate		Bupropion (Bupropion)	SS		ORAL
ORAL		Increased		Ethanol (Ethanol)	SS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day 10 YR	Asthenia	Consumer	Wellbutrin Sr	PS	Glaxosmithkline	
150MG Twice		Depressed Level Of Consciousness		Bupropion Hydrochloride	SS	Glaxosmithkline	ORAL
per day	10 YR	Disturbance In Attention Motor Dysfunction		Lithium	C	Glaxosmithkline	ORAL
		Pharmaceutical Product Complaint Tremor					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/24/04ISR Number: 4458122-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0526801A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other		Asthenia					
3 WK		Choking					
		Distractibility					
		Early Morning Awakening					
		Insomnia					
		Lethargy					
		Memory Impairment					
		Muscle Tightness					
		Nausea					
		Sleep Disorder					
		Speech Disorder					
		Tinnitus					
		Tongue Paralysis					
		Tremor					
		Vision Blurred					
		Weight Decreased					

Date:09/24/04ISR Number: 4458131-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0338492A

Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Health	Zyban	PS	Glaxosmithkline	ORAL
Life-Threatening		Arthralgia	Professional				
150MG See		Asthenia					
		Blood Immunoglobulin E					
		Increased					
		Cough					
		Dysphagia					
		Dyspnoea					
		Insomnia					
		Joint Swelling					
		Oesophageal Spasm					
		Pruritus					
		Serum Sickness					
		Urticaria Generalised					

dosage text

Date:09/24/04ISR Number: 4458133-XReport Type:Expedited (15-DaCompany Report #IE-GLAXOSMITHKLINE-B0342939A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Consumer	Zyban	PS	Glaxosmithkline	ORAL
		Death		Alcohol	SS		

Date:09/24/04ISR Number: 4458139-0Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0345594A

Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Electrocardiogram Qrs	Consumer	Zyntabac	PS	Glaxosmithkline	ORAL
300MG Per day	25 DAY						
Initial or Prolonged		Complex Shortened		Vioxx	C		
25MG Per day							
Other		Tachycardia		Atrovent	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/24/04ISR Number: 4461698-5Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #CTU 227996

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG Q AM Initial or Prolonged ORAL		Convulsion		Wellbutrin Xl 150 Mg	PS		ORAL
				Prozac	C		
				Vistaril	C		
				Depakote Er	C		

Date:09/27/04ISR Number: 4459400-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0517054A  
 Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 2 YR		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
		Anxiety					
		Completed Suicide					
		Condition Aggravated					
		Dissociation					
		Gun Shot Wound					
		Head Injury					
		Hostility					
		Insomnia					
		Irritability					
		Weight Increased					

Date:09/27/04ISR Number: 4459407-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0526854A  
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lung Neoplasm		Wellbutrin	PS	Glaxosmithkline	
Other		Rash		Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day				Ethambutol	SS	Glaxosmithkline	
				Isoniazid	SS	Glaxosmithkline	
				Pyrazinamide	SS		
				Vitamin B6	C		

Date:09/27/04ISR Number: 4459408-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0526859A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Medication Error		Wellbutrin Sr	SS	Glaxosmithkline	

Date:09/27/04ISR Number: 4459409-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0526881A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Twice		Medication Error					
per day							

Date:09/27/04ISR Number: 4459413-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0527124A

Age: Gender:Male I/FU:I

Outcome	PT
Other	Accident At Work
	Amnesia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Convulsion Road Traffic Accident	Report Source	Product	Role	Manufacturer	Route
				Zyban No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:09/27/04ISR Number: 4459436-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0345641A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	1TAB Per day	5 DAY		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged		Dry Mouth Facial Spasm Rash Generalised		No Concurrent Medication	C		

Date:09/27/04ISR Number: 4462106-0Report Type:Direct  
Age:47 YR Gender:Female I/FU:I

Company Report #CTU 228068

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropriion Sr - Watson	PS	Watson	
Hospitalization - Initial or Prolonged		Depression Disease Recurrence Pharmaceutical Product Complaint					

Date:09/27/04ISR Number: 4462134-5Report Type:Direct  
Age:38 YR Gender:Male I/FU:I

Company Report #CTU 228090

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	450MG QD			Welbutrin Xl 150mg Glaxosmithkline	PS	Glaxosmithkline	ORAL
Other		Convulsion					

ORAL

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
RESPIRATORY		Adverse Event		Serevent	PS	Glaxosmithkline	
( INHALATION)		Dyspnoea					
		1 MON		Wellbutrin	SS	Glaxosmithkline	ORAL

Age: Gender:Male I/FU:F

Outcome	PT
Other	Acquired Porphyria
	Aggression
	Agitation
	Akathisia
	Allergy To Chemicals
	Amnesia
	Arthralgia
	Asthenia
	Bipolar Disorder
	Contrast Media Reaction
	Convulsion
	Coordination Abnormal

Freedom Of Information (FOI) Report

Dose	Duration		Report Source	Product	Role	Manufacturer	Route
42	DAY	Crying Depression Depression Suicidal Disability		Wellbutrin	PS	Glaxosmithkline	ORAL
		Dizziness		Tranquilizer	SS		
		Dysarthria		Depakote	SS		
		Dysphemia		Zyprexa	SS		
		Dyspnoea		Alcohol	C		
		Electric Shock					
		Encephalopathy					
		Facial Palsy					
		Fatigue					
		Feeling Abnormal					
		Head Banging					
		Headache					
		Hyperaesthesia					
		Hypersensitivity					
		Hypoxia					
		Immune System Disorder					
		Injury					
		Insomnia					
		Intentional Self-Injury					
		Mania					
		Mastocytosis					
		Metabolic Encephalopathy					
		Muscle Spasms					
		Myalgia					
		Myoclonus					
		Nervous System Disorder					
		Nightmare					
		Overdose					
		Pain Of Skin					
		Petit Mal Epilepsy					
		Suicidal Ideation					
		Suicide Attempt					
		Temperature Regulation Disorder					
		Throat Tightness					
		Tremor					
		Vestibular Disorder					
		Vomiting					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
3 YR		Euphoric Mood					
		Formication					
		Imprisonment					
		Loss Of Employment					
		Mania					
		Memory Impairment					
		Muscle Twitching					
		Personality Change					
		Restlessness					
		Suicidal Ideation					
		Tremor					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/28/04ISR Number: 4463377-7Report Type:Expedited (15-DaCompany Report #A0525788A  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anger Depression Feeling Abnormal Headache	Consumer	Wellbutrin Tablet-Zyban (Bupropion Hydrochloride)	PS		
1 TABLET/PER DAY		Homicidal Ideation Memory Impairment Oral Intake Reduced		Wellbutrin Xl Tablet-Extended Release (Bupropion Hydrochloride)	SS		ORAL
150 MG/PER DAY/ORAL							

Date:09/28/04ISR Number: 4463388-1Report Type:Expedited (15-DaCompany Report #A0524358A  
 Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness Loss Of Consciousness Syncope Vomiting	Consumer	Bupropion Hydrochloride Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ PER DAY/ ORAL				Atenolol	C		

Date:09/28/04ISR Number: 4463391-1Report Type:Expedited (15-DaCompany Report #A0525776A  
 Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Intentional Misuse Suicide Attempt	Consumer	Wellbutrin Xl Tablet-Extended Release (Bupropion			

300 MG/ ORAL

Hydrochloride)	PS	ORAL
Amitriptyline (Formulation Unknown) (Amitriptyline)	SS	

Date:09/28/04ISR Number: 4463553-3Report Type:Expedited (15-DaCompany Report #B0345474A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Angioneurotic Oedema Skin Reaction	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

ORAL

Date:09/28/04ISR Number: 4465181-2Report Type:Direct Company Report #CTU 228208

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - THREE TIMES A Initial or Prolonged DAY		Affect Lability Drug Ineffective		Wellbutrin Sr 150 Mg	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/29/04ISR Number: 4462243-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0410501A

Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anaemia		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Chest Pain					
150MG See dosage text		Dyspepsia		Gaviscon	C	Glaxosmithkline	
		Dysphagia		Black Cohosh	C	Glaxosmithkline	
		Gastric Disorder		Echinacea	C		
		Gastrooesophageal Reflux Disease		Aloe Vera	C		
		Nasopharyngeal Disorder		Tylenol Arthritis	C	Glaxosmithkline	
				Calcium	C		
				Aspirin	C	Glaxosmithkline	
				Amitriptyline	C		
				Ginger Root	C		
				Acidophilus	C		
				Hawthorn Berry	C		
				Kava-Kava	C		

Date:09/29/04ISR Number: 4462252-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0527512A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other		Anger					
5 MON		Depression					
		Eating Disorder					
		Feeling Hot					
		Hyperhidrosis					
		Insomnia					
		Nervousness					
		Panic Attack					
		Restlessness					
		Suicidal Ideation					

Date:09/29/04ISR Number: 4462259-4Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0336854A

Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Generalised Oedema		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	24 DAY	Nephrotic Syndrome					
Other 100MG Twice		Renal Failure		Nexen	SS	Glaxosmithkline	ORAL
per day	14 DAY	Weight Increased					

Date:09/29/04ISR Number: 4462261-2Report Type:Expedited (15-DaCompany Report #IE-GLAXOSMITHKLINE-B0342939A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion		Zyban Alcohol	PS SS	Glaxosmithkline	ORAL

Date:09/29/04ISR Number: 4464122-1Report Type:Direct Company Report #CTU 228288  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Effect Decreased Pharmaceutical Product		Bupropion Xl 150mg ?	PS		ORAL
1 BID ORAL		Complaint		Buproprion Xl	SS		ORAL
3 QAM ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/29/04ISR Number: 4465736-5Report Type:Direct  
Age:35 YR Gender:Female I/FU:I

Company Report #CTU 228328

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Withdrawal Syndrome		Wellbutrin 150 Mg			
		Migraine		-100mg	PS		ORAL

SEE IMAGE

Date:09/30/04ISR Number: 4463232-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506106A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective	Consumer	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Feeling Abnormal		Wellbutrin	SS	Glaxosmithkline	
150MG Per day							
		Libido Disorder		Paxil Cr	SS	Glaxosmithkline	
12.5MG See							
dosage text		Medication Error					
		Nausea		Xanax	SS		
		Nightmare		Lexapro	SS		
		Smoker		Seroquel	C		
				Advair	C	Glaxosmithkline	
				Albuterol	C	Glaxosmithkline	

Date:09/30/04ISR Number: 4463234-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0524499A  
Age:19 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Agitation	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	2 WK						
		Headache	Professional	No Concurrent			
		Psychiatric Symptom		Medication	C		
		Suicidal Ideation					

Date:09/30/04ISR Number: 4463242-5Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0338426A  
Age:34 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Agitation
Disability	Alanine Aminotransferase
Other	Increased
	Anorexia
	Anxiety
	Aspartate
	Aminotransferase
	Increased
	Asthenia
	Autoimmune Hepatitis
	Blood Alkaline
	Phosphatase Increased
	C-Reactive Protein
	Increased
	Fatigue
	Gamma-Glutamyltransferase
	Increased
	Liver Function Test
	Abnormal
	Malaise
	Myalgia
	Nausea
	Red Blood Cell
	Sedimentation Rate

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Increased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	36 DAY	Health Professional	Zyban	PS	Glaxosmithkline	ORAL
50MG per day			Zoloft	SS		ORAL
300MG Three times per day	4 DAY		Neurontin	C		ORAL
250MG per day	53 DAY		Seroquel	C		ORAL
2U per day			Prednisolone Slow Potassium	C	Glaxosmithkline	
40MG per day			Cipramil	C		
2U At night			Zolpidem	C		
2U At night			Caltrate	C	Glaxosmithkline	
25MG per day			Imuran	C	Glaxosmithkline	

Date:10/01/04ISR Number: 4464186-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0525619A  
 Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300MG Per day	2 MON	Completed Suicide	Wellbutrin	PS	Glaxosmithkline	ORAL
				Strattera	C		

Date:10/01/04ISR Number: 4464188-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0525720A  
 Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Convulsion Wellbutrin Xl PS Glaxosmithkline ORAL  
 300MG Per day 4 DAY  
 Grip Strength Decreased  
 Mouth Injury  
 Muscle Twitching  
 Tremor  
 Vomiting

Date:10/01/04ISR Number: 4464189-0Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0525978A  
 Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day		Vaginal Haemorrhage		Glimepiride + Rosiglitazone	SS	Glaxosmithkline	ORAL
176 DAY				Effexor Xr	SS		
225MG Per day				Vitamin C	C	Glaxosmithkline	

Date:10/01/04ISR Number: 4464194-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0527277A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Zyban	PS	Glaxosmithkline	
UNKNOWN	150MG Twice						
per day				No Concurrent Medication	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/01/04ISR Number: 4464195-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0527304A  
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Chest Pain	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4 WK						
Initial or Prolonged			Pain In Extremity				

Date:10/01/04ISR Number: 4464196-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0527318A  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Apathy	Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day	4 MON						
Initial or Prolonged			Drug Ineffective	Zoloft	SS		
			Overdose	Trileptal	SS		
			Pharmaceutical Product Complaint				
			Sleep Disorder				
			Stool Analysis Abnormal				
			Suicide Attempt				
			Thinking Abnormal				

Date:10/01/04ISR Number: 4464217-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0345688A  
Age:27 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening			Eye Rolling	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	9 DAY		Loss Of Consciousness				
			Masked Facies				
			Road Traffic Accident				
			Staring				

Date:10/01/04ISR Number: 4464218-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0345690A  
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day 10 DAY	Anxiety Disorder		Zyban	PS	Glaxosmithkline	ORAL
		Circulatory Collapse Disability Dizziness Vomiting					

Date:10/01/04ISR Number: 4464225-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0346164A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1TAB Per day 4 WK	Anxiety		Zyban	PS	Glaxosmithkline	ORAL
		Depression Disturbance In Attention Hallucination Personality Change Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/01/04ISR Number: 4464227-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0346382A

Age:26 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Fall		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	11 WK	Grand Mal Convulsion					

Date:10/01/04ISR Number: 4464231-7Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0346671A

Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Circulatory Collapse Dehydration Fatigue Feeling Abnormal Syncope		Zyban	PS	Glaxosmithkline	

Date:10/01/04ISR Number: 4464241-XReport Type:Expedited (15-DaCompany Report #US-JNJFOC-20040908652

Age:32 YR Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death OROPHARINGEAL		Cardio-Respiratory Arrest		Topamax	PS		
OROPHARINGEAL		Completed Suicide		Bupropion	SS		
OROPHARINGEAL		Overdose		Venlafaxine	SS		

Date:10/01/04ISR Number: 4464243-3Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20040908721

Age:20 YR Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death OROPHARINGEAL		Completed Suicide		Topamax	PS		
OROPHARINGEAL		Intentional Misuse		Valproic Acid	SS		

Date:10/04/04ISR Number: 4465049-1Report Type:Expedited (15-DaCompany Report #PHBS2004US12809  
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Pressure Decreased Cardiac Failure		Metoprolol	PS	Novartis Sector: Pharma	
75 mg, BID		Congestive		Bupropion	SS		
150 mg, BID		Drug Interaction		Diltiazem	C		
240 mg, BID		Dyspnoea Fatigue Heart Rate Decreased Pitting Oedema Pulmonary Hilum Mass Pulmonary Vascular Disorder Respiratory Rate Increased Sinus Bradycardia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/04/04ISR Number: 4465403-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511243A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG See	Completed Suicide	Other	Wellbutrin Sr	PS	Glaxosmithkline	
dosage text							
Decreased Appetite							
Depressed Mood							
Disturbance In Attention							
Feelings Of Worthlessness							
Gun Shot Wound							
Major Depression							
Sleep Disorder							
Social Avoidant Behaviour							
Suicidal Ideation							
Treatment Noncompliance							
Tremor							

Date:10/04/04ISR Number: 4465417-8Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0528067A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1800MG Single	Overdose		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
dose							
Paraesthesia							

Date:10/04/04ISR Number: 4465426-9Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0343398A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Twice	Death	Health	Zyban	PS	Glaxosmithkline	ORAL
per day							
Multi-Organ Failure							
Professional							
Thyroxine							
Hormone Replacement							
Therapy							
Amlodipine							

Date:10/04/04ISR Number: 4465442-7Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0346880A

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150MG Twice	Myocardial Infarction		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	Stent Occlusion					

Date:10/04/04ISR Number: 4467068-8Report Type:Direct Company Report #CTU 228724

Age:29 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Grand Mal Convulsion		Wellbutrin Xl 300 Mg			
300 MG PO	Medication Error		Glaxo Smithkline	PS	Glaxo Smithkline	ORAL

Date:10/05/04ISR Number: 4466311-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0520112A

Age:40 YR Gender:Female I/FU:F

Outcome  
Life-Threatening  
Hospitalization -  
Initial or Prolonged  
Disability

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
364 DAY		Amnesia	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
		Arthralgia	Professional	Prednisone	C		ORAL
		Arthropathy		Nexium	C		ORAL
		Aura		Remicade	C		
INTRAVENOUS		Brachial Plexus Injury		Klonopin	C		ORAL
RESPIRATORY		Electroencephalogram		Advair	C	Glaxosmithkline	
(INHALATION)		Abnormal					
15MG Weekly		Grand Mal Convulsion		Methotrexate	C		ORAL
SUBCUTANEOUS	80MG Twice	Joint Dislocation		Lovenox	C		
per day	28 DAY	Neck Pain					
INTRAVENOUS		Nuclear Magnetic Resonance Imaging Brain		Multivitamin	C		
				Synvisc	C		
INTRAVENOUS		Abnormal		Synvisc	C		
INTRAVENOUS		Ovarian Cyst		Iron	C		
INTRAVENOUS		Paraesthesia		Flu Vaccine	C	Glaxosmithkline	
		Retinopathy		Lovenox	C		
				Coumadin	C	Glaxosmithkline	
				Fragmin	C		
12 DAY				Bipap	C		

Date:10/05/04ISR Number: 4466314-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0527585A

Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day 2 MON	Intentional Misuse		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged Disability		Vomiting		Simvastatin	C		

Date:10/05/04ISR Number: 4466319-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0528132A  
Age:39 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day 6 MON						
Initial or Prolonged	Fall		Ritalin	C		
	Laceration		Lithium	C	Glaxosmithkline	

Date:10/05/04ISR Number: 4466320-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0528136A  
Age:46 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Depression		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Per day 18 MON						
Initial or Prolonged	Difficulty In Walking					
	Eating Disorder					
	Headache					
	Speech Disorder					
	Suicidal Ideation					

Date:10/05/04ISR Number: 4466324-7Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0528437A  
Age: Gender: I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Blood Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/06/04ISR Number: 4467295-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0345690A  
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day	10 DAY	Anxiety Disorder	Zyban	PS	Glaxosmithkline	ORAL
			Circulatory Collapse				
			Disability				
			Dizziness				
			Vomiting				

Date:10/06/04ISR Number: 4467305-XReport Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0346847A  
 Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN			Extrapyramidal Disorder	Zyban	PS	Glaxosmithkline	
Initial or Prolonged			Overdose				
			Suicide Attempt				

Date:10/06/04ISR Number: 4467311-5Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0347143A  
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day	1 MON	Abortion Induced	Zyban	PS	Glaxosmithkline	ORAL
			Drug Exposure During Pregnancy				
			Drug Ineffective				

Date:10/06/04ISR Number: 4467312-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0347143B  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	1 MON	Chromosome Abnormality	Zyban	PS	Glaxosmithkline	
Congenital Anomaly			Congenital Hand				

Malformation  
Diaphragmatic Hernia  
Foetal Growth Retardation  
Foetal Malformation  
Gastrointestinal Disorder

Date:10/06/04ISR Number: 4467313-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0347195A  
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Twice per day	34 DAY	Cerebral Ischaemia Completed Suicide Injury Asphyxiation		Zyban Novonorm Actos Mediator Aspegic	PS C C C C	Glaxosmithkline	ORAL

Date:10/06/04ISR Number: 4470677-3Report Type:Direct Company Report #CTU 228999  
Age:47 YR Gender:Male I/FU:I

Outcome	PT
Other	Feeling Abnormal Headache Heart Rate Increased

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Other	Renal Failure	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
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Date:10/07/04ISR Number: 4468373-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0528565A  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Nephrolithiasis		Wellbutrin	PS	Glaxosmithkline	ORAL
YR	Renal Neoplasm					

Date:10/07/04ISR Number: 4468374-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0528776A  
Age:65 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
225MG Per day	Major Depression					
	Polyuria					

Date:10/07/04ISR Number: 4468376-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0528781A  
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Pelvic Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	Urinary Incontinence		Hytrin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/07/04ISR Number: 4468392-5Report Type:Expedited (15-DaCompany Report #2004016942  
Age:20 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150MG per day	Skin Discolouration		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged 100MG per day			Lamictal	SS	Glaxosmithkline	ORAL

Date:10/07/04ISR Number: 4473175-6Report Type:Direct Company Report #USP 56898  
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
TABLET, EXTENDED RELEASE	Medication Error		Wellbutrin	PS	Eon	

Date:10/08/04ISR Number: 4469806-7Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0528431A  
Age:45 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other 55 DAY	Convulsion		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
	Loss Of Consciousness		Sedative	C		ORAL

Date:10/08/04ISR Number: 4469811-0Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0344204A  
Age:37 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 9 DAY	Adverse Event	Health	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged	Aggression	Professional	Valium	SS		
	Agitation		Warfarin	SS	Glaxosmithkline	
	Amnesia		Tritace	SS		

Hostility  
Irritability  
Mental Disorder  
Overdose  
Self-Medication  
Suicide Attempt

Panadeine Forte SS Glaxosmithkline

Date:10/08/04ISR Number: 4469815-8Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0345688A  
Age:27 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Twice per day	9 DAY	Eye Rolling Loss Of Consciousness Road Traffic Accident Staring		Zyban	PS	Glaxosmithkline	ORAL

Date:10/08/04ISR Number: 4469818-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0346989A  
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG See dosage text	15 DAY	Swelling Swelling Face Urticaria Wheezing		Zyban	PS	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/08/04ISR Number: 4469823-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0347372A  
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diplopia		Zyban	PS	Glaxosmithkline	
150MG Unknown							
		Iiird Nerve Paralysis					
		Nystagmus					

Date:10/08/04ISR Number: 4471120-0Report Type:Expedited (15-DaCompany Report #2004S1002873  
Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Intentional Misuse	Consumer	Wellbutrin Xl Tablet			
Initial or Prolonged		Suicide Attempt	Other	Extended Release 300			
Other				Mg	PS	Glaxosmithkline	ORAL
300MG, ORAL							

Date:10/11/04ISR Number: 4471020-6Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0525978A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Health	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day							
		Vaginal Haemorrhage	Professional	Glimepiride + Rosiglitazone	SS	Glaxosmithkline	ORAL
177 DAY							
				Effexor Xr	SS		ORAL
225MG Per day							
				Vitamin C	C	Glaxosmithkline	

Date:10/11/04ISR Number: 4471022-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0528860A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Bupropion	PS	Glaxosmithkline	
200MG per day							

Date:10/11/04ISR Number: 4471046-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0346993A  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Chromaturia		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	1 MON	Pruritus					

Date:10/11/04ISR Number: 4471053-XReport Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0347371A  
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	
UNKNOWN		2 DAY					
		Crying					
		Delusion					
		Dry Mouth					

Date:10/12/04ISR Number: 4471574-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0529360A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fracture		Wellbutrin	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/12/04ISR Number: 4471600-8Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0347567A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Respiratory Disorder		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day		Urticaria					
Other							

Date:10/12/04ISR Number: 4472878-7Report Type:Direct Company Report #CTU 229280

Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Prozac	PS		
		Drug Ineffective		Wellbutrin	SS		
		Nausea					

Date:10/13/04ISR Number: 4472803-9Report Type:Expedited (15-DaCompany Report #200416846US

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Creatinine Increased	Consumer	Ketek	PS	Aventis Pharmaceuticals Inc.	ORAL
		Blood Urea Increased		Wellbutrin	SS		
dose: NOT PROVIDED		Hallucination, Visual					
dose: NOT PROVIDED		Hyperkinesia		Metoprolol	C		
dose: NOT PROVIDED		Restlessness					
				Ssri	C		
dose: NOT PROVIDED				Baby Aspirin	C		
dose: NOT PROVIDED							

dose: NOT PROVIDED	Allopurinol	C
dose: NOT PROVIDED	Hydrochlorothiazide	C
dose: NOT PROVIDED	Lasix	C
dose: NOT PROVIDED	Potassium Chloride	C
dose: NOT PROVIDED	Voltaren	C
dose: NOT PROVIDED	Aceon	C
dose: NOT PROVIDED	Hydrocodone	C
dose: NOT PROVIDED	Calcium	C
dose: NOT PROVIDED	Multivitamins, Plain	C
dose: 6-7	Coumadine	C
dose: NOT PROVIDED	Detrol	C

Date:10/13/04ISR Number: 4473037-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0521769A  
Age:61 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	18 DAY					
	Muscle Tightness					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/13/04ISR Number: 4473044-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0529354A

Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated		Wellbutrin Sr	PS	Glaxosmithkline	
150MG Twice							
		Glaucoma					
per day	YR			Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day				Zoloft	C		

Date:10/13/04ISR Number: 4473049-0Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0343107A

Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Abasia		Bupropion			
Initial or Prolonged		Arthralgia		Hydrochloride	PS	Glaxosmithkline	
150MG Unknown	2 DAY						
		Erythema Multiforme		Prednisolone	C	Glaxosmithkline	
30MG Per day							
		Pain					
		Type Iv Hypersensitivity					
		Reaction					

Date:10/13/04ISR Number: 4476558-3Report Type:Expedited (15-DaCompany Report #A0499051A

Age:6 MON Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Exposure Via Breast	Literature	Wellbutrin Sr			
		Milk	Health	Tablet-Controlled			
		Febrile Convulsion	Professional	Release (Bupropion			
		Grand Mal Convulsion		Hydrochloride)	PS		
TRANSMAMMARY	150 MG/PER						
		Postictal State					
DAY/							
		Restlessness					
TRANSMAMMARY							
		Upper Respiratory Tract					
		Infection					

Date:10/14/04ISR Number: 4474467-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0526711A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -		Intentional Misuse		Xanax	SS		
Initial or Prolonged		Overdose					
		Refusal Of Treatment By					
		Patient					

Date:10/14/04ISR Number: 4474479-3Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0345610A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Bradyphrenia	Health	Zyban	PS	Glaxosmithkline	ORAL
		Dizziness	Professional	No Concurrent			
		Hypotension		Medication	C		
		Syncope					
		Vision Blurred					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4475761-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0522931A

Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Glucose Increased	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	DAY	Clonic Convulsion	Professional				
		Fall		Ludiomil	C		
		Joint Dislocation		Symbyax	C		
		Loss Of Consciousness					
		Medication Error					
		Upper Limb Fracture					

Date:10/15/04ISR Number: 4475776-8Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0529972A

Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anxiety		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
Initial or Prolonged		Hypersensitivity					
per day							
		Oropharyngeal Swelling					
		Rash Generalised					
		Swollen Tongue					
		Tremor					

Date:10/15/04ISR Number: 4475789-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0344204A

Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Adverse Event	Health	Zyban	PS	Glaxosmithkline	ORAL
1TAB Twice							
Initial or Prolonged		Aggression	Professional				
per day	9 DAY						
		Agitation		Valium	SS		
		Amnesia		Warfarin	SS	Glaxosmithkline	
		Hostility		Tritace	SS		
		Irritability		Panadeine Forte	SS	Glaxosmithkline	
		Mental Disorder		Paracetamol	C	Glaxosmithkline	

Overdose  
Suicide Attempt

Date:10/15/04ISR Number: 4475802-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0347674A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Anorexia		Zyban	PS	Glaxosmithkline	ORAL
29 DAY		Dizziness		Edronax	C		
		Feeling Abnormal		Aropax	C	Glaxosmithkline	
		Mania					
		Nausea					
		Suicidal Ideation					

Date:10/15/04ISR Number: 4479229-2Report Type:Expedited (15-DaCompany Report #2004236066US  
Age:50 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Literature	Xanax (Alprazolam)			
		Intentional Misuse	Health	Tablet	PS		
			Professional	Bupropion			
				(Amfebutamone)	SS		
				Propoxyphene Hcl And			
				Acetaminophen			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Dextropropoxyphene  
Hydrochloride) SS

Date:10/15/04ISR Number: 4479315-7Report Type:Expedited (15-DaCompany Report #2004236050US  
Age:44 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse	Literature Health	Solfidin (Clonazepam)Tablet	PS		ORAL
ORAL			Professional	Xanax (Alprazolam) Tablet	SS		ORAL
ORAL				Bupropion (Amfebutamone)	SS		ORAL

Date:10/15/04ISR Number: 4479343-1Report Type:Expedited (15-DaCompany Report #A0527924A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage SEE DOSAGE		Abdominal Pain Fall Grand Mal Convulsion	Study Health Professional Other	Wellbutrin Xl Tablet-Extended Release (Bupropion Hydrochloride)	PS		ORAL
TEXT/ORAL							

Date:10/15/04ISR Number: 4507722-2Report Type:Periodic Company Report #2003116563  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 2400 MG Initial or Prolonged (TID), ORAL Other		Dystonia Eye Disorder Intentional Misuse Medication Error	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
				Bupropion			



Muscle Twitching  
Suicide Attempt

Hydrochloride  
(Bupropion  
Hydrochloride) SS  
Baclofen (Baclofen) C  
Vicodin  
(Paracetamol,  
Hydrocodone  
Bitartrate) C  
Zolpidem Tartrate  
(Zolpidem Tartrate) C  
Clonazepam  
(Clonazepam) C  
Fluoxetine  
Hydrochloride  
(Fluoxetine  
Hydrochloride) C  
Quetiapine Fumarate  
(Quetiapine  
Fumarate) C

Date:10/18/04ISR Number: 4477486-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0520111A  
Age:26 YR Gender:Female I/FU:F

Outcome PT  
Life-Threatening Arthralgia  
Complex Partial Seizures

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Confusional State Fatigue Headache	Report Source	Product	Role	Manufacturer	Route
300MG Per day	MON	Malaise	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
			Professional	Risperdal	C		
				Prozac	C		

Date:10/18/04ISR Number: 4477487-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0521279A  
Age:6 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Hallucination	Professional				
37.5MG Per day	1 WK	Medication Error		Strattera	C		
25MG per day				Clonidine	C		
.1MG At night							

Date:10/18/04ISR Number: 4477499-8Report Type:Expedited (15-DaCompany Report #200416846US  
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	2 WK	Cellulitis		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged	1 WK	Condition Aggravated		Telithromycin	SS		ORAL
800MG Per day		Dyskinesia		Antibiotic (Unspecified)	SS		ORAL
		Hallucination, Visual		Metoprolol	C		
		Parasomnia		Lexapro	C		
		Pneumonia		Baby Aspirin	C	Glaxosmithkline	
		Pyrexia		Allopurinol	C	Glaxosmithkline	
		Restlessness		Hydrochlorothiazide	C		
				Lasix	C	Glaxosmithkline	
				Potassium Chloride	C	Glaxosmithkline	
				Voltaren	C	Glaxosmithkline	
				Aceon	C		

Hydrocodone	C	
Multivitamin	C	
Coumadin	C	Glaxosmithkline
Detrol	C	
Uroxatral	C	
Calcium	C	

Date:10/18/04ISR Number: 4477524-4Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0347371A

Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	
UNKNOWN		2 DAY					
		Crying					
		Delusion					
		Dry Mouth					

Date:10/18/04ISR Number: 4477530-XReport Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0044946A

Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Intentional Misuse		Zyban	PS	Glaxosmithkline	ORAL
10TAB Single							
dose		No Adverse Effect					
		Suicide Attempt					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/18/04ISR Number: 4478471-4Report Type:Direct  
Age:26 YR Gender:Female I/FU:I

Company Report #CTU 229799

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Abnormal Behaviour		Wellbutrin Sr	PS		
1X MORNING						
Hospitalization -	Drug Ineffective		Lorazepam	SS		
1 X NIGHT						
Initial or Prolonged	Intentional Self-Injury					
Required	Overdose					
Intervention to	Suicidal Ideation					
Prevent Permanent	Suicide Attempt					
Impairment/Damage						

Date:10/18/04ISR Number: 4478516-1Report Type:Direct  
Age:57 YR Gender:Female I/FU:I

Company Report #CTU 229884

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
ONE PO BID	Drug Intolerance		Wellbutrin Sr 100 Mg	PS		ORAL
	Gastrointestinal Disorder					
	Hypersensitivity					
	Pharmaceutical Product					
	Complaint					

Date:10/19/04ISR Number: 4478447-7Report Type:Expedited (15-DaCompany Report #PHEH2004US10928  
Age:41 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Apathy		Trileptal	PS	Novartis Sector:	
Initial or Prolonged	Intentional Misuse				Pharma	
	Sleep Disorder		Zoloft	SS		
	Stool Analysis Abnormal		Wellbutrin - Slow			
	Suicide Attempt		Release	SS		ORAL
450 mg, QD	Thinking Abnormal					

Date:10/19/04ISR Number: 4479180-8Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 229912

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Pharmaceutical Product Complaint		Wellbutrin Sr (Generic)	PS		ORAL
150 MG 2		Suicidal Ideation					
TWICE BID PO							

Date:10/19/04ISR Number: 4479363-7Report Type:Direct  
Age:41 YR Gender:Female I/FU:I

Company Report #CTU 229932

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other QD X 3 D ,		Hypomania		Zyban 150 Mg Cr Tab	PS		
THEN BID		Insomnia					
		Ovarian Cyst Pelvic Pain		Accupril (Quinapril) Zoloft (Sertraline Hcl) Flovent (Gluticasone Propionate ) Inhal Albuterol Midrin (Apap-Isometheptene- Dichloral) Yasmin (Drospirenone-Ethiny	C C C C C		

Freedom Of Information (FOI) Report

1 Estradiol) C  
 Atarax (Hydroxyzine Hcl) C  
 Allegra (Fexofenadine Hcl) C  
 Flonase(Fluticasone Propionate (Nasal)) C  
 Advair Diskus (Salmeterol-Fluticasone) C  
 Lorcet-Hd (Hydrocodone-Acetaminophen) C

Date:10/19/04ISR Number: 4480633-7Report Type:Expedited (15-DaCompany Report #2004076041  
 Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	50 MG (50 MG, 1 IN 1 D),	Condition Aggravated	Health	Zoloft (Sertraline)	PS		ORAL
ORAL		Cystitis	Professional				
		Feeling Jittery					
		Urinary Retention		Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		ORAL
450 MG, ORAL				Clonazepam (Clonazepam)	SS		

Date:10/20/04ISR Number: 4479673-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0530207A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	UNKNOWN	Convulsion		Wellbutrin	PS	Glaxosmithkline	
	600MG	See					
		Overdose					

dosage text

Date:10/20/04ISR Number: 4490081-1Report Type:Expedited (15-DaCompany Report #B0347331A  
Age:56 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Cardiac Failure Congestive Drug Interaction	Literature Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
150 MG TWICE  PER DAY ORAL	Dyspnoea					
   75 MG TWICE  PER DAY	Fatigue Nodal Rhythm Respiratory Rate Increased  Sinus Bradycardia		Metoprolol (Formulation Unknown) (Metoprolol)	SS		
	Ventricular Hypertrophy		Diltiazem Hydrochloride	C		

Date:10/21/04ISR Number: 4481334-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0530398A  
Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Decreased Appetite Headache Medication Error

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Tremor Vomiting	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	4 DAY			Wellbutrin	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:10/21/04ISR Number: 4483586-0Report Type:Expedited (15-DaCompany Report #2004AL000696  
Age:15 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature	Children'S Apap Elixir (Acetaminophen) (Alpharma)	PS	Alpharma	ORAL
PO				Infant'S Apap Drops (Acetaminophen) (Alpharma)	SS	Alpharma	ORAL
PO				Bupropion	SS		ORAL
PO				Aspirin	SS		ORAL

Date:10/21/04ISR Number: 4483683-XReport Type:Expedited (15-DaCompany Report #2004AL000923  
Age:52 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose	Literature	Children'S Apap Elixir (Acetaminophen) (Alpharma)	PS	Alpharma	ORAL
PO				Infants' Apap Drops (Acetaminophen)			



PO	(Alpharma)	SS	Alpharma	ORAL
PO	Aspirin	SS		ORAL
PO	Bupropion	SS		ORAL

Date:10/22/04ISR Number: 4482394-4Report Type:Periodic Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-12542486  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
dose		Paraesthesia Swollen Tongue		Buspar	PS	Bristol-Myers Squibb Company	ORAL
increased to							
20mg/day							
,then							
30mg/day, and							
1	DAY			Keflex Wellbutrin	SS SS		ORAL ORAL

Date:10/22/04ISR Number: 4482480-9Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0348835A  
 Age: Gender: I/FU:F

Outcome	PT
Disability	Adverse Event Disability Feeling Abnormal Hypersensitivity

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Impaired Work Ability Malaise	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Zyntabac	PS	Glaxosmithkline	
Date:10/22/04		ISR Number: 4483878-5	Report Type:Direct	Company Report #CTU 230348			
Age:41 YR	Gender:Male	I/FU:I					
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Bupropion	PS		ORAL
100MG Q DAY		Pharmaceutical Product					
PO		Complaint					
Date:10/22/04		ISR Number: 4483903-1	Report Type:Direct	Company Report #CTU 230246			
Age:	Gender:Female	I/FU:I					
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Intentional Self-Injury		Wellbutrin Sr			
Life-Threatening		Pharmaceutical Product		(Generic)	PS		ORAL
150 MG 2 BID		Complaint					
PO		Suicidal Ideation					
Date:10/22/04		ISR Number: 4484569-7	Report Type:Expedited (15-Da	Company Report #2004AL000695			
Age:44 YR	Gender:	I/FU:I					
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Multiple Drug Overdose	Literature	Clonazepam	PS	Purepac	ORAL
Death				Alprazolam	SS	Purepac	ORAL
PO				Bupropion (Long			
				Acting)	SS		ORAL
PO							

Date:10/22/04ISR Number: 4484596-XReport Type:Expedited (15-DaCompany Report #2004AL000698

Age:43 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Multiple Drug Overdose	Literature	Temazepam Capsules, 30 Mg (Purepac)	PS	Purepac	ORAL
PO				Bupropion (Long Acting)	SS		ORAL

Date:10/22/04ISR Number: 4484738-6Report Type:Expedited (15-DaCompany Report #2004AL000680

Age:50 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Multiple Drug Overdose	Literature	Alprazolam Tablets Usp, 0.25mg,0.5mg,1mg & 2mg (Purepac)	PS		ORAL
PO				Propoxyphene Napsylate And Acetaminophen Tablets Usp, 100mg/650 Mg	SS		ORAL
PO				Bupropion	SS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/22/04ISR Number: 4485435-3Report Type:Expedited (15-DaCompany Report #2004AL000694  
Age:52 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Completed Suicide	Literature	Diazepam Tablets Usp, 10 Mg (Purepac)	PS	Purepac	ORAL
PO		Intentional Misuse		Bupropion	SS		ORAL
PO				Quetiapine	SS		ORAL

Date:10/25/04ISR Number: 4484225-5Report Type:Expedited (15-DaCompany Report #2004-03238  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day 10 YR	Asthenia	Consumer	Wellbutrin Sr	PS	Glaxosmithkline	
	150MG Twice per day 10 YR	Coordination Abnormal Depressed Level Of Consciousness Disturbance In Attention		Bupropion Hydrochloride	SS	Glaxosmithkline	ORAL
		Motor Dysfunction Pharmaceutical Product Complaint Tremor		Lithium	C	Glaxosmithkline	ORAL

Date:10/25/04ISR Number: 4484229-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLIN-A0530836A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day	Cerebrovascular Accident		Wellbutrin	PS	Glaxosmithkline	

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day 1	MON	Abortion Induced	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged	UNKNOWN	UNKNOWN	Drug Exposure During	Meteospasmyl	C		
UNKNOWN			Pregnancy	Primperan	C	Glaxosmithkline	
UNKNOWN			Drug Ineffective	Vitamin C	C	Glaxosmithkline	
UNKNOWN				Ginkor Fort	C		
UNKNOWN				Stresam	C		ORAL
3UNIT per day							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day 1	MON	Chromosome Abnormality	Zyban	PS	Glaxosmithkline	
Congenital Anomaly			Congenital Hand Malformation	Meteospasmyl	C		
			Dextrocardia	Primperan	C	Glaxosmithkline	
			Diaphragmatic Hernia	Vitamin C	C	Glaxosmithkline	
			Drug Exposure During	Ginkor Fort	C		
				Stresam	C		
3UNIT per day							
			Pregnancy				
			Foetal Growth Retardation				
			Foetal Malformation				
			Gastrointestinal Disorder				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/25/04ISR Number: 4484260-7Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0348835A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Adverse Event Disability Hypersensitivity Malaise		Zyntabac	PS	Glaxosmithkline	

Date:10/25/04ISR Number: 4487637-9Report Type:Expedited (15-DaCompany Report #US010405

Age:17 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Aura Coma Complex Partial Seizures Grand Mal Convulsion Parosmia Road Traffic Accident Status Epilepticus Treatment Noncompliance	Health Professional	Gabitril Wellbutrin	PS SS		

Date:10/26/04ISR Number: 4485727-8Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0349044A

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Insomnia Sleep Disorder		Zyban	PS	Glaxosmithkline	ORAL

Date:10/26/04ISR Number: 4485728-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0349140A

Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	15 DAY	Agitation Confusional State		Zyban	PS	Glaxosmithkline	ORAL

25MG Per day	15	DAY	Incoherent	Atarax	SS	ORAL
UNKNOWN			Insomnia	Alprazolam	C	
UNKNOWN				Equanil	C	
UNKNOWN				Stilnox	C	

Date:10/26/04ISR Number: 4488275-4Report Type:Expedited (15-DaCompany Report #S04-USA-04732-01  
Age:61 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour	Health	Lexapro			
		Agitation	Professional	(Escitalopram)	PS		ORAL
10 MG QD PO							
		Belligerence		Lexapro			
		Confusional State		(Escitalopram)	SS		ORAL
20 MG QD PO							
		Delusion		Wellbutrin			
		Diet Refusal		(Bupropion			
		Insomnia		Hydrochloride)	SS		ORAL
300 MG QD							
		Mania		Wellbutrin			
		Paranoia		(Bupropion			
				Hydrochloride)	SS		ORAL
150 MG QD							
				Ambien (Zolpidem			
				Tartrate)	C		
				Trazodone	C		
				Lipitor			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Atorvastatin) C  
 Aciphex (Rabeprazole Sodium) C  
 Aspirin C

Date:10/27/04ISR Number: 4487069-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0530790A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300MG Per day	4 WK	Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
Other			Death	Abilify	C		

Date:10/27/04ISR Number: 4487070-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0530819A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	1050MG Per day	0 DAY	Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
Other			Overdose				
Hospitalization - Initial or Prolonged Disability							

Date:10/27/04ISR Number: 4487076-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0531192A  
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	5 WK		Constipation	Wellbutrin	PS	Glaxosmithkline	ORAL
			Convulsion	Trileptal	C		
			Dizziness	Mysoline	C		
			Insomnia	Synthroid	C	Glaxosmithkline	
			Micturition Disorder	Premarin	C		



Date:10/27/04ISR Number: 4487088-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0334395A  
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged UNKNOWN Disability		Choking Confusional State Convulsion Headache Therapy Non-Responder Tongue Biting		Zyban Desloratadine	PS C	Glaxosmithkline	ORAL

Date:10/27/04ISR Number: 4487843-3Report Type:Direct Company Report #CTU 230570  
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150 MG BID		Convulsion		Wellbutrin 150 Mg (Bupropion Sr)	PS		

Date:10/28/04ISR Number: 4488047-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0530912A  
Age:22 YR Gender:Female I/FU:F

Outcome	PT
Other	Convulsion Medication Error

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Overdose

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
2	DAY		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:10/28/04ISR Number: 4488052-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0531408A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	4	DAY	Convulsion	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Impaired Driving Ability				

Date:10/28/04ISR Number: 4488072-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0346671A  
Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice			Dehydration	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	29	DAY	Fatigue	Professional			
			Feeling Abnormal Syncope				

Date:10/28/04ISR Number: 4488093-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0349464A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Anaphylactic Shock	Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown			Drug Interaction	Anesthesia	SS		

Date:10/28/04ISR Number: 4488527-8Report Type:Direct Company Report #CTU 230662  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Pharmaceutical Product		Budeprion Sr 150 Mg Teva-?-	PS	Teva	
150 MG TWICE		Complaint					
DAILY ORAL							

Date:10/28/04ISR Number: 4490501-2Report Type:Expedited (15-DaCompany Report #2004-BP-09934BP  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Limb Injury Oedema Peripheral	Consumer	Spiriva Handihaler (Tiotropium Bromide)	PS		
Hospitalization - Initial or Prolonged RESPIRATORY		Platelet Count Decreased					
(INHALATION)	18 MCG (18						
MCG), IH				Bextra (Valdecoxib)	SS		ORAL
PO				Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
PO				Foradil (Formoterol Fumarate)	SS		
RESPIRATORY							
(INHALATION)	IH			Prednisone	C		
				Fosamax (Alendronate Sodium)	C		
				Foltz (Folic Acid)	C		
				Glucosamine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Duoneb (Combivent) C

Date:10/29/04ISR Number: 4490523-1Report Type:Expedited (15-DaCompany Report #2004UW21313

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 10 MG QD PO		Convulsion	Foreign	Crestor	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage			Health Professional Other	Wellbutrin - Slow Release	SS		

Date:10/29/04ISR Number: 4491919-4Report Type:Direct

Company Report #CTU 230810

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Disability 20 MG 2 TIMES		Disorientation Lacunar Infarction		Methylphenidate 20 Mg	PS		ORAL
PER DA ORAL		Speech Disorder		Wellbutrin 150 Mg	SS		ORAL
150 MG 2 TIMES PER DA ORAL				Celexa Trazadone Luvox	C C C		

Date:11/01/04ISR Number: 4489942-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0531529A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 200MG Per day		Blood Amylase Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
				No Concurrent			

Date:11/01/04ISR Number: 4489968-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0349166A  
 Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Angioneurotic Oedema	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice			Head Discomfort				
per day	8	DAY	Swollen Tongue	Nexium	C		ORAL
20MG per day				Doxazosin	C		ORAL
4MG per day				Amlodipine	C		ORAL
10MG per day				Bendroflumethiazide	C	Glaxosmithkline	
UNKNOWN				Becloforte	C	Glaxosmithkline	
				Ventolin	C	Glaxosmithkline	

Date:11/02/04ISR Number: 4490664-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0521834A  
 Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Amnesia	Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day			Confusional State	Geodon	C		
60MG Twice			Grand Mal Convulsion				
per day			Incontinence	Temazepam	C		
30MG At night	2	DAY	Thinking Abnormal				
			Tongue Biting				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/02/04ISR Number: 4490666-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0524516A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	1 MON	Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Alcohol Intolerance Amnesia Convulsion Dissociation Drug Ineffective Overdose Suicidal Ideation Vision Blurred		Alcohol	C		

Date:11/02/04ISR Number: 4490670-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0528776A  
Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	225MG Per day	Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
15MG Single		Benign Prostatic Hyperplasia Major Depression		Zoloft Clonazepam Remeron	SS SS SS		
dose		Polyuria Urinary Retention					

Date:11/02/04ISR Number: 4490675-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0531761A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	600MG Unknown	Convulsion Overdose		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/02/04ISR Number: 4490676-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0531793A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
15 YR		Murder		Wellbutrin	SS	Glaxosmithkline	
150MG Per day		Therapeutic Response		Lopressor	C		
UNKNOWN		Unexpected		Prilosec	C		
				Premarin	C		
UNKNOWN	.3MG	Unknown					

Date:11/02/04ISR Number: 4490678-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0531856A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Abuser		Seizure Medication	C		

Date:11/02/04ISR Number: 4490683-2Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0347567A

Age:18 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Angioneurotic Oedema
Initial or Prolonged	Dysphagia
	Dyspnoea
	Oedema Peripheral

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Pain Respiratory Disorder Swelling Face					
per day	18 DAY	Urticaria		Zyban	PS	Glaxosmithkline	ORAL
		White Blood Cell Count Abnormal		Harmonet	C		ORAL

Date:11/02/04ISR Number: 4490688-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0349294A  
Age:56 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Unknown		Subarachnoid Haemorrhage		Zyban	PS	Glaxosmithkline	ORAL
	75MG Twice				Venlafaxine	C		ORAL
	per day							

Date:11/02/04ISR Number: 4493415-7Report Type:Expedited (15-DaCompany Report #A0531315A  
Age:29 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Intentional Misuse	Health Professional	Wellbutrin Xl Tablet-Extended Release (Bupropion Hydrochloride)	PS		ORAL
Life-Threatening Disability Required	300 MG / PER	Intervention to DAY / ORAL			Ethanol (Alcohol)	SS		
Prevent Permanent Impairment/Damage					Citalopram Hydrobromide	C		

Date:11/02/04ISR Number: 4493416-9Report Type:Expedited (15-DaCompany Report #A0531537A  
Age:28 YR Gender:Male I/FU:I



Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Required Intervention to Prevent Permanent Impairment/Damage			Drug Ineffective	Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride) Wellbutrin Xl Tablet-Extended Release (Bupropion Hydrochloride)	PS		
ORAL		4	MON		Celecoxib Mirtazapine	C C		ORAL

Date:11/02/04ISR Number: 4493421-2Report Type:Expedited (15-DaCompany Report #A0531464A  
Age:31 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Convulsion	Consumer	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL		2	YR					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/02/04ISR Number: 4493434-0Report Type:Expedited (15-DaCompany Report #A0531546A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion Gastritis Haematemesis Medication Error Migraine Pain Rectal Haemorrhage Vomiting	Consumer	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride) Wellbutrin Xl Tablet-Extended Release (Bupripion Hydrochloride)	PS		ORAL
ORAL		Weight Decreased		Lorazepam	C		

Date:11/02/04ISR Number: 4494274-9Report Type:Expedited (15-DaCompany Report #B0349472A  
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other  150 MG/ORAL		Aspartate Aminotransferase Increased  Blood Creatinine Increased Blood Urea Increased Rhabdomyolysis	Foreign Literature Health  Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

Date:11/03/04ISR Number: 4492143-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0524069A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG per day		Suicidal Ideation		Wellbutrin  No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

Date:11/03/04ISR Number: 4492157-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0531755A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	
RESPIRATORY							
( INHALATION)		Drug Abuser					

Date:11/03/04ISR Number: 4492173-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0350226A

Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown							
		Anger		Alcohol	SS		ORAL
		Disturbance In Attention					
		Impulse-Control Disorder					
		Memory Impairment					
		Road Traffic Accident					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/04ISR Number: 4492428-9Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 231193

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Required DAILY ORAL Intervention to Prevent Permanent Impairment/Damage		Aggression Agitation		Wellbutrin 100 Mg Galxio Smith Kline	PS	Galxio Smith Kline	ORAL

Date:11/03/04ISR Number: 4494805-9Report Type:Expedited (15-DaCompany Report #2004-BP-10340RO  
Age:52 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Completed Suicide Intentional Misuse	Literature	Diazepam Intensol Oral Solution (Concentrate), 5 Mg/Ml (Diazepam) Bupropion (Bupropion) Quetiapine (Quetiapine)	PS  SS SS		

Date:11/03/04ISR Number: 4494970-3Report Type:Expedited (15-DaCompany Report #2004-BP-10339RO  
Age:44 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Multiple Drug Overdose	Literature	Alprazolam Intensol [Oral Solution (Concentrate) ] 1 Mg/ Ml (Lorazepam) Bupropion (Bupropion) Clonazepam (Clonazepam)	PS  SS SS		

Date:11/03/04ISR Number: 4495701-3Report Type:Expedited (15-DaCompany Report #2004-BP-10338RO  
Age:50 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature	Alprazolam Intensol [Oral Solution (Concentrate)] 1 Mg/Ml (Alprazolam Bupropion (Bupropion) Acetaminophen W/Propoxyphene (Doxyphene)	PS  SS  SS		

Date:11/04/04ISR Number: 4493093-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0531546A  
Age:36 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Abdominal Pain Upper Convulsion Gastric Disorder Gastritis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Haematemesis				
		Medication Error				
		Migraine				
		Rectal Haemorrhage	Wellbutrin	PS	Glaxosmithkline	
		Regurgitation Of Food	Wellbutrin	SS	Glaxosmithkline	ORAL
		Vomiting	Ativan	C		
		Weight Decreased				

Date:11/04/04ISR Number: 4493108-6Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0347425A  
 Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Unknown	12 DAY		Retinopathy Hypertensive	Zyban	PS	Glaxosmithkline	ORAL
			Tinnitus	No Concurrent Medication	C		
			Visual Acuity Reduced				

Date:11/04/04ISR Number: 4493111-6Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0349671A  
 Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening							
150MG per day	22 DAY		Overdose	Zyban	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged			Personality Change				
			Suicidal Ideation				

Date:11/04/04ISR Number: 4493852-0Report Type:Expedited (15-DaCompany Report #2004UW21313  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required							
10 MG QD PO			Convulsion	Crestor	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage			Health Professional Other	Wellbutrin - Slow Release	SS		
				Wellbutrin - Slow Release	SS		

Date:11/04/04ISR Number: 4518624-XReport Type:Periodic  
Age:25 YR Gender: I/FU:I

Company Report #DSA\_24601\_2004

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 30 MG ONCE	1 DAY	Coma		Lorazepam	PS		
Initial or Prolonged 12.5 G ONCE		Overdose		Lorazepam	SS		
				Wellbutrin	SS		ORAL
PO	1 DAY			Wellbutrin	SS		
				Paxil	SS		
1220 MG ONCE	1 DAY			Paxil	SS		

Date:11/04/04ISR Number: 4679227-XReport Type:Direct  
Age: Gender: I/FU:I

Company Report #CTU 242672

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 50 ONE QAM		Therapeutic Response		Bupropione	PS		
		Unexpected With Drug Substitution					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/05/04ISR Number: 4493795-2Report Type:Expedited (15-DaCompany Report #US-ABBOTT-04P-163-0279296-00  
Age:20 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Valproic Acid	PS		ORAL
				Bupropion	SS		ORAL
				Topiramate	SS		ORAL

Date:11/05/04ISR Number: 4494299-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511243A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG See	Completed Suicide		Wellbutrin Sr	PS	Glaxosmithkline	
dosage text		Decreased Appetite					
		Depressed Mood					
		Disease Recurrence					
		Disturbance In Attention					
		Feelings Of Worthlessness					
		Gun Shot Wound					
		Major Depression					
		Sleep Disorder					
		Social Avoidant Behaviour					
		Suicidal Ideation					
		Treatment Noncompliance					
		Tremor					

Date:11/05/04ISR Number: 4494303-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0532210A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Drug Exposure During Pregnancy		Bupropion	PS	Glaxosmithkline	
		Kidney Malformation					
		Pregnancy					
		Renal Disorder					



Date:11/05/04ISR Number: 4494309-3Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0346394A  
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams		Zyban	PS	Glaxosmithkline	ORAL
1TAB Variable		Constipation					
dose	20	DAY					
		Diarrhoea					
		Dizziness					
		Dysphoria					
		Feeling Abnormal					
		Palpitations					
		Speech Disorder					

Date:11/08/04ISR Number: 4495823-7Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0329055A  
Age:46 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Balance Disorder
Initial or Prolonged	Feeling Abnormal
	Paraesthesia
	Speech Disorder
	Vertigo

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Visual Acuity Reduced

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	15 DAY		Zyban	PS	Glaxosmithkline	ORAL
40MG Per day			Pravachol	C		

Date:11/08/04ISR Number: 4495850-XReport Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0350696A  
Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other UNKNOWN		Anxiety  Depression Hallucination, Visual Hyperventilation Suicidal Ideation		Zyban	PS	Glaxosmithkline	

Date:11/08/04ISR Number: 4495852-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0350764A  
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Unknown	30 DAY	Arteriospasm Coronary		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged 10MG Per day	1 DAY	Chest Pain		Actiskenan	SS	Glaxosmithkline	ORAL
1 DAY				Efferalgan	C	Glaxosmithkline	ORAL

Date:11/08/04ISR Number: 4498732-2Report Type:Expedited (15-DaCompany Report #2004076041  
Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Condition Aggravated	Health	Zoloft (Sertraline)	PS	ORAL
50 MG (50 MG,	Cystitis	Professional			
1 IN 1 D),	Drug Effect Decreased				
ORAL	Feeling Jittery		Bupropion		
	Pollakiuria		Hydrochloride		
	Urinary Retention		(Bupropion Hydrochloride)	SS	ORAL
450 MG, ORAL			Clonazepam		
			(Clonazepam)	SS	
0.5 MG			Mirtazapine		
			(Mirtazapine)	SS	
15 MG (15 MG,					
1 IN 1 D)					

Date:11/08/04ISR Number: 4498808-XReport Type:Expedited (15-DaCompany Report #A0530762A  
Age:58 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Drug Ineffective	Consumer	Nicoderm Cq Patch			
Initial or Prolonged	Lung Carcinoma Cell Type		(Nicotine)	PS		
TRANSDERMAL	VARIABLE					
Required	Unspecified Stage I					
DOSE/TRANSERM						
Intervention to	Medication Error					
AL						
Prevent Permanent	Saliva Altered		Commit Nicotine			
Impairment/Damage	Throat Irritation		Polacrilex Lozenge,			
	Throat Tightness		2mg Lozenge 2 Mg			
			(Nicotine			
			Polacrilex)	SS		
BUCCAL	2 MG/ SINGLE					
DOSE/TRANSBUC						
CAL						

Freedom Of Information (FOI) Report

SEE IMAGE

	Wellbutrin (Bupropion Hydrochloride)	SS	ORAL
	Atorvastatin Calcium	C	
	Allegra-D	C	

Date:11/09/04ISR Number: 4496422-3Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20040908652  
Age:32 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Cardio-Respiratory Arrest		Topamax	PS		
OROPHARINGEAL						
	Completed Suicide		Bupropion	SS		
OROPHARINGEAL						
	Intentional Misuse		Venlafaxine	SS		
OROPHARINGEAL			Olanzapine	SS		

Date:11/09/04ISR Number: 4496423-5Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20040908721  
Age:20 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Acidosis		Topamax	PS		
OROPHARINGEAL						
	Blood Pressure Decreased		Valproic Acid	SS		
OROPHARINGEAL						
	Brain Oedema		Bupropion	SS		
OROPHARINGEAL						
	Coma		Clonidine	SS		
	Completed Suicide					
	Convulsion					
	Hypotension					
	Multiple Drug Overdose					
	Tachycardia					

Date:11/09/04ISR Number: 4496868-3Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0044946A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Alcohol Increased		Zyban	PS	Glaxosmithkline	ORAL
20TAB Single							
Initial or Prolonged		Convulsion					
dose							
		Intentional Misuse		Alcohol	C		ORAL
1 DAY							
		Somnolence					
		Suicide Attempt					

Date:11/09/04ISR Number: 4498293-8Report Type:Direct Company Report #CTU 231612  
 Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Urticaria		Wellbutrin Sr 150mg Tablet	PS		

Date:11/09/04ISR Number: 4499857-8Report Type:Direct Company Report #CTU 231633  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Rash		Zyban 150 Mg	PS		
Q DAY X 3							
Initial or Prolonged							
DAYS THEN BID 1 WK							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/09/04ISR Number: 4499859-1Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 231631

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Q DAY X 3		Rash Generalised		Zyban 150 Mg	PS		
DAYS THEN BID 1	WK						

Date:11/09/04ISR Number: 4502275-7Report Type:Expedited (15-DaCompany Report #S04-SWE-07224-01  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Pressure Increased Headache	Foreign Health	Cipramil (Citalopram Hydrobromide)	PS		ORAL
20 MG QD PO			Professional Other	Zyban (Bupropion Hydrochloride)	SS		ORAL
150 MG QD PO							

Date:11/09/04ISR Number: 4504356-0Report Type:Direct  
Age:23 YR Gender:Male I/FU:I

Company Report #CTU 232348

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pharmaceutical Product Complaint		Bupropion (Wellbutrin Sr)	PS		ORAL
150 MGM PO		Vomiting					
BID				Lithium Carbonate	C		
				Prolixin Decanoate	C		
				Abilify	C		
				Buspirone	C		

Date:11/10/04ISR Number: 4497818-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0393425A  
Age:17 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	20	DAY	Complex Partial Seizures		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged	20	DAY	Epileptic Aura		Gabitril	SS		ORAL
			Road Traffic Accident					
			Status Epilepticus					
			Tonic Clonic Movements					

Date:11/10/04ISR Number: 4497822-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0519165A  
Age:40 YR Gender:Female I/FU:F

Outcome	PT
Death	Agitation
	Anxiety
	Bedridden
	Completed Suicide
	Crying
	Derealisation
	Diarrhoea
	Dizziness
	Feeling Abnormal
	Gun Shot Wound
	Headache
	Insomnia
	Irritability
	Lethargy
	Nausea
	Nervousness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Pyrexia Social Avoidant Behaviour Toothache				
		Tremor Vomiting	Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/10/04ISR Number: 4497832-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0532724A  
Age: Gender: I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		PT Drug Abuser Overdose	Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	OTHER

Date:11/10/04ISR Number: 4497834-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0533112A  
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Other 150MG Per day	9 MON	PT Psoriasis	Wellbutrin Ssri	PS C	Glaxosmithkline	ORAL

Date:11/10/04ISR Number: 4497835-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0533144A  
Age: Gender: I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Other		PT Drug Abuser	Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	OTHER

Date:11/10/04ISR Number: 4497836-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0533147A  
Age: Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				



Hospitalization - Hot Flush Wellbutrin PS Glaxosmithkline ORAL  
Initial or Prolonged Mood Altered  
Nonspecific Reaction

Date:11/10/04ISR Number: 4677951-6Report Type:Direct Company Report #CTU 242643  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Anxiety		Bupropion Sr 150 Mg	PS		
Intervention to Prevent Permanent Impairment/Damage		Insomnia Tremor					

Date:11/11/04ISR Number: 4499211-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0533252A  
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Amphetamines Positive		Wellbutrin	PS	Glaxosmithkline	ORAL
6 MON		Hallucination					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/11/04ISR Number: 4499213-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLIN-A0533459A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	6 MON		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Alopecia					
		Deafness		Levoxine	C	Glaxosmithkline	
		Dermatitis Exfoliative					
		Thirst					

Date:11/12/04ISR Number: 4500096-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLIN-A0533311A  
Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Cardiac Failure		No Concurrent			
		Congestive		Medications	C		

Date:11/12/04ISR Number: 4500118-9Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLIN-B0351173A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		14 DAY		Bupropion	PS	Glaxosmithkline	
		Hyperthyroidism		Hydrochloride			
		Insomnia					

Date:11/15/04ISR Number: 4500997-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLIN-A0533749A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death				Wellbutrin	PS	Glaxosmithkline	
		Death					
		Overdose					

Date:11/15/04ISR Number: 4502033-3Report Type:Direct Company Report #CTU 231902  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG BID			Blood Glucose Decreased	Wellbutrin Sr 150 Mg Dsm Pharmaceuticals	PS	Dsm Pharmaceuticals	ORAL

ORAL

Zoloft	C
Metformin	C
Glipizide	C
Actos	C
Protonix	C
Tricor	C
Neurontin	C
Premarin	C

Date:11/15/04ISR Number: 4503716-1Report Type:Expedited (15-DaCompany Report #DEWYE183308NOV04  
Age:39 YR Gender:Female I/FU:I

Outcome	PT
Other	Antidepressant Drug Level Increased Drug Interaction Supraventricular Extrasystoles Tachycardia

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Freedom Of Information (FOI) Report

Ventricular Extrasystoles

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
SEE IMAGE, ORAL	85 DAY	Study	Trevilor (Venlafaxine Hydrochloride, Tablet, 0)	PS		ORAL
SEE IMAGE, ORAL	15 DAY		Zyban (Amfebutamone Hydrochloride, , 0)	SS		ORAL
			Zopiclone (Zopiclone)	C		
			Tibolone (Tibolone)	C		

Date:11/16/04ISR Number: 4502293-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0522931A  
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day		Blood Glucose Increased	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
	Fall DAY	Grand Mal Convulsion Joint Dislocation Loss Of Consciousness Medication Error Upper Limb Fracture	Professional	Ludiomil Symbyax	C C		

Date:11/16/04ISR Number: 4502295-2Report Type:Expedited (15-DaCompany Report #2004-03848  
Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day	3 DAY	Dizziness	Consumer	Bupropion	PS	Glaxosmithkline	ORAL

25MG Per day      Loss Of Consciousness      Atenolol      C  
Syncope  
Vomiting

Date:11/16/04ISR Number: 4502296-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0527318A  
Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 450MG Per day 4 MON Initial or Prolonged		Apathy	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Ineffective		Zoloft	SS		
		Overdose		Trileptal	SS		
		Pharmaceutical Product					
		Complaint					
		Sleep Disorder					
		Stool Analysis Abnormal					
		Suicide Attempt					
		Thinking Abnormal					

Date:11/16/04ISR Number: 4502305-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0533775A  
Age:25 YR Gender:Female I/FU:I

Outcome	PT
Other	Asthenia
	Convulsion
	Head Discomfort

Freedom Of Information (FOI) Report

Loss Of Consciousness

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1 MON			Wellbutrin XL	PS	Glaxosmithkline	ORAL

Date:11/16/04ISR Number: 4502327-1Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0356698A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Zyban	PS	Glaxosmithkline	
UNKNOWN	150MG	Twice					
per day	2	WK					
UNKNOWN		Grand Mal Convulsion					
UNKNOWN		Tongue Biting		Amitriptyline	C		
UNKNOWN				Allopurinol	C	Glaxosmithkline	
UNKNOWN				Hrt	C		
UNKNOWN				Simvastatin	C		
UNKNOWN				Enalapril	C		

Date:11/16/04ISR Number: 4502331-3Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0356825A  
 Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Consumer	Bupropion			
150MG	Twice	Hypoglycaemia		Hydrochloride	PS	Glaxosmithkline	
per day	14	DAY					
500MG	Three			Insulin Lantus	SS		
				Insulatard Novolet	SS		
				Novorapid	SS		
				Insulin Actrapid	SS		
				Metformin	SS		

times per day	Acetylcysteine	C	Glaxosmithkline
600MG Per day	Metoclopramide	C	Glaxosmithkline
10MG Three			
times per day	Prednisolon	C	Glaxosmithkline
35MG Weekly	Actonel	C	
10MG As	Paracetamol + Codeine	C	
required	Temazepam	C	
12MCG Twice	Foradil	C	
per day	Combivent	C	
500MCG Twice	Pulmicort	C	
per day	Atrovent	C	
20MG Per day	Pantozol	C	
250MG Twice	Theolair Retard	C	
per day	Lanoxin	C	Glaxosmithkline
.125MG Per			
day	Furosemide	C	Glaxosmithkline
40MG Per day	Spirolactone	C	
25MG Per day	Emcor	C	
2.5MG Per day	Coversyl	C	
4MG Per day	Lipitor	C	
20MG Per day	Spiriva	C	
18MCG Per day			





FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/16/04ISR Number: 4502791-8Report Type:Direct  
Age:80 YR Gender:Female I/FU:I

Company Report #CTU 232048

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash Maculo-Papular		Wellbutrin 150 Mg Xl	PS		ORAL
PO HS ORAL		Rash Pruritic					

Date:11/16/04ISR Number: 4503070-5Report Type:Direct  
Age:37 YR Gender:Female I/FU:I

Company Report #CTU 232128

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Anaphylactic Reaction		Welbutrin Xl 150 Mg			
Required				Glaxosmithklein	PS	Glaxosmithklein	ORAL
150 MG ONCE							
Intervention to							
PER DAY ORAL							
Prevent Permanent				Welbutrin Xl 150 Mg			
Impairment/Damage				Glaxosmithklein	SS		
				Multivitamin			
				Supplement	C		
				Glucosamine			
				Chondroitin	C		
				Alpha Lipoic Acid	C		
				Omega 3	C		
				Loratadine	C		

Date:11/16/04ISR Number: 4525687-4Report Type:Periodic  
Age:70 YR Gender:Female I/FU:I

Company Report #S04-USA-06882-01

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Interaction	Health Professional	Namenda (Memantine)	PS		
				Wellbutrin (Bupropion Hydrochloride)	SS		
150 MG QD							
				Wellbutrin (Bupropion Hydrochloride)	SS		
300 MG QD							

Aricept (Donepezil  
Hydrochloride) C

Date:11/17/04ISR Number: 4503372-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0524503A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Feeling Abnormal	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	5 DAY	Homicidal Ideation	Professional	Zoloft	C		ORAL
150MG Per day		Suicidal Ideation					

Date:11/17/04ISR Number: 4503373-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0525740A  
Age:16 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - MON		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Initial or Prolonged 15MG Per day		Grand Mal Convulsion		Abilify	C		
		Incontinence Tongue Biting					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/17/04ISR Number: 4503408-9Report Type:Expedited (15-DaCompany Report #IT-GLAXOSMITHKLINE-B0357042A  
 Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Zyban	PS	Glaxosmithkline	ORAL
300MG Per day	9 DAY	Disturbance In Attention		Nicorette	C	Glaxosmithkline	ORAL
		Fear					
		Headache					
		Ill-Defined Disorder					
		Psychotic Disorder					

Date:11/17/04ISR Number: 4504393-6Report Type:Direct Company Report #CTU 232292  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hot Flush		Wellbutrin Sr	PS		ORAL
150 MG PO BID		Insomnia					
		Pharmaceutical Product					
		Complaint					

Date:11/17/04ISR Number: 4504416-4Report Type:Direct Company Report #CTU 232296  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphthous Stomatitis		Wellbutrin	PS		ORAL
ONE TAB TWICE		Oral Intake Reduced					
A DAY ORAL		Speech Disorder					
		Weight Decreased					

Date:11/17/04ISR Number: 4524586-1Report Type:Periodic Company Report #US-JNJFOC-20040606830  
 Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Life-Threatening

Anaphylactoid Reaction

Health  
Professional

Levofloxacin  
(Levofloxacin)  
Wellbutrin  
(Bupropion  
Hydrochloride)  
Tablets

PS

SS

ORAL

ORAL

Date:11/18/04ISR Number: 4504405-XReport Type:Expedited (15-DaCompany Report #US-ROCHE-386012

Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Clonazepam	PS	Roche	
Other		Benign Prostatic		Wellbutrin	SS		ORAL
UNKNOWN		Hyperplasia		Wellbutrin	SS		ORAL
		Polyuria		Wellbutrin	SS		ORAL
		Urinary Retention		Wellbutrin	SS		ORAL
UNKNOWN				Zoloft	SS		
UNKNOWN				Remeron	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/18/04ISR Number: 4504611-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0351199A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Zyban	PS	Glaxosmithkline	ORAL
7 WK		Anxiety Depression Homicidal Ideation					

Date:11/18/04ISR Number: 4507091-8Report Type:Expedited (15-DaCompany Report #B0349825A  
Age:52 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardio-Respiratory Arrest Completed Suicide Drug Screen Positive	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL				Quetiapine (Formulation Unknown) (Quetiapine)	SS		ORAL
ORAL				Diazepam (Formulation Unknown) (Diazepam)	SS		ORAL

Date:11/18/04ISR Number: 4507097-9Report Type:Expedited (15-DaCompany Report #B0349826A  
Age:43 YR Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL				Temazepam (Formulation			

Unknown) (Temazepam) SS

ORAL

ORAL

Date:11/18/04ISR Number: 4507275-9Report Type:Expedited (15-DaCompany Report #B0349822A

Age:15 YR Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL					Aspirin (Aspirin)	SS		ORAL
ORAL					Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
ORAL					Topiramate (Topiramate)	SS		ORAL

Date:11/18/04ISR Number: 4507279-6Report Type:Expedited (15-DaCompany Report #B0349816A

Age:52 YR Gender: I/FU:I

Outcome	PT	Report Source
Death	Completed Suicide	Literature Health

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Professional

Dose	Duration	Product	Role	Manufacturer	Route
ORAL		Wellbutrin (Bupropion Hydrochloride)	PS		ORAL
ORAL		Aspirin (Aspirin)	SS		ORAL
ORAL		Paracetamol (Acetaminophen)	SS		ORAL

Date:11/18/04ISR Number: 4507280-2Report Type:Expedited (15-DaCompany Report #B0349817A  
Age:32 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Completed Suicide	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Toprimate (Formulation Unknown) (Topiramate)	SS		ORAL

Date:11/18/04ISR Number: 4507304-2Report Type:Expedited (15-DaCompany Report #B0349824A  
Age:30 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL				Citalopram (Citalopram)	SS		ORAL

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL					Paxil (Paroxetine Hydrochloride)	SS		ORAL
Date:11/18/04ISR Number: 4507305-4Report Type:Expedited (15-DaCompany Report #B0349823A Age:44 YR Gender: I/FU:I								
Death			Intentional Misuse	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL					Clonazepam (Clonazepam)	SS		ORAL
ORAL					Alprazolam (Alprazolam)	SS		ORAL

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL								



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL Valproic Acid (Valproic Acid) SS ORAL

Date:11/18/04ISR Number: 4510081-2Report Type:Expedited (15-DaCompany Report #B0349819A  
Age:50 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
				Alprazolam (Alprazolam)	SS		ORAL

Date:11/18/04ISR Number: 4510083-6Report Type:Expedited (15-DaCompany Report #B0349820A  
Age:43 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alcohol Use Completed Suicide	Literature Health Professional	Wellbutrin (Bupropion Hydrochloride)	PS		ORAL
				Ethanol (Alcohol)	SS		ORAL

Date:11/18/04ISR Number: 4510084-8Report Type:Expedited (15-DaCompany Report #B0349821A  
Age:21 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

Date:11/19/04ISR Number: 4505769-3Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0357250A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
UNKNOWN			Hallucination, Visual	Zyban	PS	Glaxosmithkline	
UNKNOWN			Nicotine Dependence	Tetralysal	C		
UNKNOWN				Iron	C		
UNKNOWN				Vitamin B12	C	Glaxosmithkline	
UNKNOWN							

Date:11/19/04ISR Number: 4505770-XReport Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0357272A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
UNKNOWN	150MG Twice		Convulsion	Zyban	PS	Glaxosmithkline	
per day			Syncope Vasovagal				

Date:11/19/04ISR Number: 4524232-7Report Type:Periodic Company Report #PHEH2004US09764  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
ORAL			Stool Analysis Abnormal	Zelnorm (Tegaserod) Tablet	PS		ORAL
			Health Professional	Wellbutrin- Slow			

Freedom Of Information (FOI) Report

ORAL  
 Release (Bupropion Hydrochloride) SS ORAL

Date:11/22/04ISR Number: 4506797-4Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0528265A  
 Age:24 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	ORAL
150MG See		Anger					
dosage text		Anxiety					
		Dyspepsia					
		Insomnia					
		Irritability					
		Mood Swings					
		Stomach Discomfort					

Date:11/22/04ISR Number: 4508967-8Report Type:Direct Company Report #CTU 232620  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Grand Mal Convulsion		Wellbutrin 200 Mg			
Initial or Prolonged				Glaxosmithkline	PS	Glaxosmithkline	ORAL
200 MG TWICE							

DAILY ORAL

Date:11/22/04ISR Number: 4509041-7Report Type:Direct Company Report #CTU 232548  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Confusional State		Wellbutrin Xl 300	PS		ORAL
1TABLET Q AM							
Hospitalization -		Grand Mal Convulsion					
PO							
Initial or Prolonged							
Required							

Intervention to  
Prevent Permanent  
Impairment/Damage

Date:11/22/04ISR Number: 4510269-0Report Type:Direct  
Age: Gender: I/FU:I

Company Report #USP 56965

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
TABLET, EXTENDED RELEASE		Medication Error		Budeprion Sr	PS	Teva	
TABLET, EXTENDED RELEASE				Wellbutrin Xl	SS	Glaxosmithkline	
TABLET, EXTENDED RELEASE				Wellbutrin Sr	SS	Glaxosmithkline	

Date:11/22/04ISR Number: 4510356-7Report Type:Direct  
Age:55 YR Gender:Male I/FU:I

Company Report #USP 56945

Outcome PT  
Dizziness  
Insomnia  
Lethargy

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Medication Error

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
TABLET			Wellbutrin Sr 100 Mg	PS	Glaxo Smith Kline	
TABLET			Wellbutrin Sr 150 Mg	SS	Teva	

Date:11/22/04ISR Number: 4511217-XReport Type:Direct Company Report #CTU 232739  
 Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Confusional State Fear Feeling Abnormal		Wellbutrin Xl 150mg Glaxosmithkline	PS	Glaxosmithkline	ORAL
150MG	1 A	Feeling Hot					
DAY	ORAL	Memory Impairment Mydriasis		Budeprion Sr 150mg Teva	SS	Teva	ORAL
150MG	X2 A	Paraesthesia					
DAY	ORAL	Pharmaceutical Product Complaint Thinking Abnormal					

Date:11/22/04ISR Number: 4511254-5Report Type:Direct Company Report #CTU 232725  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Rash		Bupropion 100	PS		
1 DAILY		Urticaria					

Date:11/22/04ISR Number: 4511269-7Report Type:Direct Company Report #CTU 232707  
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia		Bupropion 150 Mg	PS		ORAL
300 MG A DAY							
ORAL							

Date:11/23/04ISR Number: 4508022-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0164463A  
 Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	1 WK						
				Blood Pressure Medication	C		

Date:11/23/04ISR Number: 4508023-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0360153A  
 Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	15 MON	Depression					
60MG per day	YR	Drug Withdrawal Syndrome		Paxil	SS	Glaxosmithkline	ORAL
10MG Twice		Gastritis		Valium	C		
per day		Panic Attack					
				Prevacid	C		
				Nexium	C		
				Percocet	C		



Hospitalization -	Convulsion	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged	Intentional Misuse	Professional	Celexa	SS		
Other			Alcohol	SS		

Date:11/23/04ISR Number: 4508027-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0390353A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Lethargy					
per day	1 YR	Muscle Twitching		Zoloft	C		
50MG Per day		Obsessive-Compulsive Personality Disorder Somnolence					

Date:11/23/04ISR Number: 4508028-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0420648A  
 Age:49 YR Gender:Male I/FU:F

Outcome	PT
	Dizziness
	Malaise
	Nausea
	Pharmaceutical Product





150MG Twice	Abnormal Behaviour	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	Arthralgia				
	Myalgia	Synthroid	C	Glaxosmithkline	ORAL
	Pain In Extremity	Atacand	C		ORAL
		Atenolol	C		ORAL
		Lipitor	C		ORAL
20MG Unknown		Singulair	C		ORAL
10MG At night		Hydrocodone	C		

Date:11/23/04ISR Number: 4508033-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0425926A  
 Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Thrombocytopenia	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Professional				
Hospitalization -						
per day 60 WK						
Initial or Prolonged			Effexor Xr	C		ORAL
150MG Per day						
Other			Adderall Xr	C		ORAL

Date:11/23/04ISR Number: 4508034-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0426935A  
 Age:53 YR Gender:Female I/FU:F

Outcome	PT
	Adverse Event
	Chest Pain

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Headache Palpitations	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	5 YR			Wellbutrin	PS	Glaxosmithkline	ORAL
				Zoloft	C		
				Doxepin	C		
				Chlorazepate			
				Dipotassium	C		
				Geodon	C		
				Premarin	C		
				Vit E	C		
				Vit C	C	Glaxosmithkline	
				Calcium	C		
				Vit D	C		
				Multivitamin	C		
				Evening Primrose Oil	C		

Date:11/23/04ISR Number: 4508036-7Report Type:Periodic  
Age:41 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0427054A

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
5 WK		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
				Heroin	SS		
				Methadone	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508037-9Report Type:Periodic  
Age:34 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0427403A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice per day	17 DAY	Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
4 DAY				Quinine	C		ORAL

100U Three				Trazodone	C	ORAL
times per day						
100U Three				Topamax	C	ORAL
times per day						

Date:11/23/04ISR Number: 4508038-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0427685A  
 Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypersensitivity		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	6	Rash	MON				

Date:11/23/04ISR Number: 4508039-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0427726A  
 Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Lamictal	SS	Glaxosmithkline	ORAL
				Prevacid	C		
UNKNOWN				Claritin	C		
				Naprosyn	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508040-9Report Type:Periodic  
Age:47 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0427844A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Dysgeusia					
per day		Increased Appetite		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
300MG Per day		Weight Decreased		Ortho Novum	C		

Date:11/23/04ISR Number: 4508041-0Report Type:Periodic  
Age:16 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0427902A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Foaming At Mouth		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Grand Mal Convulsion		Zoloft	C		
200MG Per day	513 DAY	Tongue Biting					
50MG Per day							

Date:11/23/04ISR Number: 4508042-2Report Type:Periodic  
Age:52 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428249A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Asthenia					
per day	4 MON	Libido Decreased		Wellbutrin	SS	Glaxosmithkline	
		Prostate Infection		Multivitamin With Minerals	C		
				Doxycycline	C		

Date:11/23/04ISR Number: 4508043-4Report Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428261A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MCG Twice		Nausea					
per day	6	WK		Lexapro	C		

Date:11/23/04ISR Number: 4508044-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0428270A  
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Back Pain					
per day	2	YR					
		Dry Mouth					

Date:11/23/04ISR Number: 4508045-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0428272A  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysmenorrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Vaginal Haemorrhage					
per day	4	WK		Lamisil	C	Glaxosmithkline	
250MG Twice							
per day	6	DAY					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508046-XReport Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428273A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	
WK		Hyperhidrosis		Wellbutrin	SS	Glaxosmithkline	
WK		Irritability		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Per day 2	WK	Urticaria		Ultracet	C		

Date:11/23/04ISR Number: 4508048-3Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428274A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 1	WK	Anger		Keflex	C	Glaxosmithkline	
		Constipation		Advair	C	Glaxosmithkline	
		Dizziness		Detrol	C		
		Dry Mouth		Singulair	C		
		Headache		Prempro	C		
		Heart Rate Increased		Benadryl	C	Glaxosmithkline	
		Hyperhidrosis		Aciphex	C		
		Insomnia		Celebrex	C		
		Irritability		Ambien	C		
		Middle Insomnia		Centrum Multivitamin	C		
		Pruritus		Donnatal	C		
		Somnolence		Lomotil	C		
				Oxygen Therapy	C		
				Antibiotic			
				(Unspecified)	C		

Date:11/23/04ISR Number: 4508049-5Report Type:Periodic  
Age:69 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428403A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							

per day 2 MON

Neurontin C

Date:11/23/04ISR Number: 4508050-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0428415A  
Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arrhythmia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508051-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0428416A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							

per day

Date:11/23/04ISR Number: 4508052-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0428627A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508053-7Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428685A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
Other		Ventricular Extrasystoles		Vitamin E	C		
				Folate	C		
				Calcium	C		
				Lipitor	C		
				Aspirin	C	Glaxosmithkline	
				Prozac	C		

Date:11/23/04ISR Number: 4508054-9Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0428699A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4 WK	Anxiety		Wellbutrin	SS	Glaxosmithkline	ORAL
		Drug Interaction		Wellbutrin	SS	Glaxosmithkline	ORAL
		Nervousness		Thyroid Medication	SS		
UNKNOWN		19 DAY					
		Tremor					

Date:11/23/04ISR Number: 4508055-0Report Type:Periodic  
 Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428705A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Twice		Dizziness					
per day	4 YR	No Adverse Effect					
		Pharmaceutical Product					
		Complaint					

Date:11/23/04ISR Number: 4508056-2Report Type:Periodic  
 Age:59 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0428707A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Blood Glucose Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	5 YR	Diabetes Mellitus					
				Unknown Medication	C		
				Diltiazem	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508057-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0428729A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
3 DAY		Drug Interaction		Wellbutrin	PS	Glaxosmithkline	ORAL
		Fatigue		Unknown-Otc			
		Stomach Discomfort		Medication	C		
		Tunnel Vision					

Date:11/23/04ISR Number: 4508058-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0428825A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Unknown		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL



2 MON Drug Ineffective Wellbutrin PS Glaxosmithkline ORAL

Date:11/23/04ISR Number: 4508063-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0428974A  
Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Ineffective					
per day	6 WK			Klonopin	C		

Date:11/23/04ISR Number: 4508064-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0428987A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Food Craving		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	10 YR			Fluoxetine	C		

Date:11/23/04ISR Number: 4508065-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0429003A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blister		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1 WK	Rash		Lexapro	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508066-5Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429220A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100 per day	6 WK	Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
		Insomnia					
		Therapeutic Response					
		Unexpected					

Date:11/23/04ISR Number: 4508067-7Report Type:Periodic  
Age:66 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429228A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	10 DAY	Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508068-9Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429233A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Tongue Coated		Lexapro	C		
				Buspar	C		
				Multivitamin	C		

Date:11/23/04ISR Number: 4508069-0Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429243A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -	7 DAY	Erythema Multiforme		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Pruritus		Nexium	C		
Other		Urticaria		Lexapro	C		

Date:11/23/04ISR Number: 4508070-7Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429247A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	

Date:11/23/04ISR Number: 4508071-9Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429252A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508072-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429265A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3 WK						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508073-2Report Type:Periodic  
Age:20 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429489A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Disturbance In Attention		Wellbutrin	PS	Glaxosmithkline	ORAL
2 MON		Insomnia		Sonata	C		

Date:11/23/04ISR Number: 4508074-4Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429496A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day	2 YR			Ambien	C		
				Celebrex	C		
				Zyrtec	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508075-6Report Type:Periodic  
Age:42 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0429501A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Three times per day	7 YR	Insomnia					
		Nausea		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Unknown		Somnolence		Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Unknown	2 DAY						

Date:11/23/04ISR Number: 4508076-8Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429512A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Abdominal Pain Upper  
 Wellbutrin PS Glaxosmithkline ORAL  
 150MG Twice  
 per day 1 MON  
 Zoloft C  
 Fentanyl C  
 Excedrin C  
 Diazepam C

Date:11/23/04ISR Number: 4508077-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0429517A  
 Age: YR Gender:Female I/FU:I  
 Outcome PT Report Source Product Role Manufacturer Route  
 Dose Duration Amnesia Wellbutrin PS Glaxosmithkline ORAL  
 Other 150MG Twice  
 per day Syncope

Date:11/23/04ISR Number: 4508078-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0429523A  
 Age: Gender:Female I/FU:I  
 Outcome PT Report Source Product Role Manufacturer Route  
 Dose Duration Stool Analysis Abnormal Wellbutrin PS Glaxosmithkline ORAL

Date:11/23/04ISR Number: 4508079-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0429653A  
 Age:72 YR Gender:Female I/FU:I  
 Outcome PT Report Source Product Role Manufacturer Route  
 Dose Duration Tremor Wellbutrin PS Glaxosmithkline ORAL  
 150MG Twice  
 per day 3 WK



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Diovan	C	
Diclofenac	C	
Prilosec	C	Glaxosmithkline
Zocor	C	
Plavix	C	
Estradiol	C	
Synthroid	C	Glaxosmithkline
Vitamin D	C	
Calcium	C	

Date:11/23/04ISR Number: 4508080-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0429655A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Nausea					
per day				Claritin	C		

Date:11/23/04ISR Number: 4508081-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0429663A  
 Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abnormal Dreams		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Insomnia					
per day	3 MON			Allegra	C		

Date:11/23/04ISR Number: 4508082-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0429666A  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Dysuria					
per day	14 WK						

Escherichia Infection  
Pyrexia

Date:11/23/04ISR Number: 4508083-5Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429670A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	5 DAY						

Date:11/23/04ISR Number: 4508084-7Report Type:Periodic  
Age:61 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0429714A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
		Rash Vesicular		Wellbutrin	SS	Glaxosmithkline	ORAL
		Skin Irritation					
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508085-9Report Type:Periodic  
Age:57 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0429794A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice per day	4 YR	Nausea Pharmaceutical Product Complaint Stomach Discomfort Vomiting		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508086-0Report Type:Periodic  
Age:48 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0429806A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 YR	Hypertension Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Oral Contraceptives Clarinet	C C		

Date:11/23/04ISR Number: 4508087-2Report Type:Periodic  
Age:75 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0429814A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	YR	Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN		Irritability		Zoloft	SS		
UNKNOWN		Sexual Dysfunction		Effexor	SS		
UNKNOWN				Paxil	SS	Glaxosmithkline	
UNKNOWN				Lexapro	SS		
UNKNOWN				Lithium	C	Glaxosmithkline	

UNKNOWN

Thyroid

C

Date:11/23/04ISR Number: 4508088-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429839A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2	WK	Nonspecific Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508089-6Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0429969A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3	WK	Loss Of Consciousness		Wellbutrin	PS	Glaxosmithkline	ORAL
				Triphasil	C		

Date:11/23/04ISR Number: 4508090-2Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430008A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	1 MON	Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508091-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430036A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
		Crying		Unspecified			
		Depression		Medication	C		
		Dry Mouth					

Date:11/23/04ISR Number: 4508092-6Report Type:Periodic  
 Age:41 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0430077A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4 DAY	Dizziness		Lexapro	C		
		Headache		Provigil	C		
				Toprol Xl	C		
				Xanax	C		
				Protonix	C		

Date:11/23/04ISR Number: 4508093-8Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430271A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508094-XReport Type:Periodic  
 Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430277A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Deafness		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Dexedrine	C	Glaxosmithkline	
450MG Per day							

Date:11/23/04ISR Number: 4508095-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430281A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion					
450MG Per day				Ssri	C		

Date:11/23/04ISR Number: 4508096-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430284A  
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Insomnia					
150MG Per day	1 WK						

Date:11/23/04ISR Number: 4508097-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430445A  
Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Tinnitus					
200MG Per day				Celexa	C		
				Enbrel	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508098-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430468A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Painful Erection		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Three							
times per day							

Date:11/23/04ISR Number: 4508099-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430595A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
3 MON							

Date:11/23/04ISR Number: 4508100-2Report Type:Periodic  
Age:52 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0430608A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Lithium Carbonate	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508101-4Report Type:Periodic  
Age:17 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0430609A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
150MG Twice							
per day				Lexapro	C		

Date:11/23/04ISR Number: 4508102-6Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430614A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Hepatic Enzyme Increased	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2	MON					
				Topomax	C		
				Unknown Medication	C		
				Toprol	C		
				Ibuprofen	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508103-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430624A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Arthralgia	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	7	DAY					
			Back Pain				
			Chest Pain				
			Headache				
			Hot Flush				

Date:11/23/04ISR Number: 4508104-XReport Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430628A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Rash	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1	WK					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508105-1Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430641A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL
				Viagra	C		
				Aspirin	C	Glaxosmithkline	
				Vitamin	C		

Date:11/23/04ISR Number: 4508106-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430678A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day 1 YR	Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	

Date:11/23/04ISR Number: 4508107-5Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430684A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day 1 MON	Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
				Benadryl	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508108-7Report Type:Periodic  
Age:14 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0430701A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Hospitalization - Initial or Prolonged Other	Convulsion Overdose		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508109-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430706A  
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Single							
dose	1 DAY			Klonopin	C		

Date:11/23/04ISR Number: 4508110-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430709A  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1 YR						

Date:11/23/04ISR Number: 4508111-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430722A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tongue Ulceration		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	65 WK			Zoloft	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508112-9Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430723A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Tightness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 MON			Insulin	C		

Date:11/23/04ISR Number: 4508113-0Report Type:Periodic  
Age:65 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0430759A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lymphadenopathy		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice				Prednisone	C		
per day	31 DAY			Plaquenil	C		

Date:11/23/04ISR Number: 4508114-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0430853A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Insomnia		Effexor	C		
		Sleep Disorder		Klonopin	C		
		Tremor					

Date:11/23/04ISR Number: 4508115-4Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430883A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	6 YR						

Actos C  
 Glucophage C  
 Lantus C

Date:11/23/04ISR Number: 4508116-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430884A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Abuser		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508117-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0430889A  
 Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice		Insomnia					
per day	9	MON					

Amitriptyline C  
 Fish Oil C  
 Vitamin E C  
 Topamax C  
 Kcl C Glaxosmithkline  
 Parafon Forte C  
 Clonazepam C  
 Hctz C  
 Doxazosin C  
 Minipress C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Aceon C  
 Glucosamine C

Date:11/23/04ISR Number: 4508118-XReport Type:Periodic  
 Age:45 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0430893A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
300 per day	1 MON			Prozac	C		
				Trazodone	C		

Date:11/23/04ISR Number: 4508119-1Report Type:Periodic  
 Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430896A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Zyprexa	C		
				Levoxyl	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508120-8Report Type:Periodic  
 Age:27 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430901A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day		Insomnia		Propecia	C		
		Psychomotor Hyperactivity					

Date:11/23/04ISR Number: 4508121-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430939A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Diarrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL

Flatulence  
Headache  
Nausea

Date:11/23/04ISR Number: 4508122-1Report Type:Periodic  
Age:36 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430949A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyskinesia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	4	DAY					
75MG Per day				Effexor	SS		

Date:11/23/04ISR Number: 4508123-3Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430958A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	14	DAY					
		Diarrhoea		Tranxene	C		
		Insomnia		Glucovance	C		
		Nausea		Prilosec	C	Glaxosmithkline	
		Palpitations		Benicar	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508124-5Report Type:Periodic  
Age:72 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430961A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Per day	1 WK		Wellbutrin	PS	Glaxosmithkline	ORAL
				Digoxin	C	Glaxosmithkline	
				Aspirin	C	Glaxosmithkline	
				Flomax	C		

Date:11/23/04ISR Number: 4508125-7Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430967A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
				Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508126-9Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0430986A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Per day	14 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
				Lexapro	C		
				Glucovance	C		
				Benicar	C		

Date:11/23/04ISR Number: 4508127-0Report Type:Periodic  
Age:16 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431103A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other	3 DAY			Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508128-2Report Type:Periodic  
Age:26 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431164A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Per day	3	WK		Nyquil	C		

Date:11/23/04ISR Number: 4508129-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431168A  
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amenorrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice				Topamax	C		
per day	6	MON		Levoxyl	C	Glaxosmithkline	
				Zyprexa	C		

Date:11/23/04ISR Number: 4508130-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431175A  
 Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day	14	YR		Vivelle	C		
200MG Twice				Celebrex	C		ORAL
per day							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508131-2Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431188A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
4 YR				Prozac	C		

Date:11/23/04ISR Number: 4508132-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431205A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Atenolol	C		
per day	3 DAY			Thyroid Medication	C		

Date:11/23/04ISR Number: 4508133-6Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431220A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Urine Present		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day		Muscle Spasms					

Date:11/23/04ISR Number: 4508134-8Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431409A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK	Dry Mouth		Valium	C		
				Metabolife	C		
				Ginkoba	C		
				Unknown Medication	C		

Date:11/23/04ISR Number: 4508135-XReport Type:Periodic  
Age:30 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431432A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Abnormal Sensation In Eye	Wellbutrin	PS	Glaxosmithkline	ORAL
2	MON						

Date:11/23/04ISR Number: 4508136-1Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431435A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -			Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
Initial or Prolonged							
per day	5	DAY		Inderal	C		

Date:11/23/04ISR Number: 4508137-3Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0431436A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Dry Mouth	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	9	DAY	Insomnia				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508138-5Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431453A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508139-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431799A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				Klonopin	C		
				Methotrexate	C		
				Darvocet	C		
				Prednisolone	C	Glaxosmithkline	
				Flexeril	C		
				Salagen	C		
				Folic Acid	C		
				Phenergan	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508140-3Report Type:Periodic  
Age:69 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431801A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Loss Of Libido		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Neurontin	C		
per day	6	MON		Norvasc	C		
				Cardura	C		

Date:11/23/04ISR Number: 4508141-5Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431811A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

100MG Per day	Dizziness	Wellbutrin	PS	Glaxosmithkline	ORAL
		Tenormin	C		
		Avapro	C		
		Lipitor	C		

Date:11/23/04ISR Number: 4508142-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431817A  
 Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Formication					
per day	1	MON					
		Hepatic Enzyme Increased		Topamax	SS		
		Insomnia		Phentermine	SS	Glaxosmithkline	
		Nausea					
		Stress					

Date:11/23/04ISR Number: 4508143-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431821A  
 Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4	MON		Lorazepam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508144-0Report Type:Periodic  
Age:61 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431826A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Otitis Media		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:11/23/04ISR Number: 4508145-2Report Type:Periodic  
Age:36 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431837A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
Life-Threatening							
3 WK							
Other		Hypersensitivity Laryngeal Oedema Urticaria		No Concurrent Medications	C		

Date:11/23/04ISR Number: 4508146-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431971A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	6 DAY						

Date:11/23/04ISR Number: 4508147-6Report Type:Periodic  
Age:14 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431973A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
100MG See							
dosage text	2 MON						

Date:11/23/04ISR Number: 4508148-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0431975A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/23/04ISR Number: 4508149-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432093A  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:11/23/04ISR Number: 4508150-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432102A  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Vaginal Haemorrhage		Wellbutrin	PS	Glaxosmithkline	ORAL
150MCG Per							
day	2 DAY						
				Tylenol Cold + Sinus	C		
				Birth Control Pill	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508151-8Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432105A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/23/04ISR Number: 4508152-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432115A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Memory Impairment Musculoskeletal Stiffness Petit Mal Epilepsy		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508153-1Report Type:Periodic  
 Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432124A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Therapeutic Response Unexpected		Wellbutrin	PS	Glaxosmithkline	

Date:11/23/04ISR Number: 4508154-3Report Type:Periodic  
 Age:80 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0432134A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	8 WK	Dry Mouth					
25MG Per day	4 MON	Feeling Abnormal		Paxil	SS	Glaxosmithkline	ORAL
.75MG Per day		Hyperhidrosis		Synthroid	C	Glaxosmithkline	

TRANSDERMAL	Tremor	Duragesic	C	
	Vision Blurred	Bactrim	C	Glaxosmithkline
		Allopurinol	C	Glaxosmithkline
		Colchicine	C	
		Imipramine	C	

Date:11/23/04ISR Number: 4508155-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432135A  
 Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Pharmaceutical Product					
per day	1 YR	Complaint		Zyrtec	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508156-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432143A  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY	Chills		Oxycontin	C		
		Dizziness		Oxycodone Immediate			
		Hyperhidrosis		Release	C		
		Insomnia		Cipro	C	Glaxosmithkline	
				Asacol	C	Glaxosmithkline	
				Bentyl	C		
				Valium	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508157-9Report Type:Periodic  
Age:37 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432145A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Glossodynia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4	MON		Allopurinol	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508158-0Report Type:Periodic  
Age:77 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432148A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	3	DAY		Metoprolol	C		
		Nausea					

Date:11/23/04ISR Number: 4508159-2Report Type:Periodic  
Age:33 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0432150A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Diarrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	5	YR		Celexa	C		

Date:11/23/04ISR Number: 4508160-9Report Type:Periodic  
Age:29 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0432178A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day				Zoloft	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG In the							
morning	3	MON					
300MG In the				Wellbutrin	SS	Glaxosmithkline	ORAL
morning	6	DAY					
UNKNOWN	10MG	Unknown		Adderall	C		
				Lexapro	C		ORAL
				Adderall Xr	C		ORAL
20MG Per day							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	1	YR					
		Lethargy		Xanax	C		
		Somnolence		Cardura	C		
				Valium	C		
				Clonidine	C		
				Avandia	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04  
 Age:47 YR  
 Gender:Female  
 I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432272A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	100MG Per day	Euphoric Mood		Wellbutrin	PS	Glaxosmithkline	ORAL
		Fatigue		Zoloft	C		
		Insomnia		Hctz	C		
		Somnolence		Allopurinol	C	Glaxosmithkline	
				Potassium Citrate	C		
				Klor-Con	C	Glaxosmithkline	
				Ibuprofen	C	Glaxosmithkline	
				Chromium	C		

Date:11/23/04  
 Age:  
 Gender:Male  
 I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432301A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Erectile Dysfunction		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04  
 Age: YR  
 Gender:Female  
 I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432325A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Twice per day	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04  
 Age:  
 Gender:  
 I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432425A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urinary Retention		Wellbutrin	PS	Glaxosmithkline	

Date:11/23/04ISR Number: 4508167-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432430A  
Age: YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	
1	YR						

Date:11/23/04ISR Number: 4508168-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432444A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Mental Disorder					

Date:11/23/04ISR Number: 4508169-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432446A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
				Wellbutrin	C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508170-1Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0432450A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
1	YR						

Date:11/23/04ISR Number: 4508171-3Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432579A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Somnolence		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	6 WK	Weight Increased		Narcotics	C		

Date:11/23/04ISR Number: 4508172-5Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432599A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Balance Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	152 WK	Decreased Appetite		Celexa	C		
		Dizziness		Strattera	C		
		Migraine					

Date:11/23/04ISR Number: 4508173-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432605A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	3 YR			Norvasc	C		
				Aciphex	C		

Date:11/23/04ISR Number: 4508174-9Report Type:Periodic  
Age:45 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0432851A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Nausea					
per day		Vomiting		Zoloft	C		
250MG per day				Lamictal	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508175-0Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0432884A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	1 WK						

Date:11/23/04ISR Number: 4508176-2Report Type:Periodic  
Age:16 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0433130A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Visual Disturbance		Wellbutrin	PS	Glaxosmithkline	ORAL
				Adderall	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508177-4Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0433132A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG							
Alternate							
days	3	WK		Effexor	C		

Date:11/23/04ISR Number: 4508178-6Report Type:Periodic  
 Age:35 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0433140A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bronchitis		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3	WK					

Date:11/23/04ISR Number: 4508179-8Report Type:Periodic  
 Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0433142A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anaphylactic Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL
Life-Threatening Hospitalization - Initial or Prolonged Other		Dyspnoea					

Date:11/23/04ISR Number: 4508180-4Report Type:Periodic  
 Age:15 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0433258A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day	2	DAY					

Date:11/23/04ISR Number: 4508181-6Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0433268A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Blood Creatine		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Phosphokinase Increased					
per day							

Date:11/23/04ISR Number: 4508182-8Report Type:Periodic  
Age:24 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0433303A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Rash					
per day	1	MON					
437 DAY				Lexapro	C		
				Wellbutrin	C	Glaxosmithkline	
200MG Twice							
per day							

Date:11/23/04ISR Number: 4508183-XReport Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0433304A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Flushing Paraesthesia		Wellbutrin	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508184-1Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0438802A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75MG Twice per day	2 YR	Drug Ineffective Insomnia Paraesthesia		Wellbutrin Geodon Atenolol	PS C C	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508185-3Report Type:Periodic  
Age:66 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0438819A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	8 MON	Blood Pressure Increased Eructation Flatulence		Wellbutrin Azulfidine Folic Acid Lopressor Vasotec Ultracet Zocor Protonix Flexeril Ambien Clonazepam Aspirin Gas-X	PS C C C C C C C C C C C C	Glaxosmithkline Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508186-5Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0438825A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3 WK	Weight Increased		Wellbutrin None	PS C	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508187-7Report Type:Periodic  
Age:63 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0438879A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	In the	Restlessness					
morning							

Date:11/23/04ISR Number: 4508188-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439005A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Night Sweats		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Twice						
per day	2	WK					

Date:11/23/04ISR Number: 4508189-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439006A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Per day	Night Sweats					
	7	DAY					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508190-7Report Type:Periodic  
Age:76 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439036A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	4 WK	Vision Blurred		Wellbutrin	PS	Glaxosmithkline	ORAL
				Lopid	C		
				Aricept	C		
				Foltx	C		
				Vioxx	C		
				Aciphex	C		
				Multivitamin	C		
				Aspirin	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508191-9Report Type:Periodic  
Age:24 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439038A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 WK		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508193-2Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439053A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	9 MON	Vaginal Haemorrhage		Wellbutrin	PS	Glaxosmithkline	ORAL
				Effexor Xr	C		

Date:11/23/04ISR Number: 4508194-4Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439245A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Palpitations		Wellbutrin	PS	Glaxosmithkline	ORAL

Weight Decreased

Celexa

C

Date:11/23/04ISR Number: 4508195-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439248A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoaesthesia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Paraesthesia					

Date:11/23/04ISR Number: 4508196-8Report Type:Periodic  
Age:35 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0439258A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Malaise		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Unknown							

Date:11/23/04ISR Number: 4508197-XReport Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439264A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508198-1Report Type:Periodic  
Age:60 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439275A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Twice	Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
per day				Cephalexin	C	Glaxosmithkline	
				Protonix	C		
				Lisinopril	C		
				Furosemide	C	Glaxosmithkline	
				Amiodarone	C		
				Allopurinol	C	Glaxosmithkline	
				Pravachol	C		
				Coreg	C	Glaxosmithkline	
				Digoxin	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508199-3Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0439351A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	
Other		Drug Exposure Via Breast Milk					

Date:11/23/04ISR Number: 4508200-7Report Type:Periodic  
Age:63 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439409A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Twice	Condition Aggravated		Wellbutrin	PS	Glaxosmithkline	ORAL
per day				Desipramine	C		
				Klonopin	C		
				Tenormin	C		
				Vasotec	C		
				Synthroid	C	Glaxosmithkline	
				Plavix	C		

Imdur	C	
Catapres	C	
Hytrin	C	
Zantac	C	Glaxosmithkline
Lansoprazole	C	
Lipitor	C	

Date:11/23/04ISR Number: 4508201-9Report Type:Periodic  
 Age:69 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0439412A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	6	WK		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2	MON		Wellbutrin	SS	Glaxosmithkline	ORAL
UNKNOWN		2 MON		Klonopin	SS		
UNKNOWN				Xanax	SS		
				Unithroid	C	Glaxosmithkline	
				Ambien	C		
				Seroquel	C		
				Thyroid Supplement	C		
				Clonopin	C		
				Remeron	C		
				Zocor	C		



3	YR	Gastrointestinal Motility Disorder	Wellbutrin	PS	Glaxosmithkline	ORAL
			Effexor Xr	C		
			Multivitamin	C		
			Vitamin B	C		
			Vitamin C	C	Glaxosmithkline	
			Vitamin E	C		
			Iron Supplement	C		
			Calcium	C		
			Levoxyl	C	Glaxosmithkline	
			Fosamax	C		

Date:11/23/04ISR Number: 4508208-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439664A  
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Capillary Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Dizziness					
per day	3	DAY					
		Dry Mouth					
		Dyspnoea					
		Mental Status Changes					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508209-3Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0439677A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bladder Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508210-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439689A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination, Auditory		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							

Date:11/23/04ISR Number: 4508211-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0439827A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day				Lisinopril	C		

Date:11/23/04ISR Number: 4508212-3Report Type:Periodic  
Age:23 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0439835A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Diarrhoea					
per day	4 DAY	Fatigue					
		Headache					
		Rash Pruritic					

Date:11/23/04ISR Number: 4508213-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439846A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1	MON		Hormone Replacement	C		

Date:11/23/04ISR Number: 4508214-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439969A  
Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2	YR		Celexa	C		
				Proscar	C		
				Flomax	C		

Date:11/23/04ISR Number: 4508215-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0439976A  
Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea Vomiting		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508216-0Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0440023A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaemia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508217-2Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440184A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Twice		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	8 YR			Premarin	C		

Date:11/23/04ISR Number: 4508218-4Report Type:Periodic  
Age:54 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440364A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 WK	Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Propecia	C		
				Aspirin	C	Glaxosmithkline	
				Aceon	C		
				Arthrotec	C		
				Lipitor	C		

Date:11/23/04ISR Number: 4508219-6Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440368A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 WK						

Date:11/23/04ISR Number: 4508220-2Report Type:Periodic  
Age:30 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440385A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3	MON					

Date:11/23/04ISR Number: 4508221-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440391A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	5	WK					

Date:11/23/04ISR Number: 4508222-6Report Type:Periodic  
Age:63 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0440427A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lip Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice		Movement Disorder					
per day	3	MON		Adderall	C		
				Trazodone	C		
				Lexapro	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508223-8Report Type:Periodic  
 Age:43 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440559A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Twice	1	MON		Wellbutrin	PS	Glaxosmithkline	ORAL
per day				Xanax	C		
				Clonazepam	C		
				Topamax	C		
				Claritin	C		

Date:11/23/04ISR Number: 4508224-XReport Type:Periodic  
 Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440572A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75MG Twice	1	MON		Wellbutrin	PS	Glaxosmithkline	ORAL
per day				Anti-Diabetic Medication	C		ORAL

Date:11/23/04ISR Number: 4508225-1Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440590A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508226-3Report Type:Periodic  
 Age:22 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440591A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice				Wellbutrin	PS	Glaxosmithkline	ORAL

per day 2 WK

None

C

Date:11/23/04ISR Number: 4508227-5Report Type:Periodic  
Age:69 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440592A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	5 YR	Blood Creatinine		Wellbutrin	PS	Glaxosmithkline	ORAL
		Decreased		Lipitor	C		
		Blood Urea Decreased		Lotrel	C		
				Hytrin	C		
				Allopurinol	C	Glaxosmithkline	
				Lexapro	C		

Date:11/23/04ISR Number: 4508228-7Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440603A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Ear Discomfort		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	10 DAY	Muscle Tightness					
				Loestrin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508229-9Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0440612A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Priapism		Wellbutrin	PS	Glaxosmithkline	

Date:11/23/04ISR Number: 4508230-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440808A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	4 MON	Drug Ineffective Pharmaceutical Product Complaint					

Date:11/23/04ISR Number: 4508231-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0440815A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice	5 MON						
per day							

Date:11/23/04ISR Number: 4508232-9Report Type:Periodic  
Age:45 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0440822A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
Disability							
150MG Twice	9 WK						
per day							

Date:11/23/04ISR Number: 4508233-0Report Type:Periodic  
Age:48 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0440825A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Asthenia		Wellbutrin	PS	Glaxosmithkline	
150MG Twice		Dizziness					
per day	7	WK					
		Tremor		Wellbutrin	SS	Glaxosmithkline	
150MG Per day				Lithium	C	Glaxosmithkline	ORAL
				Abilify	C		
				Trazodone	C		
				Lisinopril	C		

Date:11/23/04ISR Number: 4508235-4Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441004A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
2	WK						

Date:11/23/04ISR Number: 4508236-6Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441020A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Asthenia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2	MON					
		Headache					
		Increased Appetite		Birth Control Pill	C		
		Nausea					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508237-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441126A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Paxil	C	Glaxosmithkline	
				Lexapro	C		

Date:11/23/04ISR Number: 4508238-XReport Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441156A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Chills					
per day	3 YR	Dizziness		Prilosec	C	Glaxosmithkline	
		Fatigue		Alprazolam	C		
		Nausea		Darvocet	C		

Date:11/23/04ISR Number: 4508239-1Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441157A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508240-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441479A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN	150MG Twice						
per day				Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day							

Date:11/23/04ISR Number: 4508241-XReport Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441485A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508242-1Report Type:Periodic  
Age:33 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0441527A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day		Burning Sensation		Effexor Xr	C		
UNKNOWN		MON		Kariva	C		
		Dizziness					
		Emotional Disorder					
		Erythema					
		Feeling Abnormal					
		Flushing					
		Hot Flush					
		Nausea					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508243-3Report Type:Periodic  
Age:51 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0441534A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	100MG Per day	375 DAY	Asthenia		Wellbutrin	PS	Glaxosmithkline	ORAL
	50MG Unknown	2 YR	Euphoric Mood		Zoloft	SS		ORAL
	240MG Per day		Fatigue		Verapamil	C		
			Mental Disorder					

Date:11/23/04ISR Number: 4508244-5Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441633A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	200 per day	2 YR	Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508245-7Report Type:Periodic  
Age:38 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0441649A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Twice		Pharmaceutical Product		Wellbutrin	PS	Glaxosmithkline	ORAL
	per day		Complaint					
	10MG Per day		Rash		Bextra	C		ORAL
	20MG Per day				Nexium	C		ORAL
	1TAB Per day				Microgestin Fe	C		ORAL

Date:11/23/04ISR Number: 4508246-9Report Type:Periodic  
Age:75 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441748A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Muscle Spasms		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day	1 WK			Lescol	C		
				Quinine	C		

Date:11/23/04ISR Number: 4508247-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441749A  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day		Stool Analysis Abnormal		Zoloft	C		
		Tremor		Klonopin	C		

Date:11/23/04ISR Number: 4508248-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441762A  
 Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hyperhidrosis		Wellbutrin	PS	Glaxosmithkline	
				Wellbutrin	SS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508249-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441763A  
 Age: Gender:Female I/FU:I

Outcome	PT
	Abdominal Pain Upper
	Anorexia
	Diarrhoea
	Gastrooesophageal Reflux

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Disease Headache Vomiting				
Dose	Duration		Report Source	Product	Role	Manufacturer
200MG Twice				Wellbutrin	PS	Glaxosmithkline
per day	6 MON					ORAL
				Prevacid	C	
				Seroquel	C	
				Trazodone	C	

Date:11/23/04ISR Number: 4508250-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441772A  
 Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose						
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline
150MG Twice						ORAL
per day						

Date:11/23/04ISR Number: 4508251-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441784A  
 Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose						
Other		Amnesia		Wellbutrin	PS	Glaxosmithkline
150MG Twice						ORAL
per day	8 YR	Confusional State				
				Prozac 20	C	
				Ambien	C	

Date:11/23/04ISR Number: 4508252-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0441791A  
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose						
Other		Asthma		Wellbutrin	PS	Glaxosmithkline
150MG Per day	2 WK					ORAL

Effexor Xr	C	
Proventil	C	Glaxosmithkline
Ventolin	C	Glaxosmithkline
Flovent	C	Glaxosmithkline
Allegra	C	
Ibuprofen	C	Glaxosmithkline
Baby Aspirin	C	Glaxosmithkline

Date:11/23/04ISR Number: 4508253-6Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0442007A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Ineffective					
per day	2 YR	Nausea		Trazodone	C		
		Pharmaceutical Product		Klonopin	C		
		Complaint		Estrogen Patch	C		

Date:11/23/04ISR Number: 4508254-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442040A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Pharmaceutical Product					
per day		Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508255-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442042A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Contrast Media Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508256-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442060A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
2 WK		Anxiety Feeling Abnormal					

Date:11/23/04ISR Number: 4508257-3Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442155A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	4 MON			Depakote	C		
				Neurontin	C		
				Geodon	C		
				Prevacid	C		
				Zocor	C		
				Mobic	C		
				Miralax	C		

Date:11/23/04ISR Number: 4508258-5Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442156A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Stool Analysis Abnormal		Wellbutrin Sr	PS	Glaxosmithkline	
				Wellbutrin Xl	SS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508259-7Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442162A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Accommodation Disorder					
per day		Extrasystoles					
				Prozac	C		
				Amitriptyline	C		
				Vioxx	C		
				Methotrexate	C		
				Folic Acid	C		
				Calcium	C		
				Imitrex	C	Glaxosmithkline	
				Fiorinal	C		
				Aspirin	C	Glaxosmithkline	
				Vitamin	C		

Date:11/23/04ISR Number: 4508260-3Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442170A

Outcome	PT
	Dizziness
	Drug Ineffective

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Drug Withdrawal Syndrome  
Nausea

Dose	Duration	Report Source	Product	Role	Manufacturer	Route			
300MG Per day	2 MON		Wellbutrin	PS	Glaxosmithkline	ORAL			
			Paxil	SS		ORAL			
			Wellbutrin	SS		ORAL			
						Estradiol	C		
						Hyzaar	C		
						Prilosec	C	Glaxosmithkline	
						Meclizine	C		

Date:11/23/04ISR Number: 4508261-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442174A  
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Dizziness					
per day	2 WK	Dry Mouth		Tenormin	C		
		Insomnia		Zocor	C		
				Aspirin	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508262-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442176A  
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	6 WK						

Date:11/23/04ISR Number: 4508263-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442330A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	5 WK						

Date:11/23/04ISR Number: 4508264-0Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442354A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK	Tremor					

Date:11/23/04ISR Number: 4508265-2Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442438A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Libido Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
2 YR				Effexor	SS		
UNKNOWN							

Date:11/23/04ISR Number: 4508266-4Report Type:Periodic  
Age:57 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442508A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1 MON			Xanax	C		

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Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508267-6Report Type:Periodic  
Age:66 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0442558A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 WK	Agitation Anorexia Constipation Depression Dry Mouth Insomnia Weight Decreased		Wellbutrin Altace Niaspan Actonel Calcium Fish Oil	PS C C C C C	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508268-8Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442561A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 WK	Drug Ineffective Dry Mouth Influenza		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508269-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442562A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression		Wellbutrin Sr Pain Medications	PS C	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508270-6Report Type:Periodic  
Age:30 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0442654A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Pharmaceutical Product  
per day 18 MON  
Complaint

Antibiotics C

Date:11/23/04ISR Number: 4508271-8Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442655A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	8 YR			Celexa	C		

Date:11/23/04ISR Number: 4508272-XReport Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442772A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Confusional State		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	3 WK	Insomnia					
		Irritability		Valium	C		
				Vicodin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508273-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442779A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day	1 WK	Dyspepsia Flatulence Muscle Spasms Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508274-3Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442810A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 WK	Abdominal Distension Drug Ineffective Periorbital Oedema		Wellbutrin	PS	Glaxosmithkline	ORAL
				Patanol Eye Drops	C		
				Flonase	C	Glaxosmithkline	
				Nasalcrom	C		
				Benadryl	C	Glaxosmithkline	
				Allegra	C		
				Docusate	C		
				Zelnorm	C		
				Protonix	C		
				Prednisone	C		
				Creon	C		
				Pentasa	C	Glaxosmithkline	
				Ambien	C		
				Cytotec	C		
				Miacalcin Nasal Spray	C		
				Unspecified Medication	C		

Date:11/23/04ISR Number: 4508275-5Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442976A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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100MG Per day	2	WK	Abdominal Pain	Wellbutrin	PS	Glaxosmithkline	ORAL
			Diarrhoea				
			Gastrooesophageal Reflux Disease				

Date:11/23/04ISR Number: 4508276-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442996A  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1	MON	Arthralgia	Wellbutrin	PS	Glaxosmithkline	ORAL
			Hypoaesthesia	Amitriptyline	C		
			Insomnia	Xanax	C		
			Muscle Tightness				

Date:11/23/04ISR Number: 4508277-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0443000A  
 Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2	YR	White Blood Cell Count	Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN			Increased	Unspecified Inhaler	C		
				Promethazine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Bextra C

Date:11/23/04ISR Number: 4508278-0Report Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443008A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	7	WK					

Date:11/23/04ISR Number: 4508279-2Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443199A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Influenza Like Illness					
per day	2	MON					
		Insomnia		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Irritability		Atenolol	C		
		Parosmia		Altace	C		
		Tremor		Lamictal	C	Glaxosmithkline	
				Zonegran	C		
				Flexeril	C		
				Omeprazole	C	Glaxosmithkline	
				Mobic	C		
				Tricor	C		

Date:11/23/04ISR Number: 4508280-9Report Type:Periodic  
Age:39 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0443202A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	5	WK					
				Provigil	C		

Betaseron

C

Date:11/23/04ISR Number: 4508281-0Report Type:Periodic  
Age: YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0443208A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
		Depressed Mood					
		Feeling Abnormal		Zoloft	C		
		Headache		Multivitamin	C		
		Memory Impairment		Flax Seed Oil	C		
		Nasopharyngitis					
		Vision Blurred					

Date:11/23/04ISR Number: 4508282-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443341A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
8 YR		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL



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Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508283-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443381A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pruritus Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508284-6Report Type:Periodic  
 Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443389A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	1 YR	Depression Irritability		Wellbutrin	PS	Glaxosmithkline	ORAL
2 WK		Nervousness Sexual Dysfunction		Wellbutrin	SS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508285-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443390A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
MON		Dizziness Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508286-XReport Type:Periodic  
 Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443476A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	5 WK	Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
				Ultram	C		
				Sudafed	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508287-1Report Type:Periodic  
Age:39 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0443496A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	142	DAY					

Date:11/23/04ISR Number: 4508288-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443511A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
		Sexual Dysfunction					
per day	2	YR		Synthroid	C	Glaxosmithkline	
		Weight Decreased					

Date:11/23/04ISR Number: 4508289-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443524A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
400MG Twice		Blood Pressure Systolic		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Increased					

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Date:11/23/04ISR Number: 4508290-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443647A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Insomnia					
per day	15 DAY						

Date:11/23/04ISR Number: 4508291-3Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443754A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	1 MON	Pharmaceutical Product Complaint					

Date:11/23/04ISR Number: 4508292-5Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443771A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Epistaxis		Wellbutrin Sr	PS	Glaxosmithkline	

Date:11/23/04ISR Number: 4508293-7Report Type:Periodic  
Age:48 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0443782A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Hiv Treatments (Unspecified)	C		
				Lorazepam	C		
				Electroconvulsive Therapy	C		

Date:11/23/04ISR Number: 4508294-9Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443785A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	3	MON		Effexor Xr	C		
				Synthroid	C	Glaxosmithkline	
				Neurontin	C		
				Clonazepam	C		

Date:11/23/04ISR Number: 4508295-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443929A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin Sr	PS	Glaxosmithkline	
				Adderall	C		

Date:11/23/04ISR Number: 4508296-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443939A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
150MG Twice							
per day							

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Date:11/23/04ISR Number: 4508297-4Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443943A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	5 YR			Morphine	C		
				Estrogen	C		
				Synthroid	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508298-6Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0444026A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression		Wellbutrin	PS	Glaxosmithkline	ORAL
		Delusion		Wellbutrin	SS	Glaxosmithkline	ORAL
		Flashback					
		Hallucination					
		Medication Error					
		Paranoia					

Date:11/23/04ISR Number: 4508299-8Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0444110A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Three							
times per day	7 YR	Insomnia					
		Sleep Disorder		Hormone Replacement			
				Therapy	C		
				Blood Pressure			
				Medication	C		

Date:11/23/04ISR Number: 4508300-1Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0444127A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperaesthesia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3	WK		Paxil	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508301-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0444128A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cystitis		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Twice							
per day	3	MON					
		Insomnia					
		Pollakiuria					

Date:11/23/04ISR Number: 4508302-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0444130A  
 Age:39 YR Gender:Female I/FU:I

Outcome	PT
	Abnormal Dreams
	Anger
	Anxiety
	Depression
	Fatigue

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Insomnia Irritability	Report Source	Product	Role	Manufacturer	Route
150MG Per day				Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	6 DAY			Wellbutrin Xl	SS	Glaxosmithkline	ORAL
				Xanax	C		
				Calcium	C		

Date:11/23/04ISR Number: 4508303-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0444161A  
 Age:37 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1 MON	Drug Interaction					
500MG Per day		Insomnia		Kaletra	SS		
300MG Per day		Irritability		Viread	C		
450MG Per day		Mania		Combivir	C	Glaxosmithkline	
10MG Twice				Oxandrin	C		
per day							
10MG Per day				Fosinopril	C		
				Furosemide	C	Glaxosmithkline	
25MG Per day				Metoprolol	C		
100MG Per day				Trazodone	C		
20MG Per day				Zyprexa	C		
20MG Per day				Paxil	C	Glaxosmithkline	
2PUFF Four				Combivent	C		
times per day							

2PUFF Twice	Flovent	C	Glaxosmithkline
per day			
100MG As	Viagra	C	
required			
	Lomotil	C	
	Creon	C	
	Multivitamin	C	
600MG Per day	N-Acetyl-L-Cysteine	C	Glaxosmithkline
250MG Per day	Alpha Lipoic Acid	C	
150MG Per day	Coenzyme Q10	C	
400MG Per day	Milk Thistle	C	
1G Per day	Carnitor	C	
1TAB Per day	Fiber	C	
	Omega 3	C	Glaxosmithkline

Date:11/23/04ISR Number: 4508304-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0490763A  
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorgasmia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Libido Decreased					
per day	6	MON		Wellbutrin	SS	Glaxosmithkline	ORAL
100MG Twice							
per day	2	YR		Lexapro	C		
				Lotensin	C	Glaxosmithkline	
				Vioxx	C		
				Neurontin	C		
				Robaxin	C		
				Oxycontin	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508306-2Report Type:Periodic  
Age:41 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0490791A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Scratch					
per day	DAY	Swelling Face		Depo-Provera	C		
		Urticaria		Xanax	C		

Date:11/23/04ISR Number: 4508307-4Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0490843A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Early Morning Awakening		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Per day	2 YR			Klonopin	C		
				Tegretol	C		
				Abilify	C		
				Elavil	C	Glaxosmithkline	
				Darvocet	C		

Date:11/23/04ISR Number: 4508308-6Report Type:Periodic  
Age:78 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0490845A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	12 WK			Synthroid	C	Glaxosmithkline	
				Flonase	C	Glaxosmithkline	
				Prevacid	C		
				Citrucel	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508309-8Report Type:Periodic  
Age:20 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0490969A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Agitation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Per day	1	MON					
		Insomnia					

Date:11/23/04ISR Number: 4508310-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0490983A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508311-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0490984A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	18	MON					

Date:11/23/04ISR Number: 4508312-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0491306A  
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hypertension		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	35	WK		Lipitor	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Premarin C  
 Calcium C

Date:11/23/04ISR Number: 4508313-XReport Type:Periodic  
 Age:30 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0491328A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	4	MON					
50MG As				Imitrex	C	Glaxosmithkline	ORAL
required							
5MG As				Zomig	C		ORAL
required							
SUBCUTANEOUS	6MG As			Zanaflex	C		ORAL
				Imitrex	C	Glaxosmithkline	
required							

Date:11/23/04ISR Number: 4508314-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491336A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day		Urticaria		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Provigil	C		
				Hormones	C		
				Vasotec	C		

Date:11/23/04ISR Number: 4508315-3Report Type:Periodic  
 Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491344A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Libido Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	5 YR						

Date:11/23/04ISR Number: 4508316-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0491345A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
150MG Twice							
per day	2 YR			Protease Inhibitor (Unspecified)	C		
				Interferon	C		

Date:11/23/04ISR Number: 4508317-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0491355A  
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK			Zoloft	C		

Date:11/23/04ISR Number: 4508318-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0491580A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508319-0Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491697A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2 WK Initial or Prolonged		Heart Rate Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508320-7Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491699A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
50MG Per day	3 YR	Hypoaesthesia Hypoaesthesia Oral		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508321-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491712A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Stevens-Johnson Syndrome		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508322-0Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491713A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	4 YR	Insomnia		Wellbutrin Zoloft	PS C	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508323-2Report Type:Periodic  
Age:37 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0491743A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Loss Of Consciousness					
per day	50	DAY		Alcohol	SS		
		Myalgia					
		Paraesthesia					
		Tremor					

Date:11/23/04ISR Number: 4508324-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0491772A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
		Euphoric Mood					

Date:11/23/04ISR Number: 4508325-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0491882A  
Age: YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Tinnitus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	1	MON		Advil	C	Glaxosmithkline	



150MG Twice		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1	MON		Buspar	C		
Date:11/23/04ISR Number: 4508330-XReport Type:Periodic			Company Report #US-GLAXOSMITHKLINE-A0491911A				
Age: YR	Gender:Male	I/FU:I					
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Dysuria		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Prostatic Disorder		Saw Palmetto	C		
per day							
Date:11/23/04ISR Number: 4508331-1Report Type:Periodic			Company Report #US-GLAXOSMITHKLINE-A0491915A				
Age: YR	Gender:Female	I/FU:I					
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Zoloft	C		
per day	5	WK		Seroquel	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508332-3Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491931A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 per day	10 YR	Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
		Weight Increased		Zoloft	C		
				Fosamax	C		

Date:11/23/04ISR Number: 4508333-5Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491994A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
4 YR		Bipolar I Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
		Confusional State		Levoxyl	C	Glaxosmithkline	
		Disturbance In Attention		Cytomel	C	Glaxosmithkline	
		Headache		Zyrtec	C	Glaxosmithkline	
		Insomnia		Allergy Shots	C		
		Logorrhoea		E-Vista	C		
		Memory Impairment		Ditropan	C		
		Obsessive-Compulsive Disorder		Vioxx	C		
		Pharmaceutical Product Complaint					

Date:11/23/04ISR Number: 4508335-9Report Type:Periodic  
Age:21 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0492013A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	7 DAY	Flatulence		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Pharmaceutical Product Complaint		Prozac	C		

Date:11/23/04ISR Number: 4508336-0Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492243A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Feeling Jittery		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 YR			Prinivil	C		
				Premarin	C		
				Estratest	C		

Date:11/23/04ISR Number: 4508337-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0492562A  
 Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Depression		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
300MG Unknown	1 MON			Doxycyclin	C		
		Increased Appetite					
		Irritability					
		Urticaria					

Date:11/23/04ISR Number: 4508338-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0492581A  
 Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Tremor		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
250MG							
Variable dose	7 MON			Celexa	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508339-6Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492598A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice per day	5 YR	Diarrhoea Headache Nausea Vomiting		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508340-2Report Type:Periodic  
Age:6 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492607A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day	4 DAY	Vomiting		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Strattera	C		

Date:11/23/04ISR Number: 4508341-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492683A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508342-6Report Type:Periodic  
Age:39 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0492886A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Drug Withdrawal Syndrome		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
3MG Per day		Headache		Klonopin	SS		
		Insomnia Nausea Palpitations Tremor					

Weight Decreased

Date:11/23/04ISR Number: 4508343-8Report Type:Periodic  
Age:57 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492889A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Pruritus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	4 WK	Rash					
		Urticaria		Zithromax	C		
				Proscar	C		

Date:11/23/04ISR Number: 4508344-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492890A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Lymphadenopathy		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 WK						
				Methocarbamol	C		
				Guaifenesin	C		
				Unknown Supplement	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508345-1Report Type:Periodic  
Age:11 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0492892A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2 YR			Adderall	C		

Date:11/23/04ISR Number: 4508346-3Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493453A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Epistaxis		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	8 MON			Flonase	SS	Glaxosmithkline	
		Hypertension		Hydrochlorothiazide	C		
		Palpitations		Premarin	C		
				Tiazac	C	Glaxosmithkline	
				Altace	C		
				Vioxx	C		
				Maxair	C		
				Advair	C	Glaxosmithkline	
				Nasonex	C		
				Synthroid	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508347-5Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493454A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
3 WK							

Date:11/23/04ISR Number: 4508348-7Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493472A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG	Per day	Myalgia		Tamoxifen	C		
				Melatonin	C		
				Antihistamine	C		

Date:11/23/04ISR Number: 4508349-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0493491A  
 Age:13 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea Medication Error Nausea Yawning		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508350-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0493575A  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hepatic Enzyme Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Twice			Buspar	C		
per day	5 YR			Glucophage	C		
				Synthroid	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vitamins C

Date:11/23/04ISR Number: 4508351-7Report Type:Periodic  
Age:57 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493578A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice	3 WK	Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day				Lortab	C		
				Soma	C		
				Xanax	C		
				Sleep Aid	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508352-9Report Type:Periodic  
Age:40 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0493579A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown	11 DAY	Urticaria		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
25MG Unknown				Paxil	C	Glaxosmithkline	ORAL
UNKNOWN	100MG Unknown			Toprol Xl	C		

Date:11/23/04ISR Number: 4508353-0Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493584A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508354-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493775A

Outcome Dose Duration PT Weight Increased Report Source Product Wellbutrin Sr Role PS Manufacturer Glaxosmithkline Route ORAL

Date:11/23/04ISR Number: 4508356-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0493916A  
Age:38 YR Gender:Female I/FU:I

Outcome Dose Duration PT Drug Ineffective Report Source Product Wellbutrin Sr Role PS Manufacturer Glaxosmithkline Route ORAL  
150MG Per day 1 MON

Date:11/23/04ISR Number: 4508357-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0493931A  
Age: Gender:Female I/FU:I

Outcome Dose Duration PT Rash Report Source Product Wellbutrin Role PS Manufacturer Glaxosmithkline Route ORAL  
100MG Twice per day 3 WK

Date:11/23/04ISR Number: 4508358-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0493942A  
Age: Gender:Female I/FU:I

Outcome Dose Duration PT Heart Rate Increased Insomnia Medication Error Report Source Product Wellbutrin Role PS Manufacturer Glaxosmithkline Route ORAL  
100MG Per day 1 MON



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508359-1Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493947A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2	WK		Wellbutrin	PS	Glaxosmithkline	ORAL
3	MON			Paxil	SS	Glaxosmithkline	ORAL
				Celebrex	C		
				Allegra	C		
				Multivitamin	C		
				Calcium Supplement	C		
				Vitamin E	C		
				Simvastatin	C		
				Prempro	C		
				Sleep Aid	C	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508360-8Report Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493948A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	4	MON		Wellbutrin	PS	Glaxosmithkline	ORAL
				Antihypertensive	C		

Date:11/23/04ISR Number: 4508361-XReport Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493952A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice				Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2	MON					
				Lisinopril	C		
				Diclofenac	C		
				Synthroid	C	Glaxosmithkline	
				Hctz	C		
				Xanax	C		
				Magnesium Oxide	C		
				Dhea	C		

Vitamins C  
Glucosamine C

Date:11/23/04ISR Number: 4508362-1Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493967A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vomiting		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3	WK					

Date:11/23/04ISR Number: 4508363-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493971A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
225MG Per day	1	MON					
		Fatigue		Wellbutrin	SS	Glaxosmithkline	ORAL
100MG Per day	2	MON					
		Palpitations		Lexapro	C		
		Vomiting		Oxizole	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508364-5Report Type:Periodic  
 Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493973A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Per day	1	MON		Prempro	C		
				Bextra	C		
				Oxycontin	C		
				Zanaflex	C		

Date:11/23/04ISR Number: 4508365-7Report Type:Periodic  
 Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0493976A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG		Disturbance In Attention					
Variable dose	62	WK		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
150MG Per day		Dizziness		Celexa	C		
		Euphoric Mood		Clarinx	C		
		Headache					
		Insomnia					
		Irritability					
		Mood Swings					
		Screaming					

Date:11/23/04ISR Number: 4508366-9Report Type:Periodic  
 Age:56 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0494062A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphagia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1	DAY		Estrogen	C		
		Feeling Hot					
		Rash					
		Rash Papular					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
		Anorexia		Wellbutrin	SS	Glaxosmithkline	ORAL
		Crying		Neurontin	C		
		Diarrhoea		Trileptal	C		
		Drug Ineffective		Klonopin	C		
		Hot Flush		Unspecified Inhaler	C		
		Migraine					
		Nervousness					
		Vision Blurred					
		Weight Decreased					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508372-4Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494247A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Twice		Dry Mouth		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	2 YR	Insomnia		Ambien	C		

Date:11/23/04ISR Number: 4508373-6Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494267A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	4 MON	Insomnia Nervousness Social Phobia					

Date:11/23/04ISR Number: 4508374-8Report Type:Periodic  
Age:20 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494270A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	11 MON	Weight Decreased		Yasmin	C		

Date:11/23/04ISR Number: 4508375-XReport Type:Periodic  
Age:59 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494280A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150TAB Twice		Vitamin D Deficiency		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

per day 1 YR

Vitamin D	C	
Risperdal	C	
Avandia	C	Glaxosmithkline
Axid	C	
Nexium	C	
Baby Aspirin	C	Glaxosmithkline
Calcitrol	C	
Provigil	C	
Effexor	C	

Date:11/23/04ISR Number: 4508376-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0494335A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Tremor					

Date:11/23/04ISR Number: 4508377-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0494471A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gynaecomastia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Famotidine	C		
				Protonix	C		

7 YR

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508378-5Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494482A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1MG Twice per day	1 MON	Alopecia Feeling Jittery		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508379-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494545A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
MON		Hypersensitivity Stomatitis Urticaria		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508380-3Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494680A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 WK	Weight Increased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Celexa	C		

Date:11/23/04ISR Number: 4508381-5Report Type:Periodic  
Age:17 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0494715A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day	3 YR	Blood Prolactin Increased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Zyprexa	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Diarrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
300 per day		Flatulence		Nexium	C		
		Pharmaceutical Product		Amaryl	C		
		Complaint		Plavix	C		
				Unspecified Drug	C		
				Allopurinol	C	Glaxosmithkline	
				Glucophage Xr	C		
				Topral	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Diarrhoea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Per day		Pharmaceutical Product		Zoloft	C		
		Complaint		Lorazepam	C		
				Benzaclin	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508384-0Report Type:Periodic  
 Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495037A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complex Partial Seizures		Wellbutrin	PS	Glaxosmithkline	ORAL
2 YR							

Date:11/23/04ISR Number: 4508385-2Report Type:Periodic  
 Age:40 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495047A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Erectile Dysfunction		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	1 MON			Atenolol	C		
				Lovastatin	C		
				Plavix	C		
				Lisinopril	C		
				Famotidine	C		
				Aspirin	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508386-4Report Type:Periodic  
 Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0495335A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Three		Tremor					
times per day							

Date:11/23/04ISR Number: 4508387-6Report Type:Periodic  
 Age:78 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0495336A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Unknown		Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508388-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495351A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1	MON					

Date:11/23/04ISR Number: 4508389-XReport Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495353A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	5	WK					

Date:11/23/04ISR Number: 4508390-6Report Type:Periodic  
Age:26 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495356A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
300MG Per day	11	MON					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508391-8Report Type:Periodic  
Age:13 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0495528A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	8	MON					

Date:11/23/04ISR Number: 4508392-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495529A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN				Effexor	SS		

Date:11/23/04ISR Number: 4508393-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495530A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain Dysmenorrhoea Menorrhagia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508394-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495552A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Unknown		Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Pharmaceutical Product Complaint					

Date:11/23/04ISR Number: 4508395-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0495574A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508396-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0495938A  
Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

150MG Three  
times per day 2 YR

Date:11/23/04ISR Number: 4508397-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0495962A  
Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Weight Decreased					

450MG Per day 7 MON

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508398-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495973A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoacusis		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Tinnitus					
per day	2 YR			Valtrex	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508399-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496025A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							

Date:11/23/04ISR Number: 4508400-6Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496033A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharyngolaryngeal Pain		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Throat Tightness					
per day	3 WK			Prozac	C		
				Estrace	C		
				Ambien	C		
				Tylenol	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508401-8Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496040A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2 YR						

Date:11/23/04ISR Number: 4508402-XReport Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496044A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Twice			Blood Prolactin Increased	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day							

Date:11/23/04ISR Number: 4508403-1Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496057A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3 DAY		Nervousness	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Thyroid Medication	C		

Date:11/23/04ISR Number: 4508404-3Report Type:Periodic  
Age:25 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496078A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Paraesthesia	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	6 WK						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508406-7Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496229A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day				Wellbutrin	C	Glaxosmithkline	ORAL
300MG per day							

Date:11/23/04ISR Number: 4508407-9Report Type:Periodic  
 Age:17 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496279A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1 MON			Ambien	C		
				Wellbutrin	C	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508408-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496281A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 YR	Stool Analysis Abnormal		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508409-2Report Type:Periodic  
 Age:41 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496295A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Arthralgia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	1 YR	Back Pain					

150MG Per day	Fall		Effexor Xr	C		ORAL
	Grand Mal Convulsion					
	Loss Of Consciousness					
	Neck Pain					
Date:11/23/04	ISR Number: 4508410-9	Report Type:Periodic	Company Report #US-GLAXOSMITHKLINE-A0496310A			
Age:	Gender:Male	I/FU:F				
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Anger		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
Date:11/23/04	ISR Number: 4508411-0	Report Type:Periodic	Company Report #US-GLAXOSMITHKLINE-A0496344A			
Age:34 YR	Gender:Female	I/FU:I				
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Tonsillitis		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	4	MON				
Date:11/23/04	ISR Number: 4508412-2	Report Type:Periodic	Company Report #US-GLAXOSMITHKLINE-A0496496A			
Age:	Gender:Female	I/FU:I				
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Claustrophobia		Paxil	PS	Glaxosmithkline	ORAL
	Headache		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
200MG In the						
morning	4	MON				
19-Aug-2005	12:44	PM				
Page: 2939						



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508413-4Report Type:Periodic  
Age:31 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496507A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Chest Discomfort		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day		Panic Reaction					
				Albuterol	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508414-6Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496634A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Per day	1 MON	Flushing		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Prempro	C		

Date:11/23/04ISR Number: 4508415-8Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496637A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Twice		Somnolence		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 YR			Effexor Xr	C		

Date:11/23/04ISR Number: 4508416-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496655A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 DAY	Feeling Abnormal		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Tinnitus		Strattera	C		
				Lorazepam	C		

Date:11/23/04ISR Number: 4508417-1Report Type:Periodic  
Age:25 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496658A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Twice		Eructation					
per day	2	Nausea		Seroquel	C		
				Trileptal	C		

Date:11/23/04ISR Number: 4508418-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496741A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphagia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Unknown							

Date:11/23/04ISR Number: 4508419-5Report Type:Periodic  
Age:42 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496756A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Mania					
100MG Twice		Tinnitus		Wellbutrin	SS	Glaxosmithkline	
per day	3			Klonopin	C		
				Lamictal	C	Glaxosmithkline	
300MG Per day	3			Lithium	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Ativan C  
 Buspar C  
 Inderal C

Date:11/23/04ISR Number: 4508420-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0496767A  
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG See							
dosage text	10 DAY						

Date:11/23/04ISR Number: 4508421-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0496784A  
 Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abdominal Pain Upper		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Anxiety		Wellbutrin	SS	Glaxosmithkline	ORAL
		Hypomania		Lamictal	SS	Glaxosmithkline	
		Insomnia					
		Palpitations					
		Tremor					

Date:11/23/04ISR Number: 4508422-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0496954A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day							

Date:11/23/04ISR Number: 4508423-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0496960A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Per day 9 MON	Pharyngolaryngeal Pain	Wellbutrin	PS	Glaxosmithkline	ORAL
		Protonix	C		
		Imitrex	C	Glaxosmithkline	
		Darvocet	C		

Date:11/23/04ISR Number: 4508424-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497085A  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice		Foreign Body Trauma		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day		Pharmaceutical Product					
		Complaint					

Date:11/23/04ISR Number: 4508425-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497093A  
 Age:8 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Hallucination		Wellbutrin	PS	Glaxosmithkline	ORAL
1 YR				Risperdal	C		ORAL
2MG At night							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508426-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497145A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 YR		Erythema		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Eye Irritation					
		Eyelid Oedema					
		Headache					
		Hypersensitivity					
		Rash Macular					

Date:11/23/04ISR Number: 4508427-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0497168A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pruritus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508428-6Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0497179A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Anorexia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Anxiety					
		Confusional State					
		Depression					
		Dizziness					
		Drug Withdrawal Syndrome					
		Insomnia					
		Muscle Tightness					
		Palpitations					
		Panic Attack					

Date:11/23/04ISR Number: 4508430-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0497373A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus Urticaria		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508431-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497603A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth Dysgeusia Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508432-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497741A  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1	YR		Benicar	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508433-XReport Type:Periodic  
 Age:41 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0497758A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100MG Twice		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1 MON	Euphoric Mood					
		Hypomania		Depakote	C		
		Insomnia		Lithium	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508434-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497770A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Alopecia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day							

Date:11/23/04ISR Number: 4508435-3Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497773A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day				Adderall	C		

Date:11/23/04ISR Number: 4508436-5Report Type:Periodic  
 Age:51 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0497786A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day	3 DAY	Headache		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Insomnia		Lipitor	C		

Date:11/23/04ISR Number: 4508437-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497800A  
 Age:35 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 MON	Insomnia					
		Migraine		Nexium	C		

Date:11/23/04ISR Number: 4508438-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497824A  
 Age:45 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day				Pravachol	C		

Date:11/23/04ISR Number: 4508439-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0497974A  
 Age:45 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1 YR			Glucovance	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508440-7Report Type:Periodic  
Age:66 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0497994A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	300MG per day		Anaphylactoid Reaction	Wellbutrin	PS	Glaxosmithkline	ORAL
Other	500MG per day		Angioneurotic Oedema	Levaquin	SS		
			Rash				

Date:11/23/04ISR Number: 4508441-9Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498162A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Irritable Bowel Syndrome	Wellbutrin	PS	Glaxosmithkline	ORAL
	150MG Twice per day	22 WK		Fosamax	C		
				Premarin	C		

Date:11/23/04ISR Number: 4508442-0Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498164A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Halitosis	Wellbutrin	PS	Glaxosmithkline	ORAL
	150MG Twice per day	4 MON					

Date:11/23/04ISR Number: 4508443-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498178A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Agitation	Wellbutrin	PS	Glaxosmithkline	ORAL
	200MG Twice						

per day

Toprol Xl C  
Nicotine Patch C Glaxosmithkline

Date:11/23/04ISR Number: 4508444-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498179A  
Age:53 YR Gender:Male I/FU:I

Outcome PT Report Source Product Role Manufacturer Route  
Dose Duration Fatigue Wellbutrin PS Glaxosmithkline ORAL  
150MCG Twice  
Sexual Dysfunction  
per day 12 DAY

Trazodone C  
Lipitor C

Date:11/23/04ISR Number: 4508445-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498339A  
Age:37 YR Gender:Male I/FU:I

Outcome PT Report Source Product Role Manufacturer Route  
Dose Duration Fatigue Wellbutrin Sr PS Glaxosmithkline ORAL  
100MG Per day  
Pharmaceutical Product  
Complaint  
Effexor Xr C

Date:11/23/04ISR Number: 4508446-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498343A  
Age: YR Gender:Male I/FU:I

Outcome PT Report Source Product Role Manufacturer Route  
Dose Duration Feeling Abnormal Wellbutrin Sr PS Glaxosmithkline ORAL  
150MG Per day 1 WK  
Vitamins C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Antioxidants C

Date:11/23/04ISR Number: 4508447-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498344A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508448-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498348A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

150MG Twice

per day 4 YR

Date:11/23/04ISR Number: 4508449-3Report Type:Periodic  
Age:38 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0498350A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL

Hospitalization - 15 MON  
Initial or Prolonged Grand Mal Convulsion

Tessalon Perles SS

Date:11/23/04ISR Number: 4508450-XReport Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498552A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

200MG Twice

per day

Date:11/23/04ISR Number: 4508451-1Report Type:Periodic  
Age:71 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498750A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
7 MON				Chemotherapy	C		

Date:11/23/04ISR Number: 4508452-3Report Type:Periodic  
Age:22 MON Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498772A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Faeces Discoloured		Wellbutrin Sr	PS	Glaxosmithkline	
100MG Per day 2 MON		Vomiting		Prevacid	C		
				Levsin	C		

Date:11/23/04ISR Number: 4508453-5Report Type:Periodic  
Age: YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0498856A

Outcome	PT
Other	Amnesia
	Dry Mouth
	Insomnia
	Panic Attack
	Petit Mal Epilepsy
	Restlessness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	14 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
			Thyroid Medication	C		
			Buspar	C		

Date:11/23/04ISR Number: 4508454-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498874A  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	17 WK			Spironolactone	SS		ORAL
4 WK				Synthroid	C	Glaxosmithkline	
				Vitamins	C		

Date:11/23/04ISR Number: 4508455-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499193A  
 Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	9 MON						

Date:11/23/04ISR Number: 4508456-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499210A  
 Age: YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -							
150MG Twice							

Initial or Prolonged  
per day 1 WK

Date:11/23/04ISR Number: 4508457-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499226A  
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day		Headache Insomnia Migraine					

Date:11/23/04ISR Number: 4508458-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499406A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day 3 MON		Headache Nausea		Prozac Aspirin	C C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508459-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499448A  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Twitching		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Sleep Disorder					
per day 4 MON		Weight Decreased		Buspar	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Cardizem Cd C Glaxosmithkline

Date:11/23/04ISR Number: 4508460-2Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499455A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Muscle Twitching		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	3 WK	Swelling Face					

Date:11/23/04ISR Number: 4508461-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499458A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
1 WK		Dizziness		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508462-6Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499627A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	1 YR			Lipitor	C		

Date:11/23/04ISR Number: 4508463-8Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499808A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	4 DAY	Anxiety		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508464-XReport Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499809A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	2	YR		Oral Contraceptives	C		

Date:11/23/04ISR Number: 4508465-1Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499839A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2	MON		Ambien	C		

Date:11/23/04ISR Number: 4508466-3Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499842A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	10	MON		Lisinopril	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508467-5Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499849A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Dizziness					
150MG Twice		Weight Gain Poor					
per day							

Date:11/23/04ISR Number: 4508468-7Report Type:Periodic  
Age:38 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0499860A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Stomach Discomfort		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	2 YR						

Date:11/23/04ISR Number: 4508469-9Report Type:Periodic  
Age:23 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499862A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decreased Appetite		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Twice		Depression					
per day		Sleep Disorder					

Date:11/23/04ISR Number: 4508470-5Report Type:Periodic  
Age:63 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499865A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Ineffective					
per day	6 WK						

Nausea

Xanax

C

Date:11/23/04ISR Number: 4508471-7Report Type:Periodic  
Age:27 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0499870A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Twice			Stool Analysis Abnormal	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day							

Date:11/23/04ISR Number: 4508472-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #04-00071

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day			Tinnitus	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Nystatin + Triamcinolone Acetonide	SS		
TOPICAL	.1%DS	Unknown					

Date:11/23/04ISR Number: 4508473-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500645A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Twice			Fatigue	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	1	YR					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508475-4Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500661A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	7	YR					
				Diazepam	C		
				Risperdal	C		
				Trazodone	C		

Date:11/23/04ISR Number: 4508476-6Report Type:Periodic  
 Age:19 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500662A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	3	WK					
		Rash		Wellbutrin	SS	Glaxosmithkline	ORAL
100MG Per day	0	DAY					

Date:11/23/04ISR Number: 4508477-8Report Type:Periodic  
 Age:32 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500670A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
Disability							
200MG Twice							
per day	3	MON					
				Zyprexa	C		
20MG Per day							

Date:11/23/04ISR Number: 4508478-XReport Type:Periodic  
 Age:47 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0500690A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

100MG Twice  
per day 6 MON  
Disturbance In Attention  
Drug Ineffective

Wellbutrin Sr PS Glaxosmithkline ORAL

Date:11/23/04ISR Number: 4508479-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0500695A  
Age: YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508480-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0500714A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

150MG Twice  
per day 8 WK  
Thinking Abnormal

Tamoxifen C  
Toprol Xl C

Date:11/23/04ISR Number: 4508481-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0500715A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

150MG Twice  
per day  
Zoloft C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508482-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500842A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain Headache		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508483-3Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500849A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cough		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice per day							

Date:11/23/04ISR Number: 4508484-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0500895A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cerebrovascular Accident		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
Hospitalization - 200MG Twice Initial or Prolonged per day							

Date:11/23/04ISR Number: 4508485-7Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #2004-00809

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
100MG Per day 15 DAY							
UNKNOWN							
Topamax C							

Date:11/23/04ISR Number: 4508486-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0500935A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	

Date:11/23/04ISR Number: 4508487-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0501052A  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Rash Macular					
per day	3 WK			Amoxicillin	C	Glaxosmithkline	
				Guaifen	C		

Date:11/23/04ISR Number: 4508488-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0501058A  
Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
300MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508489-4Report Type:Periodic  
 Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501060A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 MON	Constipation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508490-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501062A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	10 DAY	Joint Swelling		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Glyburide	C		
				Avandia	C	Glaxosmithkline	
				Diovan	C		
				Synthroid	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508491-2Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501066A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		Bedridden Drug Interaction		Wellbutrin Sr Pain Medication	PS C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508492-4Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501072A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 WK	Anxiety		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

300MG Per day	Drug Ineffective	Wellbutrin Xl	SS	Glaxosmithkline	ORAL
	Nausea	Benicar	C		
	Vomiting	Norvasc	C		
		Fish Oil	C		

Date:11/23/04ISR Number: 4508493-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0501103A  
 Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Headache		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Unknown						

Date:11/23/04ISR Number: 4508494-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0501174A  
 Age:63 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Nightmare		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice						
	Tremor					
per day	14 DAY		Nexium	C		

Date:11/23/04ISR Number: 4508495-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0501202A  
 Age:54 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Disability	Deafness		Wellbutrin	PS	Glaxosmithkline	
			Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day	2 MON					
			Concerta	C		
3 WK						



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Levoxyl C Glaxosmithkline

Date:11/23/04ISR Number: 4508496-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501228A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dry Mouth		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Unknown		Hyperhidrosis		Effexor	C		

Date:11/23/04ISR Number: 4508497-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501409A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Stool Analysis Abnormal		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508498-5Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501414A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice per day	4 DAY			Ritalin Concerta	C C		

Date:11/23/04ISR Number: 4508499-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501431A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypertension		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508500-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0501443A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Twice		Tremor					
per day				Zoloft	C		

Date:11/23/04ISR Number: 4508501-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0502027A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amenorrhoea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
3 MON							

Date:11/23/04ISR Number: 4508502-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0502028A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
Other							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508503-6Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502163A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/23/04ISR Number: 4508504-8Report Type:Periodic  
 Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502177A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
3 YR							

Date:11/23/04ISR Number: 4508505-XReport Type:Periodic  
 Age:71 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0502179A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	9 YR	Dry Skin		Atenolol	SS		
UNKNOWN				Enalapril	C		
				Synthroid	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508506-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502196A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Single							
dose	1 DAY						

Date:11/23/04ISR Number: 4508507-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502197A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR						

Date:11/23/04ISR Number: 4508508-5Report Type:Periodic  
Age: YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0502365A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
150MG Twice							
per day							

Date:11/23/04ISR Number: 4508509-7Report Type:Periodic  
Age:45 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0502373A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Grand Mal Convulsion					
150MG Twice							
per day	9 WK						
				Alcohol	SS		
				Zoloft	C		
200MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508510-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502563A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	30 DAY						

Date:11/23/04ISR Number: 4508511-5Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502583A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flatulence		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	2 YR			Synthroid	C	Glaxosmithkline	
				Xanax	C		
				Allegra D	C		
				Paxil	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508512-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502632A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Zyban	C	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508513-9Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502657A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flatulence		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	2 YR			Lipitor	C		
				Xanax	C		
				Clonidine	C		
				Hctz	C		

Date:11/23/04ISR Number: 4508514-0Report Type:Periodic  
Age:55 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0502715A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Unknown				Effexor	C		ORAL
150MG Twice							
per day	2	YR		Protonix	C		
UNKNOWN				Volmax	C	Glaxosmithkline	
UNKNOWN				Procrit	C		
UNKNOWN				Premarin	C		
UNKNOWN							

Date:11/23/04ISR Number: 4508515-2Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502866A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Dizziness					
per day	1	WK		Estrogens	C		
		Headache		Synthroid	C	Glaxosmithkline	
		Myalgia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508516-4Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502869A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1TAB Twice		Abnormal Dreams		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	5	DAY					
150MG Per day	5	DAY		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Dry Mouth		Arimidex	C		
		Gingival Pain		Propranolol	C		
		Glossodynia		Hyzaar	C		
		Headache		Diclofenac	C		
		Mood Altered		Zometa	C		
		Nausea		Aciphex	C		
		Oedema Peripheral		Clonidine	C		
				Synthroid	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508517-6Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502876A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown		Nausea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508518-8Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502878A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Dizziness		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	5	DAY					
		Dry Mouth					
		Headache		Prilosec	C	Glaxosmithkline	
		Nausea					
		Vision Blurred					

Date:11/23/04ISR Number: 4508519-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0502889A  
Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Wellbutrin Xl	SS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508520-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0502924A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508521-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0502958A  
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day 1 WK							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508522-XReport Type:Periodic  
Age:10 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502963A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	18 MON	Emotional Disorder	Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508523-1Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502984A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	100MG Per day	1 WK	Vision Blurred	Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508524-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503162A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	1 YR		Vaginal Mycosis	Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508525-5Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503198A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Life-Threatening	1 WK	Grand Mal Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
Disability			Skin Laceration	Prozac	C		
Other				Ibuprofen	C	Glaxosmithkline	
				Claritin	C		
				Nicotine Patch	C	Glaxosmithkline	
				Axid	C		
				Vicodin	C		
				Vitamin B Complex	C		
				Trazodone	C		
				Klonopin	C		
				Ambien	C		

Date:11/23/04ISR Number: 4508526-7Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503221A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1	DAY	Blood Pressure Increased	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
			Rash	Blood Pressure Medication	C		
			Tinnitus	Cholesterol	C		
				Bc	C	Glaxosmithkline	ORAL
				Sinus Medication	C		

Date:11/23/04ISR Number: 4508527-9Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503232A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	5	DAY	Affect Lability	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
			Depression	Biaxin	C		
			Dyspnoea	Albuterol	C	Glaxosmithkline	
			Muscle Spasms	Aciphex	C		
			Swelling Face				
			Swollen Tongue				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508528-0Report Type:Periodic  
 Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503444A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Feeling Of Relaxation Personality Change		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508529-2Report Type:Periodic  
 Age:55 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0503447A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	4 YR	Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Nexium	C		

Date:11/23/04ISR Number: 4508530-9Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503449A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 WK	Agitation Anxiety Depression Nausea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Allegra	C		

Date:11/23/04ISR Number: 4508531-0Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503450A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Tachycardia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508532-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0503461A  
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2	WK					

Date:11/23/04ISR Number: 4508533-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0503508A  
Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	10	DAY					

Date:11/23/04ISR Number: 4508534-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0503510A  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Pruritus					
per day	1	WK		Levaquin	SS		
				Plavix	SS		
UNKNOWN							



Other Convulsion Wellbutrin PS Glaxosmithkline ORAL

150MG See

dosage text 8 DAY

Date:11/23/04ISR Number: 4508539-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0503831A  
Age:16 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tongue Eruption		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2	WK		None	C		

Date:11/23/04ISR Number: 4508540-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0503845A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Twice		Dry Mouth					
per day	2	YR					

Date:11/23/04ISR Number: 4508541-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0503852A  
Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3	WK					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508542-5Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504063A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day		Dysarthria		Wellbutrin	PS	Glaxosmithkline	ORAL
				Depakote	C		
				Ambien	C		
				Luvox	C		

Date:11/23/04ISR Number: 4508543-7Report Type:Periodic  
 Age:50 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504069A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	1 DAY	Muscle Twitching		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508544-9Report Type:Periodic  
 Age:24 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504071A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	3 YR	Dry Mouth		Wellbutrin	PS	Glaxosmithkline	ORAL
		Halitosis					
		Libido Increased		Ambien	C		

Date:11/23/04ISR Number: 4508545-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504225A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Overdose		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508546-2Report Type:Periodic  
Age:28 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504301A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ejaculation Failure		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK			None	C		

Date:11/23/04ISR Number: 4508547-4Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504573A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypothermia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Twice							
per day				Avelox	C		
				Sinus Medication	C		

Date:11/23/04ISR Number: 4508548-6Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504789A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
2 MON							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508549-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504810A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination		Wellbutrin No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508550-4Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0504832A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
8 DAY		Depressed Level Of Consciousness		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508551-6Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504888A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Twice per day	61 WK	Abdominal Pain Upper Menstrual Disorder Nausea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
1 WK				Wellbutrin Xl	SS	Glaxosmithkline	ORAL
				Levoxyl	C	Glaxosmithkline	
				Atenolol	C		
				Soma	C		
				Imitrex	C	Glaxosmithkline	
				Ibuprofen	C	Glaxosmithkline	
				Tylenol W/ Codeine	C		

Date:11/23/04ISR Number: 4508552-8Report Type:Periodic  
Age:32 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505099A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Agitation Wellbutrin Sr PS Glaxosmithkline ORAL

150MG Twice

per day 4 DAY

Date:11/23/04ISR Number: 4508553-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505139A  
Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mixed Connective Tissue		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Disease					
per day	4	YR					

Date:11/23/04ISR Number: 4508554-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505262A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort Dyspnoea Eye Pruritus Hypersensitivity Nasal Discomfort Ocular Hyperaemia Pruritus Sneezing Somnolence Throat Irritation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508555-3Report Type:Periodic  
Age:66 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505382A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	10 DAY	Dry Mouth Headache		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Klonopin	C		
				Lopid	C		
				Timoptic	C		
				Xalatan	C		

Date:11/23/04ISR Number: 4508556-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505418A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice per day	3 YR	Breast Discharge		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508557-7Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505419A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	1 YR	Feeling Abnormal Increased Appetite Insomnia Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
				Detrol La	C		
				Yasmin	C		

Date:11/23/04ISR Number: 4508558-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505455A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Crying

Wellbutrin Sr  
Zoloft

PS  
C

Glaxosmithkline

ORAL

12 YR

Date:11/23/04ISR Number: 4508559-0Report Type:Periodic  
Age:60 YR Gender:Female I/FU:F

Company Report #2004-01334

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Twice		Abdominal Distension		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	1	Flatulence					
UNKNOWN		Frequent Bowel Movements		Zoloft	C		
		Headache					
		Nausea					

Date:11/23/04ISR Number: 4508560-7Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505564A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Per day	6	Blood Cholesterol		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Increased		Sulfasalazine	C		
		Blood Triglycerides					
		Increased					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508561-9Report Type:Periodic  
Age:66 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505578A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day				Xanax	C		
				Coumadin	C	Glaxosmithkline	
				Cosaar	C		

Date:11/23/04ISR Number: 4508562-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505609A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Per day	5 MON						
		Myalgia		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
150MG Per day	5 MON						
				Buspar	C		
				Herbal Supplements	C		
				Fish Oil	C		
				Vitamin Supplements	C		

Date:11/23/04ISR Number: 4508563-2Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505933A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anaemia		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
150MG Twice		Leukopenia					
per day	9 MON			Mvi	C		

Date:11/23/04ISR Number: 4508564-4Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505939A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	50MG Per day	2 WK	Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
					Zantac	C	Glaxosmithkline	
					Prilosec	C	Glaxosmithkline	
					Topamax	C		
					Zoloft	C		
					Allegra	C		

Date:11/23/04ISR Number: 4508565-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505972A  
 Age:28 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Twice		Fatigue		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
	per day	3 WK	Rash Pruritic					

Date:11/23/04ISR Number: 4508566-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505974A  
 Age:36 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	100MG Per day	3 WK	Swollen Tongue		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
					No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508567-XReport Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505984A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
75MG Per day	6 YR			Chemotherapy	C		
				Effexor	C		
				Tamoxifen	C		

Date:11/23/04ISR Number: 4508568-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506015A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Wellbutrin Sr	PS	Glaxosmithkline	
2 WK		Urticaria					

Date:11/23/04ISR Number: 4508569-3Report Type:Periodic  
Age:13 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506125A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bulimia Nervosa		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Per day		Therapeutic Response		Paxil Cr	SS	Glaxosmithkline	ORAL
37.5MG Per day		Decreased					

Date:11/23/04ISR Number: 4508570-XReport Type:Periodic  
Age:10 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506221A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Unknown		Constipation					
		Visual Disturbance					

Date:11/23/04ISR Number: 4508571-1Report Type:Periodic  
Age:37 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506226A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
400MG Twice							
per day	5	WK		Zoloft	C		
				Abilify	C		

Date:11/23/04ISR Number: 4508572-3Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506227A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2	WK		No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508573-5Report Type:Periodic  
Age:65 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506375A

Outcome	PT
Blood Pressure Increased	
Tinnitus	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Visual Acuity Reduced

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	6 MON		Wellbutrin	PS	Glaxosmithkline	ORAL
2 YR			Zoloft	C		
			Combipatch	C		

Date:11/23/04ISR Number: 4508574-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506376A  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Twitching		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice				Strattera	C		
per day				Armour Thyroid	C		

Date:11/23/04ISR Number: 4508575-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506484A  
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypotrichosis		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice				Prevacid	C		
per day	8 YR			Estrace	C		
				Ziac	C		
				Bextra	C		
				Guiafenesisin-Dm	C		

Date:11/23/04ISR Number: 4508576-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506509A  
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Per day		Insomnia		Effexor Xr	SS		ORAL
150MG Per day	2 YR			Protonix	C		
				Yasmin	C		

Date:11/23/04ISR Number: 4508577-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506519A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508578-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506520A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Insomnia					
per day	1 WK	Night Sweats		Zoloft	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508579-6Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0506579A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/23/04ISR Number: 4508580-2Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506812A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	4 YR						
		Drug Withdrawal Syndrome		Synthroid	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508581-4Report Type:Periodic  
Age:81 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506858A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vomiting		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK						
				Norvasc	C		
				Lisinopril	C		
				Atenolol	C		
				Clonidine	C		
				Paroxetine	C	Glaxosmithkline	
				Plavix	C		
				Pepcid	C		
				Nitro Patch	C	Glaxosmithkline	
				Insulin	C		

Date:11/23/04ISR Number: 4508582-6Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506922A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Twice	Restlessness	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	6 MON	Oral Contraceptive	C		

Date:11/23/04ISR Number: 4508583-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506928A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Blood Pressure Increased					
per day		Heart Rate Increased					

Date:11/23/04ISR Number: 4508584-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506941A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Dyspnoea		Inh	C	Glaxosmithkline	
		Yawning					

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Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508585-1Report Type:Periodic  
Age:47 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0506949A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Rectal Haemorrhage		Zyrtec	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508586-3Report Type:Periodic  
Age:33 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0506958A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	55 DAY	Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Stomach Discomfort		No Concurrent Medication	C		
		Urticaria					

Date:11/23/04ISR Number: 4508587-5Report Type:Periodic  
Age:39 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0506966A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Per day	2 WK	Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Nasopharyngitis		Concerta	C		
		Nausea					
		Skin Wrinkling					

Date:11/23/04ISR Number: 4508588-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507162A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Headache		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day		Nausea					
		Tremor					

Date:11/23/04ISR Number: 4508589-9Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507169A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day				Celexa	C		
				Lisinopril	C		
				Wellbutrin Xl	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508590-5Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507189A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hot Flush		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Hyperhidrosis					
per day	2 DAY			Vitamins	C		

Date:11/23/04ISR Number: 4508591-7Report Type:Periodic  
Age:53 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507196A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Psychomotor Hyperactivity		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Per day	2 DAY			Sam-E	C		

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Date:11/23/04ISR Number: 4508592-9Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507263A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	3 WK	Chest Pain Dizziness Pruritus Rash Urticaria		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508593-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507273A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Twice per day	8 MON	Drug Screen Positive		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Seroquel	C		

Date:11/23/04ISR Number: 4508594-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0507282A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Per day	30 MON	Depression Purging		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508595-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507283A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypersensitivity		Wellbutrin Sr	PS	Glaxosmithkline	

Date:11/23/04ISR Number: 4508596-6Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507454A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menstrual Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day				Alprazolam	C		

Date:11/23/04ISR Number: 4508597-8Report Type:Periodic  
Age:39 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0507562A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Visual Disturbance		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3 WK						

Date:11/23/04ISR Number: 4508598-XReport Type:Periodic  
Age:51 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0507695A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	4 DAY	Emotional Disorder					

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Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508599-1Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507907A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Suicidal Ideation		Bupropion	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	4	WK					

Date:11/23/04ISR Number: 4508600-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507938A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diplopia Dissociation Dizziness Vision Blurred		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508601-7Report Type:Periodic  
Age:62 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508032A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Plavix	SS		ORAL
				Xanax	C		ORAL
15	YR						

Date:11/23/04ISR Number: 4508602-9Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508052A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menstrual Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3	WK					

Date:11/23/04ISR Number: 4508603-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508085A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	3 MON	Fluid Retention		Synthroid	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508604-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508210A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
8 DAY		Dry Mouth		Cenestin	C		
UNKNOWN		Feeling Abnormal					
		Headache					
		Photophobia					
		Therapeutic Response					
		Unexpected					

Date:11/23/04ISR Number: 4508605-4Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508353A

Outcome	PT
	Agitation
	Anger

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Freedom Of Information (FOI) Report

Insomnia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 WK		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508606-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508428A  
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
20 MON		Cataract		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508607-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508433A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Twice per day	2 YR	Stool Analysis Abnormal		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Prozac C

Date:11/23/04ISR Number: 4508608-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508459A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 WK	Increased Appetite		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Weight Increased		Paxil	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508609-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508463A  
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Myalgia		Amitriptyline	C		
		Neck Pain		Perphenazine	C	Glaxosmithkline	
				Lamictal	C	Glaxosmithkline	
				Zoloft	C		
				Amitriptyline	C		

Date:11/23/04ISR Number: 4508610-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508566A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508611-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508745A  
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Ineffective					
per day	3 YR			Bupropion Hydrochloride	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508612-1Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508768A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia Parosmia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508613-3Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508786A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscular Weakness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Unknown	2 WK			Botox Injection	SS		
				Primidone	C		
				Diazepam	C		
				Oxycodone	C		

Date:11/23/04ISR Number: 4508614-5Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508959A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3 WK						

Date:11/23/04ISR Number: 4508615-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508978A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Migraine		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Five		Overdose					
times per day	2 YR			Cordarone	C		
				Ativan	C		

Neurontin C  
Coumadin C Glaxosmithkline

Date:11/23/04ISR Number: 4508616-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508982A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hunger		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150TAB Per		Weight Increased					
day	8	MON		Prozac	C		
				Vit B12	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508617-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0509131A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	

Date:11/23/04ISR Number: 4508618-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0509236A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	9	MON		Estrace	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Provera C  
 Prenatal Vitamins C

Date:11/23/04ISR Number: 4508619-4Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509246A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Twice			Hepatic Enzyme Increased	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	6	WK					
				Nabumetone	C	Glaxosmithkline	
				Prednisone	C		

Date:11/23/04ISR Number: 4508620-0Report Type:Periodic  
 Age:55 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509446A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3	MON	Blood Pressure Increased	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Asa	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508621-2Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509454A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Drug Ineffective	Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508622-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509550A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Asthenia	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
			Headache	Wellbutrin Xl	C	Glaxosmithkline	ORAL
				Zoloft	C		

Date:11/23/04ISR Number: 4508623-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509566A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Depressed Mood		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Lacrimation Increased		Klonopin	C		ORAL
.5MG Three							
times per day							
75MG At night				Effexor	C		ORAL

Date:11/23/04ISR Number: 4508624-8Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509624A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Ineffective					
per day	2	WK		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
300MG Per day	1	WK		Yasmin	C		
		Nausea		Metformin	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508626-1Report Type:Periodic  
Age:53 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509653A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	1 YR	Pruritus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Bupropion Hydrochloride	SS	Glaxosmithkline	
				Niacin	C		
				Crestor	C		
				Lanoxin	C	Glaxosmithkline	
				Coumadin	C	Glaxosmithkline	
				Atenolol	C		

Date:11/23/04ISR Number: 4508627-3Report Type:Periodic  
Age:49 YR Gender:Female I/FU:F

Company Report #2004-01832

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day		Drug Withdrawal Headache		Bupropion	SS	Glaxosmithkline	ORAL
		Headache Pharmaceutical Product Complaint					

Date:11/23/04ISR Number: 4508628-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509781A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Unknown	YR	Drug Exposure During Pregnancy Drug Exposure Via Breast Milk		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508629-7Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509817A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2	MON					
		Nightmare		Lorazepam	C		
		Thyroid Function Test		Donnatal	C		
		Abnormal		Synthroid	C	Glaxosmithkline	
				Nexium	C		

Date:11/23/04ISR Number: 4508630-3Report Type:Periodic  
Age:59 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0509819A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Jittery		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1	MON					
		Insomnia		Advair	C	Glaxosmithkline	
				Serevent	C	Glaxosmithkline	
				Celebrex	C		
				Allegra	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508631-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509964A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
7	WK						

Date:11/23/04ISR Number: 4508632-7Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510010A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	6 DAY	Medication Error		Zyban	SS	Glaxosmithkline	ORAL
150MG Per day	6 DAY	Psychomotor Hyperactivity		No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508633-9Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #2004-01906

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Bupropion	PS	Glaxosmithkline	ORAL
150MG Per day				Multivitamin	C		ORAL

Date:11/23/04ISR Number: 4508635-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510194A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vomiting		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 DAY			No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508636-4Report Type:Periodic  
Age:70 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0056952A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	11 DAY	Bruxism		Elavil	C	Glaxosmithkline	
		Feeling Jittery		Zoloft	C		
		Nervousness		Paxil	C	Glaxosmithkline	
				Pepcid	C		
				Antacids	C		
				Tylenol	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508638-8Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0365399A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Feeling Hot					

Date:11/23/04ISR Number: 4508639-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0371708A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2 WK			Celexa	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508640-6Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0381524A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	1 YR		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Oral Contraceptive	C		ORAL
				Asthma Medication	C		
UNKNOWN							

Date:11/23/04ISR Number: 4508641-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0411787A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Twice			Wellbutrin	PS	Glaxosmithkline	ORAL
	per day	2 MON					

Date:11/23/04ISR Number: 4508642-XReport Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0441170A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion					

Date:11/23/04ISR Number: 4508643-1Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0441171A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Rash					

Date:11/23/04ISR Number: 4508644-3Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510208A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anger		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Disorientation					
per day	31 DAY	Disturbance In Attention		Wellbutrin Xl	SS	Glaxosmithkline	
		Fatigue		Metformin	C		
		Irritability		Oral Contraceptives	C		
		Mood Altered					
		Nausea					

Date:11/23/04ISR Number: 4508645-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510311A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Bupropion	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508646-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510409A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice				No Concurrent Medication	C		
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508647-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510526A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Unknown				No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508648-0Report Type:Periodic  
Age:16 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510536A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Ambien	C		
per day	2 YR						

Date:11/23/04ISR Number: 4508649-2Report Type:Periodic  
Age:15 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510719A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	1 MON	Nervousness		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Tremor		Zyprexa	C		

Date:11/23/04ISR Number: 4508650-9Report Type:Periodic  
Age:65 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510730A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disturbance In Attention		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Twice		Mood Swings		Seroquel	C		
per day	2 YR	Psychomotor Hyperactivity					

Date:11/23/04ISR Number: 4508651-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510738A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Therapeutic Response		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Two		Decreased					
times per							
week							

Date:11/23/04ISR Number: 4508652-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510751A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Weight Increased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
300MG Per day				Celexa	C		

Date:11/23/04ISR Number: 4508653-4Report Type:Periodic  
Age:61 YR Gender:Female I/FU:F

Company Report #2004-02103

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Depression		Bupropion			
		Emotional Disorder		Hydrochloride	PS	Glaxosmithkline	ORAL
100MG Per day				Wellbutrin Sr	C	Glaxosmithkline	ORAL
100MG Per day	5 YR						



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508654-6Report Type:Periodic  
 Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510907A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Ineffective					
per day	1 YR	Sexual Dysfunction		Birth Control Pill	C		ORAL

Date:11/23/04ISR Number: 4508655-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511275A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	7 DAY	Headache		Percocet	C		
				Tylenol Pm	C		

Date:11/23/04ISR Number: 4508656-XReport Type:Periodic  
 Age:25 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0511435A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr	PS	Glaxosmithkline	
Other		Grand Mal Convulsion					

Date:11/23/04ISR Number: 4508657-1Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511460A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Drug Ineffective					
per day							

Date:11/23/04ISR Number: 4508658-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0511561A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Increased Tendency To		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
6	MON	Bruise Irritability					

Date:11/23/04ISR Number: 4508659-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0511573A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anhedonia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
350MG Per day		Anxiety Tinnitus		Benzodiazepam Effexor Xr Beconase Klonopin	C C C C	Glaxosmithkline	NASAL

Date:11/23/04ISR Number: 4508660-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0511574A

Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Unknown	MON						

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150MG Twice		Arthralgia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	1	Rash					
			WK				

Date:11/23/04ISR Number: 4508667-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0512032A  
Age:11 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Unknown		Menorrhagia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508668-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0512067A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation Therapeutic Response Decreased		Wellbutrin	PS	Glaxosmithkline	

Date:11/23/04ISR Number: 4508669-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0512073A  
Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective Pharmaceutical Product					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Complaint

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	4 YR		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
			Synthroid	C	Glaxosmithkline	
			Allegra	C		

Date:11/23/04ISR Number: 4508670-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0512086A  
 Age:14 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day Other	2 MON	Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508671-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0512231A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
0 DAY		Pruritus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	6 WK			Wellbutrin Xl	SS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508672-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0512232A  
 Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Liver Function Test Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508673-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512337A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression Pharmaceutical Product Complaint		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508674-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512387A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Per day							

Date:11/23/04ISR Number: 4508675-3Report Type:Periodic  
Age:74 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512395A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
2	WK			Wellbutrin	SS	Glaxosmithkline	ORAL
2	WK			Valium	C		
				Remeron	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508676-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512416A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Drug Ineffective	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	1 WK		Dry Mouth				
				Fluoxetine	C		
				Lipitor	C		
				Diazepam	C		
				Metoprolol	C		

Date:11/23/04ISR Number: 4508677-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512420A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Insomnia	Wellbutrin	PS	Glaxosmithkline	ORAL
per day							

Date:11/23/04ISR Number: 4508678-9Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512621A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Photosensitivity Reaction	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
			Urticaria	Bupropion	SS	Glaxosmithkline	ORAL
				Xanax	C		

Date:11/23/04ISR Number: 4508679-0Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512633A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Pruritus	Wellbutrin Sr	PS	Glaxosmithkline	ORAL

per day		Rash					
TOPICAL				Ortho-Evra		C	
Date:11/23/04ISR Number: 4508680-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0512636A							
Age:	Gender:	I/FU:F					
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
Other							
Date:11/23/04ISR Number: 4508681-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0512881A							
Age:	Gender:Female	I/FU:I					
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Rash					
per day				Allegra		C	
Date:11/23/04ISR Number: 4508682-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0512965A							
Age:	Gender:Male	I/FU:I					
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nervousness		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Weight Decreased		Xanax	SS		
				Effexor Xr	C		
6	MON						



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508683-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512966A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Tremor		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/23/04ISR Number: 4508684-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513047A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Unknown	22 WK						

Date:11/23/04ISR Number: 4508685-6Report Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513051A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gynaecomastia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							
				Lisinopril	C		
				Rhinocort	C		
				Protonix	C		

Date:11/23/04ISR Number: 4508686-8Report Type:Periodic  
Age:41 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513268A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Tinnitus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Zestril C  
 Protonix C  
 Trazodone C  
 Klonopin C  
 Zocor C

Date:11/23/04ISR Number: 4508687-XReport Type:Periodic  
 Age:41 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513274A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorgasmia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Twice		Erectile Dysfunction					
per day	1 YR			Effexor	C		
				Clonazepam	C		
				Atenolol	C		
				Flomax	C		
				Morphine Sulfate	C	Glaxosmithkline	
				Levoxyl	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508688-1Report Type:Periodic  
 Age:24 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513294A

Outcome	PT
Other	Hallucination
	Paranoia
	Therapeutic Response
	Unexpected

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Weight Decreased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	1 YR		Wellbutrin	PS	Glaxosmithkline	ORAL
			No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508689-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0513573A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice per day	54 MON	Drug Screen False Positive					

Date:11/23/04ISR Number: 4508690-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0513616A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK	Rash Pruritic Urticaria		No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508693-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0513652A  
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - 150MG Twice Initial or Prolonged per day	2 YR	Thrombocytopenia		Tegretol	C		

Date:11/23/04ISR Number: 4508694-7Report Type:Periodic  
Age:26 YR Gender:Female I/FU:F

Company Report #2004-02446

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Unknown	31 DAY	Abdominal Distension Abdominal Pain		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
UNKNOWN	10MG Twice per day	Arthralgia Diarrhoea		Bentyl	C		
UNKNOWN		Muscle Spasms		Singular	C		
UNKNOWN	40MG Unknown	Pharmaceutical Product Complaint Pyrexia		Ortho Novum Nexium	C C		
UNKNOWN		Vomiting		Lomotil	C		
UNKNOWN	50MG Per day			Topamax	C		
400MG Twice per day				Chromium	C		
UNKNOWN	37.5U Unknown			Phentermine	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508698-4Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513792A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75MG Three times per day	1 WK	Constipation		Wellbutrin Wellbutrin	PS SS	Glaxosmithkline Glaxosmithkline	ORAL ORAL
				Fosamax	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508699-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513804A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Rash Pruritic		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 WK			No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508700-XReport Type:Periodic  
 Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513812A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Eye Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	5 DAY			Allegra	C		

Date:11/23/04ISR Number: 4508701-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513817A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3 WK	Muscle Twitching		Wellbutrin	PS	Glaxosmithkline	ORAL
				Seroquel	C		

Date:11/23/04ISR Number: 4508702-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513832A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75MG Per day		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
		Urticaria		Lexapro	C		

Date:11/23/04ISR Number: 4508703-5Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513834A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3	WK		No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508704-7Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514148A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3	WK		Prometrium	C		
				Estradiol	C		

Date:11/23/04ISR Number: 4508705-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514275A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508706-0Report Type:Periodic  
Age:23 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514339A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	3 DAY	Therapeutic Response		Wellbutrin	PS	Glaxosmithkline	ORAL
		Unexpected		Vistaril	C		
				Unknown Medication	C		

Date:11/23/04ISR Number: 4508707-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514467A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Unknown	Hypersensitivity		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Pruritus					

Date:11/23/04ISR Number: 4508708-4Report Type:Periodic  
Age:73 YR Gender:Female I/FU:F

Company Report #2004-02596

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Single	Ear Pruritus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
	dose	Erythema					
	500MG Single	Feeling Hot		Naproxen	SS		ORAL
	dose	Pruritus					
	600MG Single	Pruritus Generalised		Mucinex	SS		ORAL
	dose	Rash Macular					
	UNKNOWN			Zovirax	C	Glaxosmithkline	
				Aspirin	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508713-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514525A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	
Other		Mania					
150MG Twice							
per day				Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day	2	WK		Geodon	C		
				Depakote	C		

Date:11/23/04ISR Number: 4508714-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514656A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Celexa	SS		
UNKNOWN							

Date:11/23/04ISR Number: 4508715-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514665A  
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	5	WK		No Concurrent Medication	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508716-3Report Type:Periodic  
 Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514693A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Per day	2 WK			Wellbutrin	PS	Glaxosmithkline	ORAL
		Haematochezia					
		Nausea					

Date:11/23/04ISR Number: 4508717-5Report Type:Periodic  
 Age:29 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0514845A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	133 DAY			Bupropion	PS	Glaxosmithkline	ORAL
Other		Grand Mal Convulsion					

Date:11/23/04ISR Number: 4508718-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514867A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Twice				Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day		Insomnia					

Date:11/23/04ISR Number: 4508719-9Report Type:Periodic  
 Age:29 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514883A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Per day	4 DAY			Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Tinnitus					
				No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508720-5Report Type:Periodic  
 Age:26 YR Gender:Female I/FU:I

Company Report #2004-02598

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia Eye Pruritus Parosmia		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508721-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0515129A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction Libido Increased		Wellbutrin Sr Klonopin Cholesterol Reducing Agent	PS C C	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508722-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0515142A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal		Wellbutrin Wellbutrin	PS SS	Glaxosmithkline Glaxosmithkline	ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508723-0Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515144A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MGK Twice		Chest Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		Heart Rate Irregular					
		Hot Flush		Birth Control Pill	C		

Date:11/23/04ISR Number: 4508724-2Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515162A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 MON	Hypertension		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Intervertebral Disc		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
per day	5 YR	Protrusion					
		Lung Infection		Advair	C	Glaxosmithkline	
		Weight Increased		Albuterol	C	Glaxosmithkline	
				Buspar	C		
				Theophylline	C		
				Flexeril	C		
				Klonopin	C		
				Lisinopril	C		
				Trazodone	C		
				Clarinex	C		
				Allegra	C		
				Benadryl	C	Glaxosmithkline	
				Tavist	C		
				Ortho Tri-Cyclen	C		

Date:11/23/04ISR Number: 4508725-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #2004-02699

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypersensitivity		Bupropion			

150MG Per day	Stomach Discomfort	Hydrochloride	PS	Glaxosmithkline	ORAL	
Date:11/23/04ISR Number: 4508726-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0515387A						
Age:75 YR Gender:Male I/FU:I						
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
150MG Per day	1 YR	Dizziness	Wellbutrin	PS	Glaxosmithkline	ORAL
		Therapeutic Response	Hytrin	C		
		Unexpected	Lescol	C		
			Prinivil	C		

Date:11/23/04ISR Number: 4508727-8Report Type:Periodic Company Report #2004-02824						
Age: Gender:Male I/FU:F						
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
150MG Twice		Agitation	Bupropion Sr	PS	Glaxosmithkline	ORAL
per day		Therapeutic Response				
		Decreased				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508728-XReport Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515598A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	1 MON	Tinnitus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508729-1Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0515686A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dyspnoea Yawning		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508730-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515725A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Unknown		Diarrhoea Nausea Vomiting		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508731-XReport Type:Periodic  
 Age:61 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0515733A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	16 DAY	Erythema Gingival Pain Lip Blister Mouth Ulceration		Wellbutrin Sr Toprol Isordil	PS C C	Glaxosmithkline	ORAL

Pain In Extremity

Niaspan

C

Zetia

C

Date:11/23/04ISR Number: 4508732-1Report Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0515759A

Age:36 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Depression		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	4 DAY			No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508734-5Report Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0515831A

Age: Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 YR		Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Unspecified Medications	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508735-7Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515849A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropion	PS	Glaxosmithkline	ORAL
150MG Twice		Therapeutic Response					
per day		Decreased					
150MG Twice				Wellbutrin	C	Glaxosmithkline	ORAL
per day	YR						

Date:11/23/04ISR Number: 4508736-9Report Type:Periodic  
 Age:79 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0515905A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Drug Ineffective					
per day	10 WK	Insomnia					
				Blood Pressure Medication	C		
				Unknown Medication	C		

Date:11/23/04ISR Number: 4508737-0Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515908A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr	PS	Glaxosmithkline	NASAL
		Medication Error					

Date:11/23/04ISR Number: 4508738-2Report Type:Periodic  
 Age:77 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515910A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	1 DAY	Insomnia					

Coreg	C	Glaxosmithkline
Proscar	C	
Neurontin	C	
Furosemide	C	Glaxosmithkline
Monopril	C	
Potassium	C	
Lanoxin	C	Glaxosmithkline
Nitrodur Patch	C	Glaxosmithkline
Insulin	C	
Hemocyte	C	
Amitriptyline	C	

Date:11/23/04ISR Number: 4508739-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0516103A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1	MON	Bruxism		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Constipation					

Date:11/23/04ISR Number: 4508740-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0516133A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
5	MON	Menstruation Irregular		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508741-2Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #2004-02794

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG			Blood Pressure Increased	Bupropion Sr	PS	Glaxosmithkline	ORAL
Variable dose	25	DAY	Constipation				
			Feeling Cold	Nicorette	C	Glaxosmithkline	ORAL
			Heart Rate Increased	Mvi	C		
			Insomnia	Calcium	C		
			Panic Attack	Magnesium	C		
			Pharmaceutical Product Complaint	Aspirin	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508742-4Report Type:Periodic  
Age:15 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516223A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Mania	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/23/04ISR Number: 4508743-6Report Type:Periodic  
Age:22 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516226A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Rash	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day				Propecia	C		

Date:11/23/04ISR Number: 4508744-8Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516253A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	15	MON					
		Loss Of Libido		Fosamax	C		
				Effexor	C		

Date:11/23/04ISR Number: 4508745-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0516437A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	2	WK					
				No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508746-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0516626A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Twice		Pharmaceutical Product					
per day	4	YR					
		Complaint		Xanax	C		
				Synthroid	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508747-3Report Type:Periodic  
Age:12 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0516906A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	32 DAY						

Date:11/23/04ISR Number: 4508748-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517078A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK						
				Prednisone	C		

Date:11/23/04ISR Number: 4508749-7Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517267A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	5 MON						
		Dry Mouth		Niacin	C		
				Lipitor	C		

Date:11/23/04ISR Number: 4508750-3Report Type:Periodic  
Age:41 YR Gender:Female I/FU:F

Company Report #2004-02794

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Bupropion Hcl	PS	Glaxosmithkline	ORAL
150MG See							
dosage text		Cold Sweat					
		Constipation					
		Dizziness					

Heart Rate Increased  
 Hyperhidrosis  
 Insomnia  
 Micturition Disorder  
 Panic Attack  
 Peripheral Coldness  
 Pharmaceutical Product  
 Complaint  
 Sensation Of Heaviness

Date:11/23/04ISR Number: 4508751-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0517687A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Anorexia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	6 MON	Weight Decreased		Oral Contraceptives	C		

Date:11/23/04ISR Number: 4508752-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0517771A  
 Age: Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508753-9Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0517798A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	1 YR	Adverse Drug Reaction		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Sexual Dysfunction		Viagra	C		

Date:11/23/04ISR Number: 4508754-0Report Type:Periodic  
Age:48 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0518016A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Three times per day		Muscle Spasms		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Pain In Extremity		Levoxyl	C	Glaxosmithkline	ORAL
500MG Twice per day				Metformin	C		ORAL
				Multivitamin	C		
				Calcium	C		
				Glucophage	C		ORAL
500MG Twice per day				Glucosamine + Chondroitin	C		ORAL

Date:11/23/04ISR Number: 4508755-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0518186A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Weight Increased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508756-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0518230A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	6	WK					
20MG per day				Celexa	C		

Date:11/23/04ISR Number: 4508757-6Report Type:Periodic Company Report #2004-03080  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State		Bupropion Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day		Disturbance In Attention					
UNKNOWN		Drug Ineffective		Benadryl	C	Glaxosmithkline	
		Dysarthria					
		Euphoric Mood					
		Nervousness					

Date:11/23/04ISR Number: 4508758-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0518513A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3	WK					
		Nervousness					
				No Concurrent Medication	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508759-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518545A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	2 DAY			Fluoxetine	C		

Date:11/23/04ISR Number: 4508760-6Report Type:Periodic  
 Age:56 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518551A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				No Concurrent Medication	C		
per day							

Date:11/23/04ISR Number: 4508761-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518555A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Constipation		Effexor	C		
per day	2 YR						

Date:11/23/04ISR Number: 4508762-XReport Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0518616A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
2 YR				Lipitor	C		
				Doxepin	C		

Date:11/23/04ISR Number: 4508763-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518793A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Per day	4	WK	Drug Ineffective	Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508764-3Report Type:Periodic  
Age:29 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0518820A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Twice			Abdominal Pain Upper	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day			Flatulence				
			Pharmaceutical Product	Flexeril	C		
			Complaint	Endocet	C		
			Skin Odour Abnormal	Nasacort Aq Nasal			
				Spray	C		NASAL
				Protonix	C		
				Allegra	C		
				Zoloft	C		
				Ambien	C		
				Topamax	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508765-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518822A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
VAGINAL		Urinary Tract Infection		Wellbutrin Sr	PS	Glaxosmithkline	
		Vaginal Burning Sensation Vaginal Mycosis		Trazodone	C		

Date:11/23/04ISR Number: 4508766-7Report Type:Periodic  
 Age:42 YR Gender:Male I/FU:I

Company Report #2004S1001835

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown		Sexual Dysfunction		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
10MG Per day				Lotrel	SS		ORAL
10MG Unknown				Lexapro	SS		ORAL
.2MG Twice per day				Clonidine Hydrochloride	C		ORAL
50MG Per day				Hydrochlorothiazide + Amiloride	C		ORAL
10MG Unknown				Percocet	C		ORAL
UNKNOWN	150MG Unknown			Trazodone	C		
UNKNOWN	40MG Per day			Nexium	C		
UNKNOWN	20MG Unknown			Bextra	C		
UNKNOWN	350MG Unknown			Carisoprodol	C		

Date:11/23/04ISR Number: 4508767-9Report Type:Periodic  
 Age:16 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519050A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Liver Function Test		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Abnormal					
per day	3 YR			Accutane	C		

Date:11/23/04ISR Number: 4508768-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0519185A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hot Flush		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Bupropion Hydrochloride	SS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508769-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0519217A  
 Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema		Wellbutrin	PS	Glaxosmithkline	ORAL
7 DAY		Arthralgia		No Concurrent Medication	C		
		Paraesthesia					
		Rash					
		Serum Sickness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508770-9Report Type:Periodic  
Age:72 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519230A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diplopia		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice		Tremor					
per day	3 YR			Zoloft	C		
				Plavix	C		
				Nitroglycerin	C	Glaxosmithkline	
				Terazosin	C		
				Xalatan	C		

Date:11/23/04ISR Number: 4508771-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519246A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Deafness		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
1 YR							

Date:11/23/04ISR Number: 4508772-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519332A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Effect Decreased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Wellbutrin Xl	SS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508773-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0519354A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Metrorrhagia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508774-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519367A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
8 YR		Drug Ineffective		Zoloft	C		

Date:11/23/04ISR Number: 4508775-8Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519528A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day		Pruritus		Soy	C		ORAL

Date:11/23/04ISR Number: 4508776-XReport Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519708A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Tinnitus					
per day	3 YR			Wellbutrin Xl No Concurrent Medication	SS C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508777-1Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519716A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	30 WK	Agitation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Irritability		No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508778-3Report Type:Periodic  
Age:28 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0519729A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Twice		Nystagmus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	6 MON						

Date:11/23/04ISR Number: 4508779-5Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519919A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	8 MON	Dry Skin		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Rash		Lexapro	C		

Date:11/23/04ISR Number: 4508781-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520512A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Pruritus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day		Rash					
		Urticaria		Plavix	C		

Date:11/23/04ISR Number: 4508782-5Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520729A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
450MG Per day	8 YR			K-Dur	C	Glaxosmithkline	
				Lasix	C	Glaxosmithkline	
				Ambien	C		
				Ativan	C		

Date:11/23/04ISR Number: 4508783-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520768A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
1TAB Twice							
per day	1 WK						

Date:11/23/04ISR Number: 4508784-9Report Type:Periodic  
Age:20 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0520921A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

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Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508785-0Report Type:Periodic  
Age:60 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521295A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Migraine		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Per day							

Date:11/23/04ISR Number: 4508786-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521433A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sensory Disturbance		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508787-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521535A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Alcohol	C		
150MG Per day							

Date:11/23/04ISR Number: 4508788-6Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521539A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Per day	6 MON	Stomach Discomfort		No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508789-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521546A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

100MG Twice Hypoaesthesia Wellbutrin Sr PS Glaxosmithkline ORAL  
per day 2 YR Skin Discolouration

Date:11/23/04ISR Number: 4508790-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0521556A  
Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Headache					
per day	10	MON		Ephedra	C		
				Allegra	C		

Date:11/23/04ISR Number: 4508791-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0521582A  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
Other		Heart Rate Increased					
		Tremor					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508792-8Report Type:Periodic  
Age:17 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521634A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	4 YR			No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508793-XReport Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521637A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice				Coffee	C		
per day	1 WK						

Date:11/23/04ISR Number: 4508794-1Report Type:Periodic  
Age:67 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521664A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Per day	8 MON	Depression		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
150MG Per day	1 MON	Dyspepsia		Prempro	C		
				Xanax	C		
				Robaxin	C		

Date:11/23/04ISR Number: 4508795-3Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521816A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							

Rash  
per day 6 DAY

Birth Control Pills C

Date:11/23/04ISR Number: 4508796-5Report Type:Periodic  
Age:78 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521840A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 MON	Blood Pressure Increased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Xanax	C		
				Zestril	C		

Date:11/23/04ISR Number: 4508797-7Report Type:Periodic  
Age:27 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0521977A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Arthritis		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day							
300MG Per day				Wellbutrin Xl	SS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508798-9Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521997A

Outcome	PT
	Anxiety
	Depression

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Dose	Duration	Drug Ineffective Fatigue	Report Source	Product	Role	Manufacturer	Route
150MG As required	3 MON			Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Klonopin	C		
				Imitrex	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508799-0Report Type:Periodic Company Report #S04-USA-04583-01  
Age:58 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT Drug Screen Positive	Report Source	Product	Role	Manufacturer	Route
300MG Per day				Wellbutrin Sr	PS	Glaxosmithkline	ORAL
10MG Per day				Lexapro	SS		ORAL
				Hydrochlorothiazide	C		

Date:11/23/04ISR Number: 4508801-6Report Type:Periodic Company Report #2004-03577  
Age:75 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT Abdominal Pain Constipation Cough Diarrhoea Nausea Weight Decreased	Report Source	Product	Role	Manufacturer	Route
150MG Per day	39 DAY			Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
UNKNOWN	99 DAY			Mobec	SS		
UNKNOWN				Topamax	SS		
				Asa	C	Glaxosmithkline	
				Calcium	C		
				Vitamin C	C	Glaxosmithkline	
				Vitamin E	C		
				Centrum Silver	C		
				Osteo Biflex	C		

Date:11/23/04ISR Number: 4508802-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0522134A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr No Concurrent Medication	PS C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508803-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0522140A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Sr	PS	Glaxosmithkline	ORAL
300MG	Unknown						

Date:11/23/04ISR Number: 4508804-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0522257A  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
150MG	Per day 2 YR			Lipitor	C		
				Lasix	C	Glaxosmithkline	
				Atenolol	C		
				Zestril	C		
				Zoloft	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochlorothiazide C  
 Prilosec C Glaxosmithkline  
 Loratadine C

Date:11/23/04ISR Number: 4508805-3Report Type:Periodic  
 Age:41 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522438A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
300MG Twice							
per day	9	MON					

Date:11/23/04ISR Number: 4508806-5Report Type:Periodic  
 Age:47 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522459A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	3	YR					
		Insomnia		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
150MG Per day							
		Lethargy		Avalide	C		
				Multivitamin	C		

Date:11/23/04ISR Number: 4508807-7Report Type:Periodic  
 Age:25 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522480A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dental Caries		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	18	MON					
		Insomnia					
		Salivary Hypersecretion		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Per day	6	MON					
				Blood Thinner	C		
				Ultram	C		
				Celebrex	C		

Date:11/23/04ISR Number: 4508808-9Report Type:Periodic  
Age:71 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522567A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
300MG Per day				Wellbutrin Xl	SS	Glaxosmithkline	ORAL
				Paxil	C	Glaxosmithkline	ORAL
				Xanax	C		

Date:11/23/04ISR Number: 4508809-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522605A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Eyelid Pain					
per day		Nausea					

Date:11/23/04ISR Number: 4508810-7Report Type:Periodic  
Age:30 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522624A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Eye Pain					
per day		Nausea		No Concurrent			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Medication C

Date:11/23/04ISR Number: 4508811-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522802A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Condition Aggravated	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	2	MON	Drug Ineffective	Wellbutrin Xl	SS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508812-0Report Type:Periodic  
Age:31 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522820A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Erythema Multiforme	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice			Oedema Peripheral	Toprol Xl	C		
per day	3	WK	Urticaria	Levothyroxin	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508813-2Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522824A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
18	MON		Anxiety	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508814-4Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523053A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Twice			Abdominal Discomfort	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
			Anxiety				
per day	2	YR					
			Dry Eye	Wellbutrin Xl	SS	Glaxosmithkline	ORAL
300MG Per day	6	MON					
			Headache	Lorazepam	C		
			Tremor				

Date:11/23/04ISR Number: 4508815-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0523082A  
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Granuloma Annulare	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2	YR					
				No Concurrent Medication	C		

Date:11/23/04ISR Number: 4508816-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0523326A  
 Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Insomnia	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	4	MON					
			Toothache				
				Glucophage	C		
				Hctz	C		
				Calcium	C		
				Magnesium	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Zinc C  
 Unknown Medication C

Date:11/23/04ISR Number: 4508817-XReport Type:Periodic  
 Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523481A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Twice			Crying	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	2 WK		Drug Toxicity				
			Dysgeusia	Oral Contraceptives	C		
			Ill-Defined Disorder				
			Insomnia				
			Medication Error				
			Rash				

Date:11/23/04ISR Number: 4508818-1Report Type:Periodic  
 Age:76 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523630A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day			Aggression	Wellbutrin	PS	Glaxosmithkline	ORAL
			Amnesia	Vitamins	C		
			Blood Pressure Increased				

Date:11/23/04ISR Number: 4508819-3Report Type:Periodic  
 Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523641A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Per day			Insomnia	Wellbutrin	PS	Glaxosmithkline	ORAL
			Paraesthesia				

Date:11/23/04ISR Number: 4508820-XReport Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523767A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508821-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0523821A  
 Age:53 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice	1 YR	Nausea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day		Pharmaceutical Product					
		Complaint		Allegra	C		
		Pruritus		Stool Softener	C		
				Aciphex	C		
				Estraderm	C		

Date:11/23/04ISR Number: 4508822-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0524243A  
 Age:77 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice	9 WK	Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day				Tylenol #3	C		
				Prevacid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydralazine	C	
Lotensin	C	Glaxosmithkline
Neurontin	C	
Isosorbide	C	

Date:11/23/04ISR Number: 4508823-5Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524244A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Drug Exposure During	Wellbutrin Sr	PS	Glaxosmithkline	
100MG Twice							
			Pregnancy				
per day	9	MON					
			Lethargy	Prenatal	C		
				Anti-Nausea Drugs	C		
				Phenergan	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508824-7Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524535A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Drug Ineffective	Wellbutrin Sr	PS	Glaxosmithkline	

Date:11/23/04ISR Number: 4508825-9Report Type:Periodic  
 Age:31 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524745A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Headache	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	42	MON					
			Lethargy	Ambien	C		
				Calcium	C		
				Vitamin E	C		

Date:11/23/04ISR Number: 4508826-0Report Type:Periodic  
 Age:29 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0524746A

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
150MG Per day	MON	Genital Pruritus Female		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Vaginitis Bacterial		Spironolactone	C		

Date:11/23/04ISR Number: 4508827-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0524771A  
 Age: Gender:Male I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
150MG Twice		Blood Pressure Increased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	5 MON			Geodon	C		

Date:11/23/04ISR Number: 4508828-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525238A  
 Age:50 YR Gender:Male I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
100MG Twice		Tinnitus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	11 DAY			No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508829-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525425A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:11/23/04ISR Number: 4508830-2Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525452A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Smoker		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice		Stomach Discomfort					
per day	1 YR						

Date:11/23/04ISR Number: 4508831-4Report Type:Periodic  
Age:55 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0525729A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Papular		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
300MG Per day	5 DAY						

Date:11/23/04ISR Number: 4508832-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525801A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							
				Tramadol	C		
				Prevacid	C		
				Ziac	C		

Date:11/23/04ISR Number: 4508833-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525825A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Drug Ineffective	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day				Lisinopril	C		
				Aspirin	C	Glaxosmithkline	
				Zocor	C		
				Vitamin	C		
				Insulin	C		

Date:11/23/04ISR Number: 4508834-XReport Type:Periodic  
Age:17 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0525827A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Pruritus	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2	MON		Clonidine	SS		
UNKNOWN			3 DAY				

Date:11/23/04ISR Number: 4508835-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526003A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
900MG Per day			Accidental Overdose	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
			Tremor				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508836-3Report Type:Periodic  
Age:51 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0526197A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day				Lipitor	C		

Date:11/23/04ISR Number: 4508837-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526689A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
300MG Per day	6 YR	Somnolence		Celexa	C		

Date:11/23/04ISR Number: 4508838-7Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526709A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Twice				No Concurrent Medication	C		
per day	2 MON						

Date:11/23/04ISR Number: 4508839-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #2004/04109

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Bupropion Sr	PS	Glaxosmithkline	ORAL
150MG Single		Dizziness					
dose		Insomnia		Librium	C		
5MG As							

required

Toprol XL C

50MG Per day

Norvasc C

5MG Per day

Date:11/23/04ISR Number: 4508840-5Report Type:Periodic  
Age:47 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526866A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	5 YR			Depakote	C		

Date:11/23/04ISR Number: 4508841-7Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526879A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alanine Aminotransferase		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Increased		Klonopin	C		
400MG Per day	2 YR	Blood Alkaline Phosphatase Increased					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508842-9Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527096A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Acne		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Twice							
per day	3	MON					
				Cytomel	C	Glaxosmithkline	
				Levoxyl	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508843-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527116A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hot Flush		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Per day	1	WK					
		Insomnia					
		Menstruation Irregular					

Date:11/23/04ISR Number: 4508847-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527290A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Concerta	C		
200MG Per day				Diet Pill			
				Unspecified	C		

Date:11/23/04ISR Number: 4508848-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527298A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
200MG Twice							

per day 2 MON

No Concurrent Medication C

Date:11/23/04ISR Number: 4508849-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0527323A  
Age:56 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Anger		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	5	MON					
		Anxiety					
		Confusional State		Nicorette	C	Glaxosmithkline	
		Euphoric Mood					
		Irritability					
		Paranoia					

Date:11/23/04ISR Number: 4508852-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0527395A  
Age:38 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Pruritus		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
per day	17	DAY					
		Rash					
		Urticaria		Lexapro	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508853-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527397A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	56 WK			Paxil Cr	C	Glaxosmithkline	

Date:11/23/04ISR Number: 4508854-5Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527407A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Per day	2 DAY						
		Fatigue		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
150MG Per day	1 DAY						
		Palpitations		Singulair	C		
		Paraesthesia		Zoloft	C		
				Sam-E	C		

Date:11/23/04ISR Number: 4508855-7Report Type:Periodic  
 Age:20 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527423A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
100MG Twice							
per day	6 WK			Concerta	C		
				Medrol	C		
				Alcohol	C		

Date:11/23/04ISR Number: 4508856-9Report Type:Periodic  
 Age:28 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527583A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	100MG	Per day 3 YR	Drug Ineffective Mania		Wellbutrin	PS	Glaxosmithkline	ORAL
					Lamictal	C	Glaxosmithkline	
					Depakote	C		
					Provigil	C		
					Seroquel	C		

Date:11/23/04ISR Number: 4508858-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0527592A  
 Age:52 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG	Twice per day 2 YR	Blood Pressure Increased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508859-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0527625A  
 Age:57 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG	Per day 2 YR	Crying		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
			Depression		Synthroid	C	Glaxosmithkline	
			Emotional Disorder		Hormone Patch	C		
UNKNOWN			Tremor		Prevacid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508860-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527709A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alanine Aminotransferase Increased Aspartate Aminotransferase Increased		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508861-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527984A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200MG Twice per day	3 YR	Abdominal Pain Pain In Extremity		Bupropion Sr	PS	Glaxosmithkline	ORAL
				Wellbutrin Xl Adderall Estradiol Buspar Lexapro	SS C C C C	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508862-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528268A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL

Date:11/23/04ISR Number: 4508863-6Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0528543A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Agitation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Anxiety

per day

Unknown

C

Date:11/23/04ISR Number: 4508903-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0531408A

Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	4 DAY	Bite		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Convulsion Impaired Driving Ability Medication Error					

Date:11/23/04ISR Number: 4508904-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0532210A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Double Ureter Drug Exposure During Pregnancy Renal Disorder Vesicoureteric Reflux		Bupropion	PS	Glaxosmithkline	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04ISR Number: 4508914-9Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0534690A

Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Chest Discomfort		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	16 DAY	Muscle Spasms Tachycardia Yawning					

Date:11/23/04ISR Number: 4508918-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0534891A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation Anger Belligerence Delusional Disorder, Persecutory Type Emotional Disorder Psychotic Disorder		Wellbutrin	PS	Glaxosmithkline	

Date:11/23/04ISR Number: 4512050-5Report Type:Expedited (15-DaCompany Report #2004-122401-NL

Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 15 MG ONCE		Anxiety	Health	Remeron	PS		
Intervention to 225 MG DAILY		Benign Prostatic Hyperplasia	Professional	Wellbutrin	SS		ORAL
Prevent Permanent ORAL Impairment/Damage 300 MG DAILY		Major Depression Polyuria		Wellbutrin	SS		ORAL
ORAL 450 MG DAILY		Urinary Retention		Wellbutrin	SS		ORAL

ORAL

Wellbutrin

SS

ORAL

300 MG DAILY

ORAL

Sertraline

Hydrochloride

SS

50 MG DAILY

Clonazepam

SS

Date:11/24/04ISR Number: 4510234-3Report Type:Expedited (15-DaCompany Report #US-ROCHE-379171

Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anger		Pegasys	PS	Roche	
SUBCUTANEOUS							
		Back Pain		Copegus	SS	Roche	ORAL
		Blindness		Wellbutrin	SS		ORAL
FIRST							
THERAPY.		Depression					
		Dyspnoea		Wellbutrin	SS		ORAL
SECOND							
THERAPY.		Headache					
		Myalgia		Atenolol	C		ORAL
		Road Traffic Accident		Bextra	C		ORAL
		Tinnitus		Zanaflex	C		ORAL
		Visual Acuity Reduced					
		Weight Decreased					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/24/04ISR Number: 4510433-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0534478A  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
18 MON							
		Bursitis		Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day	6 MON						
		Headache		Prozac	C		
		Joint Effusion		Depakote	C		
		Uterine Haemorrhage					

Date:11/24/04ISR Number: 4510462-7Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0351173A  
Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hyperthyroidism	Health	Bupropion			
		Insomnia	Professional	Hydrochloride	PS	Glaxosmithkline	
UNKNOWN	150MG	See					
dosage text	39 DAY	Palpitations					

Date:11/24/04ISR Number: 4513176-2Report Type:Expedited (15-DaCompany Report #USA-2004-0013994  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Aortic Atherosclerosis	Consumer	Oxycontin Tablets			
Other		Brain Oedema	Health	(Oxycodone			
		Cardiac Arrest	Professional	Hydrochloride) Cr			
		Coma	Other	Tablet	PS		
		Coronary Artery		Oxyir (Oxycodone			
		Atherosclerosis		Hydfrochloride)	SS		
MG							
		Cough		Propoxyphene			
		Drug Dependence		(Dextropropoxyphene)	SS		
		Dyspnoea		Bupropion			
		Emphysema		(Amfebutamone)	SS		
		Feeling Abnormal		Gabapentin			
		Goitre		(Gabapentin)	SS		
		Multiple Drug Overdose		Acetaminophen			

Accidental  
Pneumonia  
Pneumonia Aspiration  
Polysubstance Abuse  
Pulmonary Congestion  
Pulmonary Oedema  
Respiratory Depression  
Resuscitation

(Paracetamol)

SS

Date:11/24/04ISR Number: 4513745-XReport Type:Expedited (15-DaCompany Report #HQWYE557919NOV04  
Age:62 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Agitation
Initial or Prolonged	Cerebral Atrophy
	Clumsiness
	Confusional State
	Depressed Level Of
	Consciousness
	Diarrhoea
	Drug Interaction
	Electrocardiogram Qt
	Prolonged
	Faecal Incontinence

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Gait Disturbance Hallucinations, Mixed Hyperhidrosis	Literature	Effexor (Venlafaxine Hydrochloride, Unspec)	PS		
75 MG 1X PER	1 DAY	Insomnia Lethargy Myoclonus  Serotonin Syndrome					
300 MG, 1X	PER 1 DAY	Transaminases Increased Urinary Incontinence		Bupropion (Amfebutamone, )	SS		
50 MG 1X PER	1 DAY			Sertraline (Sertraline, )	SS		
				Piracetam (Piracetam)	C		

Date:11/24/04ISR Number: 4514772-9Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 233006

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG BID		Hospitalization - Initial or Prolonged Disability ORAL		Effexor Xr 75 Mg Wyeth	PS	Wyeth	ORAL
100 MG BID				Wellbutrin Sr 100 Mg	SS		ORAL

Date:11/26/04ISR Number: 4527147-3Report Type:Periodic  
Age:18 YR Gender:Female I/FU:I

Company Report #04P-163-0273168-00

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion Depression Eye Movement Disorder Fall	Consumer Health Professional	Depakote Er 500mg (Depakote Er) (Divalproex Sodium) (Divalproex Sodium)			
500 MG, 2 IN					PS		ORAL
1 D, ORAL		Intentional Misuse					
2 MG, 1 IN 1		Weight Increased		Risperidone	SS		ORAL
D, ORAL							
ORAL				Venlafaxine Hydrochloride	SS		ORAL
ORAL				Bupropion Hydrochloride	SS		ORAL
				Strattera	C		
				Levothyroxine Sodium	C		
				Quetiapine	C		

Date:11/29/04ISR Number: 4512271-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0526859A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL
				Wellbutrin Sr	SS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/29/04ISR Number: 4512276-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0535150A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Agitation Anxiety Drug Withdrawal Syndrome Restlessness Sleep Disorder Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/29/04ISR Number: 4512287-5Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0347425A

Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG		Retinopathy Hypertensive Tinnitus		Zyban	PS	Glaxosmithkline	ORAL
Variable dose	12 DAY	Visual Acuity Reduced		No Concurrent Medication	C		

Date:11/29/04ISR Number: 4512303-0Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0350118A

Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day	10 DAY	Contusion Vasculitis		Zyban	PS	Glaxosmithkline	ORAL
1TAB Per day				Cartia	C	Glaxosmithkline	ORAL
RESPIRATORY (INHALATION)				Actonel Bricanyl	C C		ORAL
1G per day				Unknown Drug Naproxin	C C		

Panamax	C	Glaxosmithkline
Sinequan	C	
Spiriva	C	
Symbicort	C	
Unknown Drug	C	
Unknown Drug	C	

Date:11/29/04ISR Number: 4512310-8Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0356698A  
 Age:58 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Grand Mal Convulsion					
per day	16	DAY					
		Tongue Biting		Amitriptyline	C		
UNKNOWN	2MG	Per day					
		Tonic Clonic Movements		Allopurinol	C	Glaxosmithkline	
UNKNOWN	100MG	Per day					
				Hrt	C		
UNKNOWN				Simvastatin	C		
UNKNOWN	10MG	Per day					
UNKNOWN	10MG	Per day		Enalapril	C		

Date:11/29/04ISR Number: 4512314-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0357170A  
 Age:69 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Glomerulonephritis		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	9	WK					
		Hypertension		Bendrofluazide	C	Glaxosmithkline	ORAL
2.5MG per day		Nephrotic Syndrome		Frusemide	C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Doxazosin	C	ORAL
Perindopril	C	ORAL

Date:11/29/04ISR Number: 4512317-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0357185A  
 Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	8 DAY	Asthenia		Zyban	PS	Glaxosmithkline	ORAL
		Chest Discomfort		Ipratropium Bromide	C		
	40MCG Three times per day	Dry Mouth					
	250MG Three times per day	Feeling Abnormal		Cefaclor	C		ORAL
	250MG Three times per day 20 DAY	Nausea					
	250MCG As required	Tremor		Ventolin	C	Glaxosmithkline	
		Vision Blurred					

Date:11/30/04ISR Number: 4513534-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498016A  
 Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	36 DAY	Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Gastric Perforation		Celexa	C		
	106 DAY	Palpitations		Wellbutrin Sr	C	Glaxosmithkline	ORAL
				Ativan	C		

Date:11/30/04ISR Number: 4513535-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0502854A  
 Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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300MG Per day 4 DAY	Rash Generalised	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Guaifenesin	C		
		Albuterol	C	Glaxosmithkline	
		Advil	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513536-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0502870A  
 Age:22 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Vision Blurred		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 109 DAY			Bactrim Ds	C	Glaxosmithkline	ORAL
1U Per day						

Date:11/30/04ISR Number: 4513538-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505370A  
 Age: Gender: I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Rash		Wellbutrin Xl	PS	Glaxosmithkline	

Date:11/30/04ISR Number: 4513539-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505372A  
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 1 WK			Amerge	C	Glaxosmithkline	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513540-1Report Type:Periodic  
Age:53 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0505385A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	52 DAY			Covera	C		
				Lipitor	C		
				Diovan Hct	C		
				Glucophage	C		
				Vicodin Es	C		
				Mobic	C		

Date:11/30/04ISR Number: 4513541-3Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0505547A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharmaceutical Product		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Complaint					

Date:11/30/04ISR Number: 4513542-5Report Type:Periodic  
Age:80 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0507160A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysarthria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK	Tremor		Synthroid	C	Glaxosmithkline	
				Zestril	C		

Date:11/30/04ISR Number: 4513543-7Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0508054A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Lexapro	C		

Date:11/30/04ISR Number: 4513544-9Report Type:Periodic  
Age:42 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0508653A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	0 DAY	Dizziness		Hctz	C		
		Tinnitus		Atenolol	C		

Date:11/30/04ISR Number: 4513545-0Report Type:Periodic  
Age:30 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0515883A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Erythema Multiforme		Wellbutrin	PS	Glaxosmithkline	ORAL
3 WK							
Initial or Prolonged		Rash		Birth Control Pills	C		
Other							

Date:11/30/04ISR Number: 4513546-2Report Type:Periodic  
Age:45 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0516258A

Outcome  
Hospitalization -  
Initial or Prolonged

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG In the morning		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Three times per day				Dilantin	C		
200MG At night				Trazodone	C		
15MG At night				Zyprexa	C		
1TAB Per day				Aspirin	C	Glaxosmithkline	
75MG Per day				Plavix	C		

Date:11/30/04ISR Number: 4513547-4Report Type:Periodic  
Age:15 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0517857A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
6 MON		Leukopenia					

Date:11/30/04ISR Number: 4513548-6Report Type:Periodic  
Age:58 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0518073A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3 WK	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Anger		Humalog	C		
		Anorexia		Evista	C		
		Anxiety		Vitamins	C		
		Crying					

Dizziness  
 Dysgeusia  
 Emotional Disorder  
 Fungal Infection  
 Hyperhidrosis  
 Insomnia  
 Myalgia  
 Nausea  
 Tremor  
 Weight Decreased

Date:11/30/04ISR Number: 4513549-8Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0518097A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
Disability		Coordination Abnormal		Cardura	C		ORAL
Other				Thalidomide	C		
2MG per day							

Date:11/30/04ISR Number: 4513550-4Report Type:Periodic  
 Age:29 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0518423A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Drug Ineffective		Oral Contraceptive	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513551-6Report Type:Periodic  
Age:31 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0518798A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 MON			Adderall	C		

Date:11/30/04ISR Number: 4513552-8Report Type:Periodic  
Age:34 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0518819A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Asthenia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Drug Ineffective		Unknown Medication	C		
		Headache		Effexor	C		
		Nausea		Vistaril	C		
		Sensation Of Heaviness		Vitamins	C		

Date:11/30/04ISR Number: 4513554-1Report Type:Periodic  
Age:28 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0519027A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Tachycardia					

Date:11/30/04ISR Number: 4513555-3Report Type:Periodic  
Age:51 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0519365A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	MON						
Disability		Pharmaceutical Product		No Concurrent			
Other		Complaint		Medication	C		
		Suicidal Ideation					

Date:11/30/04ISR Number: 4513556-5Report Type:Periodic  
Age:13 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0520321A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Abnormal Behaviour		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513557-7Report Type:Periodic  
Age:37 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0520375A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day	3 MON	Irritability		Unknown Medication	C		
		Pharmaceutical Product Complaint					
		Stool Analysis Abnormal					

Date:11/30/04ISR Number: 4513558-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520513A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK	Dizziness		Xanax	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Fiorinal 3 C

Date:11/30/04ISR Number: 4513559-0Report Type:Periodic  
Age:50 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520520A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypoventilation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK	Insomnia		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513560-7Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520526A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6 MON	Depression		Omega 3	C	Glaxosmithkline	
		Insomnia		Selenium	C		
				Niacin	C		
				Vitamin B6	C		
				Vitamin B Complex	C		

Date:11/30/04ISR Number: 4513561-9Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520527A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	7 MON			Buspar	C		
				Zyprexa	C		

Date:11/30/04ISR Number: 4513562-0Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520537A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 DAY	Urticaria		Claritin	C		

Date:11/30/04ISR Number: 4513563-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0520549A  
 Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Mouth Ulceration		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 MON	Urticaria		Lexapro	C		ORAL
30MG Per day				Xanax	C		ORAL
.15MG Per day				Provigil	C		ORAL
100MG In the morning				Allegra	C		
				Levoxyl	C	Glaxosmithkline	
				Celebrex	C		
				Cytomel	C	Glaxosmithkline	
				Zocor	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513564-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520552A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513565-6Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520553A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	13 DAY	Dry Mouth		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Nausea		Synthroid	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513566-8Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520558A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	1 YR	Abdominal Discomfort		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dysphonia		Nexium	C		
		Fatigue		Prednisolone	C	Glaxosmithkline	
		Muscle Tightness					
		Musculoskeletal Stiffness					
		Tremor					

Date:11/30/04ISR Number: 4513567-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520714A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Unknown		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513568-1Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0520734A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vomiting		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG	Unknown						

Date:11/30/04ISR Number: 4513569-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520744A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	

Date:11/30/04ISR Number: 4513570-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520747A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG	Per day 8 WK	Visual Disturbance		Lorazepam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513571-1Report Type:Periodic  
Age:61 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520758A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Per day	4 MON	Sleep Apnoea Syndrome	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Ativan	C		
				Flomax	C		
				Advair	C	Glaxosmithkline	
				Claritin	C		
				Flonase	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513572-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520762A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Blood Pressure Increased	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513573-5Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520774A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Per day	18 MON	Agitation	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Anxiety	Combivent	C		
			Chest Pain	Norvasc	C		
			Chills	K-Dur	C	Glaxosmithkline	
			Hyperhidrosis	Glucotrol Xl	C		
				Glucophage	C		
				Demadex	C		
				Estratest Hs	C		
				Aygestin	C		
				Methadone	C	Glaxosmithkline	
				Ativan	C		
				Ambien	C		

Date:11/30/04ISR Number: 4513574-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520897A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Sexual Dysfunction		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513575-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0520945A  
 Age:44 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3 WK	Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513576-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0520949A  
 Age:49 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1 WK	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Anxiety		Xanax	C		
		Hot Flush					
		Nervousness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513577-2Report Type:Periodic  
Age:49 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0520966A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Xanax	C		
				Zofran	C	Glaxosmithkline	
				Unknown Medication	C		
UNKNOWN							

Date:11/30/04ISR Number: 4513578-4Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520980A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Breast Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	17 DAY			Wellbutrin Sr	SS	Glaxosmithkline	ORAL
150MG Twice		Nervousness					
per day	1 YR			Advil	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513579-6Report Type:Periodic  
Age:33 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0521096A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Menstruation Irregular		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	11 DAY			Crestor	C		

Date:11/30/04ISR Number: 4513580-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521105A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hyperhidrosis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513581-4Report Type:Periodic  
Age:55 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0521112A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	7 DAY		Oedema Genital	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Vulval Oedema	No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513582-6Report Type:Periodic  
Age:18 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0521116A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -			Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							
Initial or Prolonged				Lexapro	C		ORAL
10MG Per day	238 DAY						
Other							

Date:11/30/04ISR Number: 4513583-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521222A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Ageusia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Claritin -D	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513584-XReport Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0521280A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513585-1Report Type:Periodic  
Age:69 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521284A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	4 WK	Cold Sweat		Prozac	C		
		Nausea		Cardizem	C	Glaxosmithkline	
		Nervousness		Prilosec	C	Glaxosmithkline	
				Bextra	C		
				Tricor	C		
				Premarin	C		
				Zelnorm	C		

Date:11/30/04ISR Number: 4513586-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521298A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Muscle Relaxant	C		

Date:11/30/04ISR Number: 4513587-5Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521301A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	6 DAY			Zyprexa	C		

Date:11/30/04ISR Number: 4513588-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521317A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin Xl No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513589-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521408A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Therapeutic Response		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Unknown		Decreased					

Date:11/30/04ISR Number: 4513590-5Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0521430A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sexual Dysfunction Somnolence		Wellbutrin Xl Venlafaxine	PS SS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513591-7Report Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521521A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	6 DAY	Epistaxis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513592-9Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521532A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	2 MON	Cystitis Interstitial		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Lexapro	C		
				Imitrex	C	Glaxosmithkline	
				Topomax	C		
				Flexeril	C		

Date:11/30/04ISR Number: 4513593-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521562A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	9 MON	Chest Discomfort		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Throat Tightness		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513594-2Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521573A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
UNKNOWN				Levoxyl	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513595-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521635A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3	DAY	Neck Pain	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513596-6Report Type:Periodic  
Age:68 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521639A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2	MON	Anxiety	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Blood Pressure Increased	Paxil	C	Glaxosmithkline	
			Dry Mouth	Glucosamine	C		
			Dyspnoea	Tylenol	C	Glaxosmithkline	
			Myalgia	Multivitamin	C		
			Oral Pain	Maalox	C		
			Swollen Tongue	Gaviscon	C	Glaxosmithkline	
			Vision Blurred	Folic Acid	C		
				Visine Tears	C		
				Ranitidine	C	Glaxosmithkline	
				Ditropan	C		
				Lisinopril	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Atenolol C

Date:11/30/04ISR Number: 4513597-8Report Type:Periodic  
Age:19 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521646A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1 DAY	Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Neurontin	C		

Date:11/30/04ISR Number: 4513598-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521649A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 WK	Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513599-1Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521655A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 WK	Rash Generalised		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Prozac	C		
				Tagamet	C	Glaxosmithkline	
				Multivitamin	C		

Date:11/30/04ISR Number: 4513600-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521657A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 MON	Cough		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Drug Interaction		Robitussin Dm	SS		
				No Concurrent			

Date:11/30/04ISR Number: 4513601-7Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521659A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day							
		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Amnesia		Triamterene	C	Glaxosmithkline	
		Blood Pressure Increased		Advil Cold + Sinus	C		
		Fatigue					
		Headache					
		Muscular Weakness					

Date:11/30/04ISR Number: 4513602-9Report Type:Periodic  
Age:61 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521665A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	7 WK						
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Colchicine	C		
				Aspirin	C	Glaxosmithkline	
				Flomax	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513603-0Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521820A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	1	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513604-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521838A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513605-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521964A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
				Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513606-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521966A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
5 WK				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513607-8Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521967A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3 MON	Drug Ineffective Paranoia		Wellbutrin Xl No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513608-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0521969A  
 Age:47 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Local Swelling		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
UNKNOWN	10MG Unknown	Paraesthesia Paraesthesia Oral		Lexapro	C		

Date:11/30/04ISR Number: 4513609-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0521971A  
 Age:32 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	4 DAY	Headache		Wellbutrin Xl No Concurrent Medication	PS C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513610-8Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521975A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	4 DAY			Flonase	C	Glaxosmithkline	
				Allegra D	C		

Date:11/30/04ISR Number: 4513611-XReport Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521978A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Balance Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	23 WK	Tremor		Klonopin	C		
				Zoloft	C		
				Verapamil	C		
				Topamax	C		
				Resperidol	C		

Date:11/30/04ISR Number: 4513612-1Report Type:Periodic  
Age:21 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521983A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphagia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK	Oedema Peripheral					
		Urticaria					

Date:11/30/04ISR Number: 4513613-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522095A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hyperhidrosis					
		Nausea					

Date:11/30/04ISR Number: 4513614-5Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522108A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	5 DAY	Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Headache		Hyzaar	C		
				Zoloft	C		
				Yasmin	C		

Date:11/30/04ISR Number: 4513615-7Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522119A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
per day	1 WK	Insomnia					
		Nausea		Climara	C		
		Vertigo					
		Vestibular Neuronitis					
		Vomiting					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513616-9Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522132A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	6 MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Trazodone	C		

Date:11/30/04ISR Number: 4513617-0Report Type:Periodic  
Age:56 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522143A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	8 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
Other	150MG Per day			Effexor Xr	C		ORAL

Date:11/30/04ISR Number: 4513619-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522220A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	2 WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Synthroid	C	Glaxosmithkline	
				Vitamins	C		
				Atenolol	C		

Date:11/30/04ISR Number: 4513620-0Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522221A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513621-2Report Type:Periodic  
Age:52 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522225A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day	WK	Abdominal Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged 20MG Per day		Pancreatitis		Lexapro	C		
				Unspecified Medications	C		

Date:11/30/04ISR Number: 4513622-4Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522228A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3 WK	Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513623-6Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522238A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 MON	Dyspepsia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Fatigue		Klor Con	C	Glaxosmithkline	
		Headache		Atenolol	C		
				Hctz	C		
				Estratest Hs	C		
				Effexor Xr	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

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Date:11/30/04ISR Number: 4513624-8Report Type:Periodic  
Age:26 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522240A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2	WK		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513625-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522247A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	7	WK		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513626-1Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522248A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:11/30/04ISR Number: 4513627-3Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522254A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3	DAY		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513628-5Report Type:Periodic  
Age:55 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522262A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3	DAY	Dry Mouth	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Lipitor	C		
				Atenolol	C		
				Zetia	C		
				Plavix	C		

Date:11/30/04ISR Number: 4513629-7Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522265A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	2	MON	Lymphadenopathy	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Pruritus	Zoloft	C		
			Rash	Alprazolam	C		
				Bextra	C		
				Albuterol	C	Glaxosmithkline	
				Zyrtec	C	Glaxosmithkline	
				Astelin	C		
				Flonase	C	Glaxosmithkline	
				Temazepam	C		

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Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513630-3Report Type:Periodic  
Age:43 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522395A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	450MG Per day	57 DAY	Convulsion	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513631-5Report Type:Periodic  
Age:24 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522439A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day	7 DAY	Agitation	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Fatigue Headache Irritability	No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513632-7Report Type:Periodic  
Age:42 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522445A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	6 MON		Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513633-9Report Type:Periodic  
Age:57 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522448A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	300MG Per day	6 MON	Grand Mal Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged			Urinary Incontinence	Hctz	C		
			Vomiting	Glucophage	C		

Date:11/30/04ISR Number: 4513634-0Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522460A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Sexual Dysfunction		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513635-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0522464A  
Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Irritability		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513636-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0522472A  
Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2	WK						

Date:11/30/04ISR Number: 4513637-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0522478A  
Age:45 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG	Per day	Cough					

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Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513638-8Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522599A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG	Unknown	Dry Mouth Insomnia					

Date:11/30/04ISR Number: 4513639-XReport Type:Periodic  
Age:50 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522629A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Allergic		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG	Per day	Tinnitus		Naprosyn	SS		
UNKNOWN				Adderall	C		

Date:11/30/04ISR Number: 4513640-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522634A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG	Per day 1 WK			No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513641-8Report Type:Periodic  
Age:51 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522635A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG	Per day 3 MON	Diarrhoea Nausea		Unspecified Medication	C		ORAL
60MG	Per day 35 DAY						

Lexapro	C
Ativan	C
Restoril	C
Soma	C
Codeine	C
Premarin	C
Fosamax	C
Plaquenil	C
Prednisone	C
Humira	C
Norco	C
Actonel	C
Mvi	C
Vitamin D	C
Vitamin E	C

Date:11/30/04ISR Number: 4513643-1Report Type:Periodic  
 Age:49 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522647A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY			Neurontin	C		
				Synthroid	C	Glaxosmithkline	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513644-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522655A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513645-5Report Type:Periodic  
Age:42 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522659A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Life-Threatening		Wellbutrin	PS	Glaxosmithkline	ORAL
MON		Grand Mal Convulsion		Alcohol	SS		
Other				Adderall	SS		
				Celexa	C		
60MG Per day							

Date:11/30/04ISR Number: 4513646-7Report Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522661A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	8 DAY	Feeling Jittery		Klonopin	C		
		Insomnia					

Date:11/30/04ISR Number: 4513647-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522663A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6 WK	Abdominal Pain Upper		Birth Control Pills	C		
		Nausea					

Date:11/30/04ISR Number: 4513648-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0522664A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
6 WK				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513649-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0522665A  
Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anger		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 21 DAY		Arthralgia		No Concurrent Medication	C		
		Asthenia					
		Dry Mouth					
		Insomnia					
		Pollakiuria					

Date:11/30/04ISR Number: 4513650-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0522769A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Acne		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

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Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513651-0Report Type:Periodic  
Age:58 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522793A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
		Pharmaceutical Product		Metformin	C		
		Complaint		Actos	C		
		Stool Analysis Abnormal		Zyrtec	C	Glaxosmithkline	
				Lipitor	C		
				L-Thyroxine	C	Glaxosmithkline	
				Estradiol	C		
				Singulair	C		
				Cortisone	C		

Date:11/30/04ISR Number: 4513652-2Report Type:Periodic  
Age:19 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522794A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	6 WK	Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Irritability					

Date:11/30/04ISR Number: 4513653-4Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522797A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
per day	3 WK	Increased Appetite					
		Pruritus		Synthroid	C	Glaxosmithkline	
				Maxzide	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513654-6Report Type:Periodic  
Age:27 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522801A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day	MON	Eye Movement Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Mastication Disorder		Lamictal	C	Glaxosmithkline	ORAL
			Tardive Dyskinesia		Lexapro	C		ORAL
					Depakote Er	C		ORAL
					Geodon	C		ORAL
215 DAY								
.5U As					Lorazepam	C		ORAL
required								

Date:11/30/04ISR Number: 4513655-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0522808A  
Age: Gender: I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513656-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0522809A  
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day	2 DAY	Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
					No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513657-1Report Type:Periodic  
Age:55 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522810A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
450MG Per day		Accidental Overdose		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
10MG Twice		Agitation		Adderall	C		ORAL
per day		Dizziness					
		Nausea					

Date:11/30/04ISR Number: 4513658-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522817A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
450MG Per day		Muscle Tightness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Adderall	C		

Date:11/30/04ISR Number: 4513659-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522826A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	4 DAY	Disorientation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Memory Impairment					
		Palpitations					
		Panic Reaction					

Date:11/30/04ISR Number: 4513660-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522895A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
4 DAY		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Swelling Face					

Urticaria

Date:11/30/04ISR Number: 4513661-3Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522903A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK	Sensation Of Blood Flow		Zantac	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513662-5Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522904A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:11/30/04ISR Number: 4513663-7Report Type:Periodic  
Age:58 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0522916A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	10 DAY	Nausea		Vioxx	C		
				Activella	C		
				Effexor	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513664-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522922A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513665-0Report Type:Periodic  
Age:58 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522929A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK			No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513666-2Report Type:Periodic  
Age:70 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0523038A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3 WK	Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Eyelid Oedema		Toprol Xl	C		
		Throat Irritation		Levothroid	C	Glaxosmithkline	
		Urticaria		Buspar	C		
				Hyzaar	C		
				Celebrex	C		
				Questran	C		

Date:11/30/04ISR Number: 4513667-4Report Type:Periodic  
Age:41 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0523041A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	26 DAY	Dyspnoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent			

Date:11/30/04ISR Number: 4513668-6Report Type:Periodic  
 Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523045A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Haemorrhage Subcutaneous	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2	MON		Gabitril	C		
				Trazodone	C		

Date:11/30/04ISR Number: 4513669-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523047A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Anorexia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day			Diarrhoea	Oral Contraceptive	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513670-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523057A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	7 WK			Wellbutrin	PS	Glaxosmithkline	ORAL
		Abnormal Dreams Acne Disturbance In Attention Headache Irritability Sleep Disorder					

Date:11/30/04ISR Number: 4513671-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523065A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 MON			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Rash		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513672-8Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0523084A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 WK			Wellbutrin	PS	Glaxosmithkline	ORAL
		Suicidal Ideation					

Date:11/30/04ISR Number: 4513673-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523165A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dysgeusia Hypersensitivity Oropharyngeal Swelling Swelling Face					

Date:11/30/04ISR Number: 4513674-1Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0523180A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513675-3Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0523205A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Wellbutrin Sr	C	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513676-5Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523320A

Outcome	PT
	Anxiety
	Mood Altered
	Nervousness
	Pharmaceutical Product
	Complaint

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tension

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	10 MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Premarin	C		
			Trazodone	C		

Date:11/30/04ISR Number: 4513677-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0523332A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthralgia Joint Swelling		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513678-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0523448A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	

Date:11/30/04ISR Number: 4513679-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0523466A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Myalgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513680-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0523473A  
 Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3 WK	Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Insomnia		Meprobamate	C		

Date:11/30/04ISR Number: 4513681-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523520A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Intolerance		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513682-0Report Type:Periodic  
Age:41 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0523523A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day							

Date:11/30/04ISR Number: 4513683-2Report Type:Periodic  
Age:58 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0523528A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hepatic Enzyme Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
Other 300MG Unknown							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513684-4Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523619A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3 WK			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dizziness					
		Hyperhidrosis		No Concurrent Medication	C		
		Tinnitus					

Date:11/30/04ISR Number: 4513685-6Report Type:Periodic  
Age:36 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0523629A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hyperhidrosis		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
1MG Per day				Hytrin	C		

Date:11/30/04ISR Number: 4513686-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523642A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4 WK						
		Convulsion		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513687-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523715A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Nervousness					

Date:11/30/04ISR Number: 4513688-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0523718A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513689-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0523721A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Libido Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513692-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0523759A  
Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 13 WK		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513693-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523763A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	9 DAY	Diplopia					
		Vision Blurred		Diazepam	C		

Date:11/30/04ISR Number: 4513694-7Report Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523779A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day	3 MON	Stool Analysis Abnormal					
				Provigil	C		
				Xanax	C		

Date:11/30/04ISR Number: 4513695-9Report Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523785A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice		Agitation					
per day	5 WK	Anxiety					
		Feeling Hot		Inderal	C		
		Heart Rate Increased		Zomig	C		
		Insomnia		Glucosamine			
		Thinking Abnormal		Chondroitin	C		

Date:11/30/04ISR Number: 4513696-0Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523787A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK	Bone Pain					
		Insomnia		Depakote	C		

Date:11/30/04ISR Number: 4513697-2Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523793A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dry Mouth		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK	Headache		Nexium	C		
		Insomnia					

Date:11/30/04ISR Number: 4513698-4Report Type:Periodic  
Age:52 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0523799A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	11 MON	Grand Mal Convulsion		Symbyax	SS		
		Tongue Biting		Fluoxetine	C		

Date:11/30/04ISR Number: 4513700-XReport Type:Periodic  
Age:18 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0523800A

Outcome	PT
Other	Dyspnoea
	Hypersensitivity
	Oropharyngeal Swelling

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Urticaria

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
5 YR			Lipitor	C		
			Ortho Tri Cyclen	C		ORAL
			Zyrtec	C	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513701-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0523801A  
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	1 MON	Irritability		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Nausea		Spiriva	C		
				Niaspan	C		
				Crestor	C		
				Mobic	C		
				Zyloprim	C	Glaxosmithkline	
				Lotensin	C	Glaxosmithkline	
				Trental	C		
				Tricor	C		

Date:11/30/04ISR Number: 4513702-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0523806A  
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	9 WK	Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
4 WK		Dehydration		Zoloft	SS		
		Diarrhoea		Klonopin	C		
		Drug Ineffective		Fioricet	C		
		Oral Pain					

Date:11/30/04ISR Number: 4513704-7Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523811A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Nausea		Detrol La	C		

Date:11/30/04ISR Number: 4513705-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523815A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oral Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Unknown							

Date:11/30/04ISR Number: 4513706-0Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523820A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Depression		Paxil Cr	C	Glaxosmithkline	
		Drug Ineffective		Metformin	C		
				Insulin 70/30	C		
				Monopril	C		
				Lopid	C		
				Lipitor	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dyazide	C	Glaxosmithkline
Singulair	C	
Albuterol	C	Glaxosmithkline
Aspirin	C	Glaxosmithkline

Date:11/30/04ISR Number: 4513707-2Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523831A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	25 DAY	Constipation		Motrin	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513708-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524051A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Contusion		Wellbutrin	PS	Glaxosmithkline	
Other		Convulsion					
		Local Swelling					

Date:11/30/04ISR Number: 4513709-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524076A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Psychomotor Hyperactivity		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:11/30/04ISR Number: 4513710-2Report Type:Periodic  
Age:70 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0524202A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Palpitations		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Ambien	C		

Vioxx C  
Zocor C

Date:11/30/04ISR Number: 4513711-4Report Type:Periodic  
Age:39 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0524215A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Acne		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2 DAY		Rash					

Date:11/30/04ISR Number: 4513712-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524224A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Irritability		Paxil	SS	Glaxosmithkline	ORAL
10MG Per day		Nervousness		Klonopin	C		
		Psychomotor Hyperactivity					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513713-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524228A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	3 WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Ranitidine	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513714-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524234A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Risperdal	C		

Date:11/30/04ISR Number: 4513715-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524379A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropion	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513716-3Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524488A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	3 WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Hormone Replacement	C		

Date:11/30/04ISR Number: 4513717-5Report Type:Periodic  
Age:14 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524496A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	10	MON		No Concurrent Medication	C		
		Hallucination					

Date:11/30/04ISR Number: 4513718-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0524500A  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2	MON		No Concurrent Medication	C		
		Irritable Bowel Syndrome					
		Pollakiuria					
		Urinary Tract Infection					

Date:11/30/04ISR Number: 4513719-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0524507A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
1	MON						
		Tinnitus					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513720-5Report Type:Periodic  
Age:35 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524509A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	4 MON			Lexapro	C		

Date:11/30/04ISR Number: 4513721-7Report Type:Periodic  
Age:54 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524521A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Initial Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK	Insomnia		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513722-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524557A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Fatigue					

Date:11/30/04ISR Number: 4513723-0Report Type:Periodic  
Age:34 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524560A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression Drug Ineffective		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
150MG Twice per day	10 DAY			Depakote	C		

Date:11/30/04ISR Number: 4513724-2Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0524561A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	10 DAY						

Date:11/30/04ISR Number: 4513725-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524563A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day							

Date:11/30/04ISR Number: 4513726-6Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524575A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Unknown				Tenormin	C		
				Calcium	C		
				Magnesium	C		
				Potassium	C		
				Ergocalciferol	C		
				Insulin	C		
				Unknown Medication	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513727-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524584A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513728-XReport Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0524587A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Insomnia		Lexapro Vitamins	C C		

Date:11/30/04ISR Number: 4513729-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524663A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fall		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Gait Disturbance		Paxil	C	Glaxosmithkline	ORAL
				Unspecified Medications	C		

UNKNOWN

Date:11/30/04ISR Number: 4513730-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524670A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513731-XReport Type:Periodic  
Age:53 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0524714A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route	
Life-Threatening	300MG Per day	1	MON	Suicidal Ideation	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513732-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0524728A  
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2	DAY	Anxiety	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Myalgia	Lithium	C	Glaxosmithkline	
			Panic Attack				
			Pyrexia				

Date:11/30/04ISR Number: 4513733-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0524732A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Urticaria	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513734-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524734A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	4 DAY			No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513735-7Report Type:Periodic  
Age:76 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524737A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY	Hyperhidrosis		Amoxil	SS	Glaxosmithkline	
				Toprol	C		
				Lipitor	C		
				Nexium	C		
				Plavix	C		

Date:11/30/04ISR Number: 4513736-9Report Type:Periodic  
Age:19 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524741A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Stomach Discomfort		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 MON	Vomiting		Unspecified Medication	C		

Date:11/30/04ISR Number: 4513737-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524743A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	4 MON			Ritalin	C		

Date:11/30/04ISR Number: 4513738-2Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524763A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice			Non-Cardiac Chest Pain	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
per day	6	WK					
				Oxycontin	C		
				Percocet	C		
				Valium	C		
				Albuterol Sulfate	C	Glaxosmithkline	
				Albuterol Inhaler	C	Glaxosmithkline	
				Lipitor	C		
				Benadryl	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513739-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524770A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Libido Decreased Medication Error	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513740-0Report Type:Periodic  
Age:45 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524874A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
1	YR						

Date:11/30/04ISR Number: 4513741-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524878A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin Xl No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513763-1Report Type:Periodic  
Age:44 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0511765A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 MON	Drug Ineffective		Decongestant	C		
		Irritability		Lexapro	C		
		Nervousness		Armour Thyroid	C		
		Tension					

Date:11/30/04ISR Number: 4513764-3Report Type:Periodic  
Age:41 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0513816A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Depakote	C		ORAL
1250MG per							
day							
.5MG Three				Klonopin	C		ORAL

times per day

Date:11/30/04ISR Number: 4513765-5Report Type:Periodic  
Age:33 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0530786A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
450MG Per day	7	MON	Drug Ineffective	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Stool Analysis Abnormal	Lamictal	C	Glaxosmithkline	ORAL
7	MON						

Date:11/30/04ISR Number: 4513766-7Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0530805A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Myalgia	Wellbutrin Xl	PS	Glaxosmithkline	

Date:11/30/04ISR Number: 4513767-9Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530831A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day			Flat Affect	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Mood Swings	Zelnorm	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513768-0Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0530882A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypertension		Wellbutrin Xl Adderall	PS C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513769-2Report Type:Periodic  
 Age:35 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530888A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice per day	4 DAY	Pain In Extremity		Excedrin Pm	C		

Date:11/30/04ISR Number: 4513770-9Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530904A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Wellbutrin Xl	PS	Glaxosmithkline	

Date:11/30/04ISR Number: 4513771-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530908A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY			No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513772-2Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0530922A

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Convulsion		Wellbutrin Narcotic	PS C	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513773-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0530927A  
 Age: Gender: I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
450MG Per day		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513775-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0531025A  
 Age:39 YR Gender:Female I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
150MG Twice per day	11 DAY	Bruxism Headache Pain In Jaw		Wellbutrin Xl	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513776-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531033A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
4 YR		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Drug Interaction		Choline Inositol	SS SS		

Date:11/30/04ISR Number: 4513777-1Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0531178A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Intraocular Pressure Increased		Wellbutrin Xl No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513778-3Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531182A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	6 WK	Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Libido Decreased Social Avoidant Behaviour		Lexapro Ibuprofen	C C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513779-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531230A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	7 DAY	Dizziness Headache Pharyngolaryngeal Pain Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513781-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531264A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
7 DAY				Prozac	C		

Date:11/30/04ISR Number: 4513782-5Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531285A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	5 DAY	Palpitations		Synthroid	C	Glaxosmithkline	
		Swollen Tongue					

Date:11/30/04ISR Number: 4513783-7Report Type:Periodic  
Age:51 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0531287A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Weight Decreased		No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513784-9Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531288A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4 WK	Headache		Paxil	C	Glaxosmithkline	ORAL
20MG Unknown		Malaise		Aspirin	C	Glaxosmithkline	
		Nausea		Cold Medication	C		

Date:11/30/04ISR Number: 4513785-0Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531296A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	10 WK	Eye Swelling		Synthroid	C	Glaxosmithkline	
		Rash		Allegra D	C		

Date:11/30/04ISR Number: 4513786-2Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531333A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Therapeutic Response		Abilify	C		
		Unexpected		Univasc	C		
				Maxzide	C	Glaxosmithkline	
				Topamax	C		

Date:11/30/04ISR Number: 4513787-4Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531413A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 MON						

				Lamictal	C	Glaxosmithkline	ORAL
UNKNOWN				Lorazepam	C		
				Keppra	C		
UNKNOWN				Unknown Medication	C		

Date:11/30/04ISR Number: 4513788-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0531448A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513789-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0531450A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Myalgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513790-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0531535A  
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY	Headache					
		Hyperhidrosis					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513791-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531547A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:11/30/04ISR Number: 4513792-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531548A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							

Date:11/30/04ISR Number: 4513793-XReport Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531577A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 18 MON				Wellbutrin Sr	SS	Glaxosmithkline	ORAL
150MG Twice							
per day				Depakote	C		
				Lorazepam	C		
				Remeron	C		
				Prevacid	C		

Date:11/30/04ISR Number: 4513794-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531597A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day		Weight Decreased					

Date:11/30/04ISR Number: 4513795-3Report Type:Periodic Company Report #S04-USA-06882-01  
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Drug Interaction		Namenda	SS		
UNKNOWN				Aricept	C		
UNKNOWN	5MG Per day						

Date:11/30/04ISR Number: 4513796-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0531730A  
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 1 WK				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513797-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0531781A  
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 5 DAY				No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513798-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531819A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Decreased		Wellbutrin Xl Depakote	PS C	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513799-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531821A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Xanax	C		

Date:11/30/04ISR Number: 4513800-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531830A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Oestrogen Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Singular	C		
				Ambien	C		
				Viagra	C		
				Prednisone	C		
				Zyrtec	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513801-6Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531840A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK			No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513802-8Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531851A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
30MG Per day	1	MON					
		Asthenia		Prednisone	C		
		Crying		Amitriptyline	C		
		Feeling Jittery		Nexium	C		
		Stress		Singulair	C		
		Tremor		Amoxicillin	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513803-XReport Type:Periodic  
Age:19 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531853A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							
		Salivary Hypersecretion		Lithium	C	Glaxosmithkline	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513804-1Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531857A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG		Blood Oestrogen Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Alternate

days

Date:11/30/04ISR Number: 4513805-3Report Type:Periodic  
 Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532209A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Blister		Wellbutrin	PS	Glaxosmithkline	ORAL
UNKNOWN		Migraine 1 MON		Wellbutrin	SS	Glaxosmithkline	
		Pruritus					

Date:11/30/04ISR Number: 4513806-5Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0532256A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513809-0Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0532257A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Photosensitivity Reaction		Wellbutrin Xl Cymbalta	PS C	Glaxosmithkline	ORAL

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown	1 YR	Chest Discomfort		Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB In the morning	1 YR			Paxil	SS	Glaxosmithkline	ORAL
SUBCUTANEOUS	35UNIT In the morning			Humulin N	C		
200MG Per day				Antihypertensives	C		
20MG At night				Prednisone	C		ORAL
40MG Per day				Plaquenil	C		ORAL
				Aciphex	C		ORAL
				Detrol La	C		
				Lasix	C	Glaxosmithkline	ORAL

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	4 WK	Anorexia Dry Mouth Nausea Weight Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513812-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524917A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged			Lorazepam	C		
			Alcohol	C		

Date:11/30/04ISR Number: 4513813-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524918A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged						

Date:11/30/04ISR Number: 4513814-4Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524919A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
300MG Per day	Hot Flush		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
UNKNOWN	Paraesthesia		Xanax	C		

Date:11/30/04ISR Number: 4513815-6Report Type:Periodic  
Age: Gender:I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525030A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Night Sweats		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513816-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525034A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					

3	WK	Drug Interaction	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Metrorrhagia	Birth Control Pill	C		ORAL

Date:11/30/04ISR Number: 4513817-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525046A  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice		Medication Error					
per day	2	DAY					

Date:11/30/04ISR Number: 4513818-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525050A  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	9	MON		Paroxetine	C	Glaxosmithkline	
		Pharmaceutical Product		Synthroid	C	Glaxosmithkline	
		Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513819-3Report Type:Periodic  
 Age:42 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0525053A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
450MG Per day	7	MON	Drug Ineffective	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Stool Analysis Abnormal	Trileptal	C		
				Zoloft	C		
				Trazodone	C		

Date:11/30/04ISR Number: 4513820-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525056A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day			Cough	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513821-1Report Type:Periodic  
 Age:81 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525057A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2	DAY	Pruritus	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513822-3Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525058A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	9	WK	Weight Increased	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513823-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525072A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oropharyngeal Swelling Swelling Face Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513824-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525077A  
 Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Toothache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG	Unknown 4 MON			No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513825-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525175A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513826-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525182A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT
		Insomnia Irritability

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nervousness

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
13	DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513827-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525234A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513828-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525237A  
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4 DAY	Myalgia		Celexa	C		
				Synthroid	C	Glaxosmithkline	
				Estradiol	C		
				Maxzide	C	Glaxosmithkline	
				Xanax	C		
				Zocor	C		

Date:11/30/04ISR Number: 4513829-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525239A  
 Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	18 MON	Urticaria		Zoloft	C		

Date:11/30/04ISR Number: 4513830-2Report Type:Periodic  
Age:19 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525242A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Inappropriate Affect		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 MON			Penicillin	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513831-4Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525243A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	20 DAY			Levoxyl	C	Glaxosmithkline	
				Zocor	C		

Date:11/30/04ISR Number: 4513832-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525261A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513833-8Report Type:Periodic  
Age:32 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525407A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	20 DAY	Urticaria					

Date:11/30/04ISR Number: 4513836-3Report Type:Periodic  
Age:23 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525424A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
75MG Single		Medication Error					
dose		Pruritus		No Concurrent			
		Swollen Tongue		Medication	C		

Date:11/30/04ISR Number: 4513837-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525428A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Benign Prostatic		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hyperplasia					
		Dry Mouth					

Date:11/30/04ISR Number: 4513838-7Report Type:Periodic  
Age:51 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0525430A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	9 MON	Paraesthesia		Serzone	C		
		Tinnitus		Klonopin	C		
				Lamictal	C	Glaxosmithkline	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513842-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525573A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Unknown			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Heart Rate Increased					
		Palpitations		Ssri	C		

Date:11/30/04ISR Number: 4513843-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525576A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	450MG per day 1 YR			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Drug Ineffective					
		Stool Analysis Abnormal					

Date:11/30/04ISR Number: 4513844-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525596A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Urticaria					

Date:11/30/04ISR Number: 4513845-4Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0525608A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Rash					

Date:11/30/04ISR Number: 4513846-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525609A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day 2 MON			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Drug Ineffective					

Date:11/30/04ISR Number: 4513847-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525630A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3 WK			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dizziness					
		Insomnia		Detrol La	C		
		Mental Impairment					

Date:11/30/04ISR Number: 4513848-XReport Type:Periodic  
Age:59 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525631A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Rash Papular					
per day	28 DAY			No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513849-1Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525632A

Outcome	PT
	Headache
	Hot Flush

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nausea

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	5 WK		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513850-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525775A  
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	4 MON	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dry Eye		No Concurrent Medication	C		
		Dry Mouth					
		Pharmaceutical Product Complaint					
		Stool Analysis Abnormal					

Date:11/30/04ISR Number: 4513851-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525798A  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Unknown	4 WK	Arthralgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Rash		Baby Aspirin	C	Glaxosmithkline	
		Swelling Face		Vitamin	C		

Date:11/30/04ISR Number: 4513852-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525800A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513853-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525803A  
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3 DAY	Feeling Drunk		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Accupril	C		
				Norvasc	C		
				Synthroid	C	Glaxosmithkline	
				Lexapro	C		

Date:11/30/04ISR Number: 4513855-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525826A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513856-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525832A  
Age: Gender:Female I/FU:I

Outcome PT  
Dizziness  
Dry Mouth  
Nausea

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Pollakiuria Tinnitus Vertigo	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	24 HR			Topomax	C		
				Clarinet	C		
				Flonase	C	Glaxosmithkline	
				Astelin	C		
				Birth Control Pills	C		

Date:11/30/04ISR Number: 4513857-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525940A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Unknown							

Date:11/30/04ISR Number: 4513858-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525953A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Unknown	6 MON						

Date:11/30/04ISR Number: 4513859-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525954A  
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pollakiuria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	13 DAY						

Date:11/30/04ISR Number: 4513860-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525993A  
Age: Gender:Female I/FU:I

Outcome		PT		Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Rash			Wellbutrin	PS	Glaxosmithkline	

Date:11/30/04ISR Number: 4513861-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525994A  
 Age: Gender:Female I/FU:F

Outcome		PT		Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Nausea			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Vomiting			Zoloft	C		

Date:11/30/04ISR Number: 4513862-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525997A  
 Age:51 YR Gender:Male I/FU:I

Outcome		PT		Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Insomnia			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	5 DAY	Muscle Tightness			Nexium	C		

Date:11/30/04ISR Number: 4513863-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0526000A  
 Age: Gender:Female I/FU:I

Outcome		PT		Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Headache			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	15 WK				Insulin	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Synthroid C Glaxosmithkline

Date:11/30/04ISR Number: 4513864-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526001A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Ill-Defined Disorder		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513865-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526008A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice		Dizziness					
per day	3 WK	Influenza Like Illness		Zoloft	C		
		Nausea		Singulair	C		
		Palpitations		Prilosec	C	Glaxosmithkline	
				Multivitamin	C		
				Omega 3 Fatty Acids	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513866-1Report Type:Periodic  
Age:40 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0526012A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Stomach Discomfort		Wellbutrin Sr	C	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513867-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526013A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Prozac	C		

Date:11/30/04ISR Number: 4513868-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0526030A  
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	12 DAY			No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513869-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0526129A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
5	MON						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513870-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526134A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
10	WK						

Date:11/30/04ISR Number: 4513871-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526135A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypertension		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG	Unknown						

Date:11/30/04ISR Number: 4513872-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526138A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG	Unknown 1 MON						
UNKNOWN		Photosensitivity Reaction		Levitra	SS	Glaxosmithkline	
		2 DAY					

Date:11/30/04ISR Number: 4513873-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526179A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG	Per day 3 WK						
		Muscle Spasms		Metoprolol	C		
				Norvasc	C		
				Klonopin	C		
				Semara	C		
				Zoloft	C		

Date:11/30/04ISR Number: 4513874-0Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0526210A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:11/30/04ISR Number: 4513875-2Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526213A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 19 DAY							
				Pro-Gest	C		

Date:11/30/04ISR Number: 4513876-4Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526416A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dry Eye		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 4 MON							
		Irritability		No Concurrent			
		Stool Analysis Abnormal		Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513877-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526437A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Xanax	C		

Date:11/30/04ISR Number: 4513878-8Report Type:Periodic  
 Age:28 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526445A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 1 YR				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513879-XReport Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0526449A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513880-6Report Type:Periodic  
 Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526450A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 1 WK		Feeling Hot		Allegra	C		
				Levoxyl	C	Glaxosmithkline	
				Monopril	C		
				Hctz	C		
				Celebrex	C		
				Crestor	C		

Date:11/30/04ISR Number: 4513881-8Report Type:Periodic  
Age:45 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0526636A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
15MG	Unknown			Adderall Xr	C		ORAL
				Oral Contraceptive	C		ORAL

Date:11/30/04ISR Number: 4513882-XReport Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526661A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG	Per day 3 MON	Palpitations		Magnesium Oxide	C		
				Fish Oil	C		

Date:11/30/04ISR Number: 4513883-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526662A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513884-3Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526674A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	6 WK	Thrombophlebitis		Effexor	C		
		Superficial					

Date:11/30/04ISR Number: 4513885-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526678A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Crying					
		Weight Decreased					

Date:11/30/04ISR Number: 4513886-7Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526700A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
400MG Per day	5 WK	Drug Ineffective					

Date:11/30/04ISR Number: 4513887-9Report Type:Periodic  
Age:36 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0526704A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
238 DAY		Alopecia		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513888-0Report Type:Periodic  
Age:40 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0526706A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	8 DAY	Rash Generalised Urticaria					

Date:11/30/04ISR Number: 4513889-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0526798A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 MON	Drug Interaction		Zoloft	SS		ORAL
200MG Per day	3 YR	Vomiting					

Date:11/30/04ISR Number: 4513890-9Report Type:Periodic Company Report #A0526799A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia Hallucination		Wellbutrin Xl	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513891-0Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526800A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1 WK	Abdominal Distension Arthralgia Constipation Dizziness Flatulence Sensation Of Pressure		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513892-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526834A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	8 WK	Bruxism Toothache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513893-4Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526850A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3 DAY	Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Trazodone	C		

Date:11/30/04ISR Number: 4513894-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526862A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Prednisone	C		

Date:11/30/04ISR Number: 4513895-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526868A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513896-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526872A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Flushing		Abilify	C		
150MG Per day	1 DAY	Head Discomfort		Zyprexa	C		
		Mania		Prozac	C		
				Klonopin	C		

Date:11/30/04ISR Number: 4513898-3Report Type:Periodic  
Age:51 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0527093A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR	Constipation		Trileptal	C		
		Drug Ineffective		Lorazepam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513899-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527107A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	7 DAY	Pollakiuria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513900-9Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527121A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	2 MON	Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Muscle Spasms		No Concurrent Medication	C		
		Tinnitus					

Date:11/30/04ISR Number: 4513901-0Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527126A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3 WK	Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Xanax	C		

Date:11/30/04ISR Number: 4513902-2Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527286A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	4 MON	Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Vicodin	C		
				Lexapro	C		

Date:11/30/04ISR Number: 4513903-4Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527295A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Swelling Face		Plaquenil Ibuprofen	C C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513904-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527317A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513905-8Report Type:Periodic  
Age:48 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0527322A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513906-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527332A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
1	MON						

Date:11/30/04ISR Number: 4513907-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527387A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513908-3Report Type:Periodic  
 Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527389A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	7 WK	Anxiety		Estrogen Replacement	C		
		Tremor		Ativan	C		

Date:11/30/04ISR Number: 4513909-5Report Type:Periodic  
 Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527396A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3	WK			Paxil	SS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513910-1Report Type:Periodic  
Age:68 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527406A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	6 WK	Headache		Lisinopril	C		
				Hct	C		

Date:11/30/04ISR Number: 4513911-3Report Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527408A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	18 MON			Remeron	C		
				Depakote	C		
				Lorazepam	C		
				Prevacid	C		
				Lisinopril	C		

Date:11/30/04ISR Number: 4513912-5Report Type:Periodic  
Age:42 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0527409A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Balance Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK	Feeling Abnormal		Lipitor	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Zetia C  
 Nexium C  
 Lexapro C

Date:11/30/04ISR Number: 4513913-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0527425A  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	6	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Alopecia		Buspar	C		
				Vistaril	C		
				Celebrex	C		

Date:11/30/04ISR Number: 4513914-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0527511A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hallucination		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513915-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0527513A  
 Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Diarrhoea Nausea Vomiting		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513916-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0527514A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown	5	DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Sleep Disorder					

Date:11/30/04ISR Number: 4513917-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527538A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
1 YR		Hypersomnia					

Date:11/30/04ISR Number: 4513920-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527550A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG At		Oedema Peripheral					
night	2 DAY	Palmar Erythema					
		Pruritus					
		Tremor					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513922-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527586A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaphylactic Reaction		Wellbutrin No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513923-XReport Type:Periodic  
Age:63 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0527590A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK	Diarrhoea Hallucination		No Concurrent Medication	 C		

Date:11/30/04ISR Number: 4513924-1Report Type:Periodic  
Age:18 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0527766A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	22 DAY	Rash Urticaria		Wellbutrin Xl  No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513925-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527967A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	2 WK	Dizziness Nervousness		Wellbutrin Xl  Ibuprofen	PS  C	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513926-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527978A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 WK			Lexapro	C		

Date:11/30/04ISR Number: 4513927-7Report Type:Periodic  
Age:37 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0527981A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	9 DAY	Pruritus Throat Tightness Urticaria		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513928-9Report Type:Periodic  
Age:77 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528110A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Twice		Insomnia		Sinemet	C		
per day	9 MON			Clonazepam	C		
				Remeron	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513929-0Report Type:Periodic  
Age:77 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528118A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	1 DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dizziness		Valium	C		
		Sedation		Tenormin	C		
				Nexium	C		
				Fioricet	C		

Date:11/30/04ISR Number: 4513930-7Report Type:Periodic  
Age:37 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0528128A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dry Mouth		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513931-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0528236A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Anxiety		5-Htp	C		
		Bruxism					
		Dry Mouth					
		Insomnia					

Date:11/30/04ISR Number: 4513932-0Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528375A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Rash		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513933-2Report Type:Periodic  
Age:77 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0528385A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4	Feeling Abnormal		Sinemet	C		
				Clonazepam	C		
				Remeron	C		

Date:11/30/04ISR Number: 4513936-8Report Type:Periodic  
Age:66 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0528389A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Myalgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	6	Pain In Extremity		Gabitril	C		
				Prozac	C		
				Evista	C		
				Klonopin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513937-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528421A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6	WK					

Date:11/30/04ISR Number: 4513938-1Report Type:Periodic  
Age:51 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528423A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	4	WK					
		Drug Ineffective		Paxil	SS	Glaxosmithkline	ORAL
25MG Per day	18	MON					
		Loss Of Libido		Percocet	C		
				Prilosec	C	Glaxosmithkline	
				Lisinopril	C		
				Lipitor	C		
				Alprazolam	C		
				Ibu	C	Glaxosmithkline	
				Viagra	C		

Date:11/30/04ISR Number: 4513939-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528472A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
4	MON						

Date:11/30/04ISR Number: 4513940-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528474A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Unknown	1	YR					
		Skin Odour Abnormal					

Date:11/30/04ISR Number: 4513941-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528476A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Interaction		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Unknown							
		Tachycardia		Adderall Xr	SS		
UNKNOWN	10MG Unknown						

Date:11/30/04ISR Number: 4513942-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528508A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Twice							
per day	2 MON						

Date:11/30/04ISR Number: 4513943-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528511A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	18 MON						
		Headache					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513944-7Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLIN-A0528522A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	4 DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Nausea		Atenolol	C		
				Estrace	C		
				Klonopin	C		

Date:11/30/04ISR Number: 4513946-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLIN-A0528533A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Drug Ineffective		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513947-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLIN-A0528542A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Hypoglycaemia					

Date:11/30/04ISR Number: 4513948-4Report Type:Periodic  
Age:7 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLIN-A0528552A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	3 WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Arthralgia		Strattera	C		
		Feeling Hot		Risperdal	C		
				Hydroxyzine	C		

Date:11/30/04ISR Number: 4513949-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528557A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	6 WK			Wellbutrin Sr	SS	Glaxosmithkline	ORAL
6 WK							

Date:11/30/04ISR Number: 4513950-2Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528795A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 DAY			Lithobid	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513951-4Report Type:Periodic  
Age:79 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528802A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gastrointestinal Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK	Nausea		Coumadin	C	Glaxosmithkline	
				Tambocor	C		
				Imuran	C	Glaxosmithkline	
				Avapro	C		
				Cardura	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513952-6Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528809A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK			Synthroid	C	Glaxosmithkline	
				Xanax	C		

Date:11/30/04ISR Number: 4513953-8Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528985A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	7 DAY			Flonase	C	Glaxosmithkline	
		Paraesthesia		Tylenol	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513954-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529007A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK						
		Headache					
		Insomnia					
		Nausea					

Date:11/30/04ISR Number: 4513955-1Report Type:Periodic  
Age:39 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0529012A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG In the							
morning	5 DAY						

Date:11/30/04ISR Number: 4513956-3Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529024A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Anxiety	Wellbutrin XL	PS	Glaxosmithkline	ORAL
300MG Per day	3	WK	Depression	Unknown Medication	C		
				Activella	C		
				Trazodone	C		

Date:11/30/04ISR Number: 4513957-5Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529089A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Nightmare	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	7	DAY		Ethanol	SS		
				Actos	C		
				Accupril	C		
				Xanax	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513958-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529099A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	5 DAY			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Back Pain					
		Pain In Extremity		Blood Pressure Medication	C		
				Vitamins	C		
				Diazepam	C		

Date:11/30/04ISR Number: 4513959-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529331A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
400MG Per day				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dysgeusia					
		Nausea		Adderall Xr	C		
		Therapeutic Response		Ambien	C		
		Unexpected		Buspirone	C		
		Weight Decreased		Clarinet	C		
				Lithium Carbonate	C	Glaxosmithkline	
				Nexium	C		
				Pravachol	C		
				Tarka	C		
				Diovan	C		

Date:11/30/04ISR Number: 4513960-5Report Type:Periodic  
Age:14 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529347A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 WK			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Urticaria					
				Paxil Cr	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513961-7Report Type:Periodic  
Age:74 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529352A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Blood Glucose Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Insomnia		Diabeta	C		
				Avandia	C	Glaxosmithkline	
				Avapro	C		
				Lasix	C	Glaxosmithkline	
				Lipitor	C		

Date:11/30/04ISR Number: 4513962-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0529365A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 YR			Lisinopril	C		
				Levoxyl	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513963-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0529457A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513964-2Report Type:Periodic  
Age:41 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0529468A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	300MG Per day	Diarrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	150MG Twice	Headache		Zyban	SS	Glaxosmithkline	ORAL
	per day	Nausea					
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513965-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529474A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Three	Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	times per day	Stool Analysis Abnormal					

Date:11/30/04ISR Number: 4513966-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529478A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Diovan Hct	C		

Date:11/30/04ISR Number: 4513970-8Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0529480A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Interaction		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Increased Appetite		Periactin	SS		

UNKNOWN

Date:11/30/04ISR Number: 4513971-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529492A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 WK		Burning Sensation Pruritus Rash Visual Disturbance		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513972-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529493A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anorgasmia Erectile Dysfunction		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513973-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529496A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513974-5Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0529497A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	5 DAY	Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513975-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529499A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	3 MON	Depressed Mood		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Zoloft	C		

Date:11/30/04ISR Number: 4513976-9Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529508A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MCG Per day	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	10 WK	Depression		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513977-0Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529518A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	Bipolar Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513978-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0529520A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513979-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0529537A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 2 WK				Advair	C	Glaxosmithkline	
				Ritalin	C		

Date:11/30/04ISR Number: 4513980-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0529672A  
 Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
450MG Unknown		Myoclonus		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Unknown 128 DAY		Tremor		Zoloft	C		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513981-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529676A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Cold Sweat					
		Dyspnoea		Xanax	C		
		Headache		Allegra	C		
		Hyperhidrosis		Flonase	C	Glaxosmithkline	
		Nasal Discomfort					
		Pharyngolaryngeal Pain					
		Tachycardia					
		Tremor					

Date:11/30/04ISR Number: 4513982-4Report Type:Periodic  
Age:31 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529683A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3 WK			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dysuria					
		Rash		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513983-6Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529689A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300MG Per day	2 MON			Wellbutrin	PS	Glaxosmithkline	ORAL
		Amnesia					
		Convulsion		Astelin	C		
		Grand Mal Convulsion		Nasonex	C		
				Singulair	C		
				Clarinet	C		

Date:11/30/04ISR Number: 4513984-8Report Type:Periodic  
Age:19 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0529693A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day	6 WK	Insomnia		Testosterone Injection	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4513985-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0529699A  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 DAY	Decreased Appetite		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Feeling Jittery		Synthroid	C	Glaxosmithkline	
		Insomnia					

Date:11/30/04ISR Number: 4513986-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0529926A  
 Age:3 WK Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1 DAY	Vomiting Projectile		Wellbutrin Xl	PS	Glaxosmithkline	
				Prenatal Vitamins	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513987-3Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529927A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
4 WK		Medication Error		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Overdose Vomiting		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513989-7Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529951A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Twice per day	4 MON	Abdominal Distension Diarrhoea Flatulence		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Glucophage	C		
				Synthroid	C	Glaxosmithkline	
				Multivitamin	C		
				Magnesium	C		
				Potassium	C		
				Zinc	C		

Date:11/30/04ISR Number: 4513990-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529959A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4513991-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529993A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown		Asthma Conjunctivitis Allergic		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Migraine  
Pain

Date:11/30/04ISR Number: 4513992-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529995A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3	WK						

Date:11/30/04ISR Number: 4513993-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530002A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG	Unknown						

Date:11/30/04ISR Number: 4513994-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530003A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Postural Orthostatic Hypotension		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4513997-6Report Type:Periodic  
Age:17 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530013A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	6 DAY	Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Agitation Heart Rate Increased Palpitations		Concerta	C		

Date:11/30/04ISR Number: 4513998-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530019A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 WK	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4513999-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530030A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4514000-4Report Type:Periodic  
Age:11 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530031A

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	6 WK	Aggression		Wellbutrin	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4514001-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530042A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	6	MON	Drug Ineffective	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Insomnia	No Concurrent Medication	C		

Date:11/30/04ISR Number: 4514002-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530114A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Unknown			Dizziness	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Unknown				Wellbutrin Sr	SS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4514003-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530115A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
UNKNOWN			Tremor	Wellbutrin Xl Buspar	PS SS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4514004-1Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530192A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 DAY	Heart Rate Increased		Ambien	C		
				Elavil	C	Glaxosmithkline	
				Xanax	C		

Date:11/30/04ISR Number: 4514005-3Report Type:Periodic  
Age:30 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530197A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice		Amnesia					
per day	3 MON	Chest Discomfort		Serzone	C		
		Confusional State		Remeron	C		
		Insomnia					

Date:11/30/04ISR Number: 4514006-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530218A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN							

Date:11/30/04ISR Number: 4514007-7Report Type:Periodic  
Age:62 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530238A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	10 DAY	Insomnia		Klonopin	C		
		Polyuria		Cozaar	C		

Weight Decreased

Lipitor  
Proscar

C  
C

Date:11/30/04ISR Number: 4514008-9Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530246A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	8	WK		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4514009-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530315A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Adverse Event			

Date:11/30/04ISR Number: 4514010-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530318A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Constipation			



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4514011-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530320A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Per day			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dizziness					
		Nausea		Paxil	C	Glaxosmithkline	
				Wellbutrin	C	Glaxosmithkline	

Date:11/30/04ISR Number: 4514012-0Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530366A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Per day			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	4 DAY	Insomnia					
				Thyroid	C		

Date:11/30/04ISR Number: 4514013-2Report Type:Periodic  
Age:60 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530367A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Per day			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Abdominal Pain					
		Abnormal Dreams		Lisinopril	C		
				Toprol Xl	C		
				Lipitor	C		
				Folic Acid	C		
				Baby Aspirin	C	Glaxosmithkline	
				Multivitamin	C		

Date:11/30/04ISR Number: 4514014-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530385A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	300MG Per day			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	8 MON	Insomnia					
				Effexor	C		

Date:11/30/04ISR Number: 4514015-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530389A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK						

Date:11/30/04ISR Number: 4514016-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530485A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Wellbutrin Sr	C	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4514017-XReport Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530494A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decreased Appetite		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hangover		Amphetamine	C		
		Somnolence		Dexatrim	C	Glaxosmithkline	

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Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4514019-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530611A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day	Libido Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4514020-XReport Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530612A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day 2 MON	Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Weight Decreased		No Concurrent Medication	C		

Date:11/30/04ISR Number: 4514021-1Report Type:Periodic  
Age:32 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530618A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG Per day 10 DAY	Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:11/30/04ISR Number: 4514022-3Report Type:Periodic  
Age:19 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530631A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day 4 MON	Disturbance In Attention		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4514023-5Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0530641A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day							

Date:11/30/04ISR Number: 4514024-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0530643A  
 Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 25 DAY							
				No Concurrent Medication	C		

Date:11/30/04ISR Number: 4514025-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0530654A  
 Age:16 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day							

Date:11/30/04ISR Number: 4514026-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0530672A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							
				Wellbutrin Sr	SS	Glaxosmithkline	ORAL
				Trazodone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Zyrtec	C	Glaxosmithkline	ORAL
Advair	C	Glaxosmithkline	
Nexium	C		ORAL
Lexapro	C		ORAL

Date:11/30/04ISR Number: 4514043-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0531529A  
 Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Lipase Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Per day				Klonopin	C		ORAL
.5MCG At							
night				B Complex	C		
				Citracal	C		
				Vitamin E	C		

Date:11/30/04ISR Number: 4514047-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0535145A  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Constipation		Wellbutrin Xl	PS	Glaxosmithkline	
150MG Per day	8 DAY	Proctalgia		Concerta	C		
		Rectal Haemorrhage					
		Stool Analysis Abnormal					

Date:11/30/04ISR Number: 4514048-XReport Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0535220A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Depressed Level Of		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
300MG Per day	1 DAY	Consciousness		Ritalin	C		ORAL
		Dizziness					

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Autosomal Chromosome		Zyban	PS	Glaxosmithkline	
150MG Per day 1	MON					
Congenital Anomaly	Anomaly		Meteospasmyl	C		
	Chromosome Abnormality		Primperan	C	Glaxosmithkline	
	Congenital Hand		Vitamin C	C	Glaxosmithkline	
	Malformation		Ginkor Fort	C		
	Dextrocardia		Stresam	C		
3UNIT per day						
	Diaphragmatic Hernia					
	Drug Exposure During					
	Pregnancy					
	Foetal Growth Retardation					
	Foetal Malformation					
	Gastrointestinal Disorder					
	Trisomy 15					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/04ISR Number: 4514073-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0358400A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG	Unknown		Accommodation Disorder	Zyban	PS	Glaxosmithkline	ORAL
			Visual Acuity Reduced				

Date:11/30/04ISR Number: 4517586-9Report Type:Expedited (15-DaCompany Report #2004AL001522

Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
Initial or Prolonged							
Required							
Intervention to							
Prevent Permanent							
10 MG; X1; PO			Arteriospasm Coronary	Kadian (Morphine	PS		ORAL
Impairment/Damage			Blood Pressure Increased	Sulfate Sustained			
			Self-Medication	Release) Capsules,			
			Professional	20 Mg (Alpharma)			
			Other	(Kadian (Morphine			
				Bupropion			
				Hydrochloride	SS		ORAL
				Efferalgan	C		
150 MG; PO							

Date:12/01/04ISR Number: 4515180-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0534663A

Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
Initial or Prolonged							
4 WK			Convulsion	Wellbutrin	PS	Glaxosmithkline	
			Headache	Lamictal	SS	Glaxosmithkline	ORAL
			Insomnia	Paxil	SS	Glaxosmithkline	
			Pruritus	Progesterone	C		
			Rash				
			Road Traffic Accident				
			Urticaria				

Date:12/01/04ISR Number: 4515190-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0535450A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Convulsion Depressed Level Of Consciousness Loss Of Consciousness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:12/01/04ISR Number: 4515191-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0535456A  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Medication Error		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:12/01/04ISR Number: 4515206-0Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0357320A  
Age:36 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Anxiety Dizziness Hyperhidrosis Malaise

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Medication Error Paraesthesia Sense Of Oppression	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Zyban	PS	Glaxosmithkline	ORAL
150MG Twice per day	7 DAY						

Date:12/01/04ISR Number: 4515218-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0357799A  
Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
Life-Threatening 150MG Per day Disability UNKNOWN	5 WK	Aortic Stenosis		Amoxicillin	C	Glaxosmithkline	
UNKNOWN				Erythromycin	C	Glaxosmithkline	
UNKNOWN				Iron Sulphate	C	Glaxosmithkline	
UNKNOWN				Prednisolone	C	Glaxosmithkline	
UNKNOWN				Carboplatin	C		
INTRAVENOUS		13 WK		Etoposide	C		
INTRAVENOUS		13 WK		Ibuprofen	C	Glaxosmithkline	
UNKNOWN				Prochlorpemazine	C	Glaxosmithkline	
UNKNOWN				Chemotherapy	C		

Date:12/01/04ISR Number: 4515230-8Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0358672A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	
Other UNKNOWN	150MG Twice	Convulsion					
per day		Loss Of Consciousness					

Urinary Incontinence

Date:12/02/04ISR Number: 4516838-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0350118A  
 Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day	10 DAY	Contusion Vasculitis		Zyban	PS	Glaxosmithkline	ORAL
1TAB Per day				Cartia	C	Glaxosmithkline	ORAL
RESPIRATORY (INHALATION)				Actonel Bricanyl	C C		ORAL
1G per day				Caltrate Naprosyn	C C	Glaxosmithkline	
				Panamax Sinequan Spiriva Symbicort Ostelin (Vitamin D2) Vagifem	C C C C C C	Glaxosmithkline	

Date:12/02/04ISR Number: 4516857-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0358654A  
 Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Balance Disorder Dizziness Dizziness Postural Visual Acuity Reduced		Zyban No Concurrent Medication	PS C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/03/04ISR Number: 4518063-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0535910A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	1 WK		Serotonin Syndrome	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Initial or Prolonged	8 MON			Lexapro	C		
	8 MON			Buspar	C		

Date:12/03/04ISR Number: 4518069-2Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0346147A

Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1TAB Per day		Tremor	Zyban	PS	Glaxosmithkline	ORAL
			Vision Blurred	Contraceptive			
			Visual Acuity Reduced	Unspecified	C		ORAL

Date:12/03/04ISR Number: 4518076-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0358058A

Age:61 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG per day 0 DAY		Chest Pain	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged	40MG per day		Chills	Pravastatin	C		ORAL
			Circulatory Collapse				
			Dyspnoea				
			Headache				
			Paraesthesia				

Date:12/03/04ISR Number: 4521302-4Report Type:Expedited (15-DaCompany Report #2004-05016

Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Other	Drug Interaction	Foreign	Clomipramine		
	Electroencephalogram	Literature	Hydrochloride		
	Abnormal	Health	(Watson		
	Grand Mal Convulsion	Professional	Laboratories)		
		Other	(Clomipramine	PS	ORAL
25 MG, DAILY,					
ORAL			Bupropion		
			Hydrochloride		
			Sustained Release		
			(Bupropion		
			Hydrochloride)	SS	ORAL
SEE IMAGE					
			Risperidone		
			(Risperidone)	C	

Date:12/06/04ISR Number: 4519299-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0535383A  
Age:63 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Accidental Overdose		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice						
Initial or Prolonged	Aphasia					
per day 4 YR						
	Bradyphrenia		Vitamin	C		
	Fall		Advil	C	Glaxosmithkline	
	Immobile		Ambien	C		
	Insomnia		Calcium	C		
	Speech Disorder					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/06/04ISR Number: 4519300-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0535398A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3	WK	Medication Error				
				Thyroid	C		
				Metoprolol	C		
				Detrol La	C		
				Avapro	C		
				Paxil	C	Glaxosmithkline	
				Premarin	C		

Date:12/06/04ISR Number: 4519317-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0358402A

Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	15	DAY					
5MG per day			Vitreous Floaters	Nitrazepam	C		ORAL
				Beclomethasone	C	Glaxosmithkline	
100MCG per							
day				Salbutamol	C	Glaxosmithkline	
100MCG per							
day							

Date:12/06/04ISR Number: 4519334-5Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0358995A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eye Pain		Zyban	PS	Glaxosmithkline	
8	DAY						
			Gaze Palsy				
			Headache				

Insomnia  
Nausea

Date:12/06/04ISR Number: 4520402-2Report Type:Direct Company Report #CTU 233713  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Bupropion Sa 150 Mg	PS		ORAL
150MG 2 X		Pharmaceutical Product					
DAY ORAL		Complaint					

Date:12/06/04ISR Number: 4521764-2Report Type:Expedited (15-DaCompany Report #L04-USA-07403-24  
Age:30 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Literature	Escitalopram	PS		
		Multiple Drug Overdose	Health	Bupropion (Long			
			Professional	Acting) (Bupropion)	SS		
				Paroxetine	SS		

Date:12/07/04ISR Number: 4520318-1Report Type:Expedited (15-DaCompany Report #A03200400600  
Age:48 YR Gender:Male I/FU:F

Outcome  
Life-Threatening  
Hospitalization -  
  
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Initial or Prolonged  
Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Suicide Attempt		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
5MG See				Ambien	SS		ORAL
dosage text	9 DAY			Paroxetine Hydrochloride	C	Glaxosmithkline	ORAL
4 DAY							

Date:12/07/04ISR Number: 4520327-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0535565A  
Age:45 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day	1 WK	Incontinence		Wellbutrin	PS	Glaxosmithkline	ORAL
					Lamictal	C	Glaxosmithkline	
					Trazodone	C		

Date:12/07/04ISR Number: 4520350-8Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0359055A  
Age:40 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day	1 DAY	Angioneurotic Oedema		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged			Erythema		Zoloft	C		
UNKNOWN			Feeling Hot		Lanzo	C		
UNKNOWN			Paraesthesia Sensation Of Foreign Body Swelling Face					

Date:12/07/04ISR Number: 4523131-4Report Type:Direct  
Age:62 YR Gender:Male I/FU:I

Company Report #CTU 233742

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required	1 TAB DAILY - ORAL	Amnesia Cataract		Lipitor 200 Mg Pfizer Us	PS	Pfizer Us	ORAL
Intervention to Prevent Permanent Impairment/Damage	1 YR	Impaired Work Ability		Welbutrin Aspirin Therapy Welchol	SS C C		

Date:12/08/04ISR Number: 4521259-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0515139A  
Age:52 YR Gender:Female I/FU:F

Outcome	PT
Other	Asthenia Chest Pain Chills Cough Disorientation Dizziness Dysphagia Feeling Abnormal Headache Hyperhidrosis Ill-Defined Disorder Increased Tendency To Bruise Mucous Membrane Disorder



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	4 WK	Muscle Atrophy Muscle Spasms Musculoskeletal Disorder	Zyban	PS	Glaxosmithkline	ORAL
		Nausea				
		Nervousness				
		Pain Of Skin	Prednisone	C		
		Pneumonitis	Premarin	C		
		Pruritus	Medroxyprogesterone	C		
		Pyrexia	Benadryl	C	Glaxosmithkline	
		Tonsillar Cyst	Maxair	C		
		Tremor	Serevent	C	Glaxosmithkline	
		Urticaria	Azmacort	C		
		Vasodilatation				
		Wheezing				

Date:12/08/04ISR Number: 4521263-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0533147A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
		Balance Disorder					
		Diarrhoea					
		Dizziness					
		Hot Flush					
		Mood Altered					
		Oedema Peripheral					
		Pruritus					
		Sleep Disorder					
		Speech Disorder					
		Swelling Face					
		Swollen Tongue					
		Urticaria					

Date:12/08/04ISR Number: 4521273-0Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0350764A

Age:27 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization - Cardiospasm  
 150MG Unknown 30 DAY  
 Initial or Prolonged Chest Pain  
 10MG Per day 1 DAY  
 1 DAY

Zyban PS Glaxosmithkline ORAL  
 Actiskenan SS Glaxosmithkline ORAL  
 Efferalgan C Glaxosmithkline ORAL

Date:12/08/04ISR Number: 4521705-8Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #CTU 233876

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Depression Pharmaceutical Product		Generic Wellbutrin Sr 150 (Budeprion Sr)	PS	Teva	ORAL
1 PO BID		Complaint		Glucophage Amaryl Ativan Mobic	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/10/04ISR Number: 4523033-3Report Type:Expedited (15-DaCompany Report #US-ABBOTT-04P-163-0282509-00  
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Grand Mal Convulsion		Depakote	PS		
				Depakote	SS		
				Bupropion			
				Hydrochloride	SS		
				Promethazine			
				Hydrochloride	C		
				Metoclopramide	C		

Date:12/10/04ISR Number: 4523685-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0323413A  
Age:22 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged dosage text	10 DAY	Arthralgia		Zyban	PS	Glaxosmithkline	ORAL
		C-Reactive Protein					
		Increased					
		Drug Exposure During					
		Pregnancy					
		Face Oedema					
		Joint Swelling					
		Neutrophil Percentage					
		Increased					
		Odynophagia					
		Oedema					
		Pregnancy					
		Urticaria					
		White Blood Cell Count					
		Increased					

Date:12/10/04ISR Number: 4523701-3Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0358672A  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Twice		Convulsion		Zyban	PS	Glaxosmithkline	ORAL

Other  
per day  
300MCG per  
day

Loss Of Consciousness  
Myalgia  
Urinary Incontinence

Flixonase  
C  
Glaxosmithkline

Date:12/10/04ISR Number: 4523706-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0359029A  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Twice Hospitalization - per day	20 DAY	Alcohol Interaction		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Depressed Mood					
750MG per day		Medication Error Overdose		Alcohol Famvir	SS C	Glaxosmithkline	ORAL
		Suicidal Ideation					

Date:12/10/04ISR Number: 4523707-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0359033A  
Age:45 YR Gender:Female I/FU:F

Outcome	PT
Other	Depressive Symptom Epistaxis Malaise

Freedom Of Information (FOI) Report

Tearfulness

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
17 DAY			Zyban	PS	Glaxosmithkline	ORAL
50MCG Twice			Serevent	C	Glaxosmithkline	
per day						
400MCG Twice			Beclomethasone	C	Glaxosmithkline	
per day						
200MCG As			Salbutamol	C	Glaxosmithkline	
required						

Date:12/10/04ISR Number: 4528227-9Report Type:Expedited (15-DaCompany Report #B0358110A  
 Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Autonomic Nervous System Imbalance Cerebral Atrophy Condition Aggravated	Foreign Literature Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride)	PS		
3 WK		Coordination Abnormal Depressed Level Of Consciousness Diarrhoea		Sertraline (Formulation Unknown) (Sertraline)	SS		
3 WK		Electrocardiogram Qt Prolonged Electrocardiogram Repolarisation Abnormality Faecal Incontinence Fall Hallucination, Auditory Hallucination, Visual Insomnia Mental Status Changes		Piracetam (Formulation Unknown) (Piracetam) Venlafaxine Hydrochloride (Formulation Unknown) (Venlafaxine Fluoxetine	SS C		

Myoclonus  
Serotonin Syndrome  
Urinary Incontinence

Date:12/13/04ISR Number: 4525381-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0536385A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Overdose		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:12/13/04ISR Number: 4525389-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0536770A

Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation Homicidal Ideation		Paxil Wellbutrin	PS SS	Glaxosmithkline Glaxosmithkline	ORAL ORAL
150MG Unknown							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/13/04ISR Number: 4525413-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0359041A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	7 DAY	Aggression		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown		Emotional Disorder		Sertraline	C		ORAL
50MG per day	204 WK	Irritability		Omeprazole	C	Glaxosmithkline	ORAL
20MG per day		Thinking Abnormal		Amitriptyline	C		ORAL
10MG per day							

Date:12/13/04ISR Number: 4526736-XReport Type:Expedited (15-DaCompany Report #A118566  
Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged (80 MG)		Atrial Fibrillation Bundle Branch Block Left	Literature Health	Ziprasidone (Ziprasidone)	PS		
Required Intervention to Prevent Permanent Impairment/Damage (100 MG)		Delirium Disorientation Electrocardiogram Qrs Complex Prolonged	Professional Company Representative	Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		
(40 MG)		Electrocardiogram Qt Corrected Interval Prolonged Electrocardiogram Qt		Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	SS		
(250 MG)		Prolonged Electrocardiogram St Segment Abnormal		Valproate Semisodium (Valproate Semisodium)	SS		
		Electrocardiogram St Segment Depression Electrocardiogram T Wave Abnormal Grand Mal Convulsion Hallucination, Visual Lethargy Multiple Drug Overdose Myocardial Infarction					

Sinus Tachycardia  
Supraventricular  
Tachycardia

Date:12/13/04ISR Number: 4527063-7Report Type:Expedited (15-DaCompany Report #2004102906  
Age:62 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Alanine Aminotransferase
Initial or Prolonged	Increased
	Amnesia
	Aspartate
	Aminotransferase
	Increased
	Autonomic Nervous System
	Imbalance
	Balance Disorder
	Cerebral Atrophy
	Condition Aggravated
	Coordination Abnormal
	Depressed Level Of
	Consciousness
	Depression
	Diarrhoea
	Drug Interaction

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Electrocardiogram Qt Prolonged					
		Faecal Incontinence					
50 MG		Fall	Foreign Literature	Sertraline (Sertraline)	PS		
300 MG		Hallucinations, Mixed					
		Hyperreflexia	Health Professional	Bupropion (Bupropion)	SS		
		Insomnia					
75 MG		Myoclonus		Venlafaxine (Venlafaxine)	SS		
		Pyrexia					
		Serotonin Syndrome		Piracetam (Piracetam)	C		
		Urinary Incontinence					

Date:12/14/04ISR Number: 4526634-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0522931A  
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Glucose Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	DAY	Fall					
150MG per day	3 MON	Grand Mal Convulsion		Ludiomil	SS		
2 MON		Joint Dislocation		Symbyax	SS		
		Loss Of Consciousness					
		Medication Error					
		Upper Limb Fracture					

Date:12/14/04ISR Number: 4527312-5Report Type:Direct Company Report #CTU 234263  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150 MG PO QD		Dizziness		Wellbutrin Xl	PS		ORAL
		Drug Ineffective					
		Feeling Abnormal					
		Pharmaceutical Product					

Complaint

Date:12/14/04ISR Number: 4527326-5Report Type:Direct  
Age:47 YR Gender:Male I/FU:I

Company Report #CTU 234306

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening			Cerebrovascular Accident	Wellbutrin Xl	150mg	PS	
DAILY							
Hospitalization -							
Initial or Prolonged							
Disability							
Required							
Intervention to							
Prevent Permanent							
Impairment/Damage							

Date:12/14/04ISR Number: 4529932-0Report Type:Expedited (15-DaCompany Report #2004103296  
Age:73 YR Gender:Female I/FU:I

Outcome	PT
Other	Anxiety
	Cataract Operation
	Drug Ineffective
	Eye Disorder
	Formication
	Post Procedural

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Complication  
Vision Blurred

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
50 MG (50 MG, 1 IN 1 D), ORAL		Health Professional	Zoloft (Sertraline)	PS		ORAL
			Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		
			Paroxetine Hydrochloride (Paroxetine Hydrochloride0 Psycholeptics (Psycholeptics)	SS		
			Provella-14 (Estrogens Conjugated, Medroxyprogesterone Acetate)	C		
			Ciclosporin (Ciclosporin)	C		
			All Otehr Therapeutic Products (All Other Therapeutic Products)	C		

Date:12/14/04  
Age: Gender:Female I/FU:I

Report Type:Periodic  
Company Report #2004241306US

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
10 MG, QD, ORAL		Psychomotor Hyperactivity Weight Increased	Consumer	Bextra (Valdecoxib) Tablet	PS		ORAL
				Wellbutrin (Amfebutamone)			

10 MG, QD,

Hydrochloride) SS  
Lexapro SS

ORAL

ORAL

Multivitamins  
(Panthenol, Retinal,  
Ergocalciferol) C  
Vitamin E C

Date:12/14/04ISR Number: 4535560-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #2004236425US

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Bextra (Valdecoxib)			
Other		Convulsion	Professional	Tablet	PS		
20 MG			Company	Wellbutrin			
			Representative	(Amfebutamone Hydrochloride)	SS		



Death UNKNOWN Renal Failure Chronic 150MG Twice Zyban PS Glaxosmithkline

per day

Date:12/15/04ISR Number: 4529950-2Report Type:Direct Company Report #CTU 234438  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300 MG (2) Q Initial or Prolonged AM	Clonic Convulsion		Wellbutrin 300 Mg Am	PS		

Date:12/15/04ISR Number: 4531026-5Report Type:Expedited (15-DaCompany Report #2004AL001522  
Age:27 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Required 10 MG; X1; PO Intervention to Prevent Permanent Impairment/Damage	Arteriospasm Coronary Self-Medication	Foreign Health Professional	Kadian (Morphin Sulfate Cr)Sustained Release 20mg	PS	Alpharma	ORAL
		Other	Bupropion Hydrochloride	SS		ORAL
			Efferalgan	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/16/04ISR Number: 4528799-4Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0140463A  
 Age:27 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	10 DAY	Amnesia		Zyban	PS	Glaxosmithkline	ORAL
		Arrhythmia					
		Cardiac Arrest					
		Confusional State					
		Energy Increased					
		Euphoric Mood					
		Feeling Abnormal					
		Feeling Jittery					
		Heart Rate Increased					
		Hypersensitivity					
		Insomnia					
		Myocardial Infarction					
		Palpitations					
		Psychomotor Hyperactivity					
		Speech Disorder					

Date:12/16/04ISR Number: 4528806-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0530819A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	1050MG Per Hospitalization - day 0 DAY	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged Disability		Overdose					
Other							

Date:12/16/04ISR Number: 4528807-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0531192A  
 Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	5 WK	Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL
		Convulsion		Trileptal	C		

Dizziness  
Insomnia  
Micturition Disorder

Mysoline  
Synthroid  
Premarin

C  
C  
C

Glaxosmithkline

Date:12/16/04ISR Number: 4528818-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0537468A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2	MON	Headache Insomnia Loss Of Consciousness Nausea Vomiting					

Date:12/16/04ISR Number: 4528833-1Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0361828A

Age:38 YR Gender:Female I/FU:I

Outcome	PT
Other	Agitation Depression Feeling Of Despair

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Restlessness Schizophrenia Tearfulness	Report Source	Product	Role	Manufacturer	Route
UNKNOWN				Zyban	PS	Glaxosmithkline	
				Haldol Decanoate	C		
				Artane	C		
UNKNOWN				Tegretol	C		

Date:12/16/04ISR Number: 4530215-3Report Type:Direct Company Report #CTU 234497  
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - PRN		Abnormal Behaviour Aggression		Cylert- Pamoline-Prn	PS		
Initial or Prolonged PRN Disability PRN		Alcohol Use		Imipramine	SS		
Required Intervention to Prevent Permanent Impairment/Damage		Anxiety		Wellbutrin	SS		
		Bereavement Reaction Disturbance In Social Behaviour Feeling Of Despair Grandiosity Insomnia Irritability Loss Of Employment Mania Mental Disorder Psychomotor Hyperactivity Relationship Breakdown Stress Suicidal Ideation					

Date:12/16/04ISR Number: 4530278-5Report Type:Direct Company Report #CTU 234493  
 Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death Completed Suicide Wellbutrin Sr 150  
 Drug Withdrawal Syndrome Mg 2x Daily Glaxo  
 Smith Kline PS Glaxo Smith Kline ORAL  
 150MG 2X  
 DAILY ORAL

Date:12/16/04ISR Number: 4535995-9Report Type:Periodic Company Report #HQWYE206123SEP04  
 Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Interaction International Normalised Ratio Decreased Thrombophlebitis	Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		ORAL
SEE IMAGE				Coumadin (Warfarin Sodium,)	SS		ORAL
SEE IMAGE				Wellbutrin Xl (Bupropion Hydrochloride, )	SS		ORAL
150 MG 1X PER 1 DAY, ORAL				Zyprexa (Olanzapine) Klonopin (Clonazepam)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/17/04ISR Number: 4530688-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0536385A  
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 450MG Per day Initial or Prolonged			Grand Mal Convulsion	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Overdose					

Date:12/17/04ISR Number: 4530706-5Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0351173A  
Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN dosage text	150MG 39 DAY	See Palpitations		Bupropion Hydrochloride	PS	Glaxosmithkline	
		Hyperthyroidism Insomnia					

Date:12/17/04ISR Number: 4530721-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0359865A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly			Drug Exposure During Pregnancy Ear Disorder	Zyban	PS	Glaxosmithkline	

Date:12/17/04ISR Number: 4530738-7Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0362084A  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN UNKNOWN	150MG per day		Angioneurotic Oedema Dermatitis Allergic	Zyban Acenocoumarol	PS C	Glaxosmithkline	
	1MG Unknown		Drug Interaction Paraesthesia Pruritus				

Tongue Disorder

Date:12/20/04ISR Number: 4532018-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0537805A  
 Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Unknown 8 DAY	Facial Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Initial or Prolonged Disability		Facial Palsy		Effexor	C		
		Facial Paresis		Paxil	C	Glaxosmithkline	
Other		Hypertension		Buspar	C		

Date:12/20/04ISR Number: 4532022-4Report Type:Expedited (15-DaCompany Report #2004/02747  
 Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Crying Depression Insomnia Palpitations Panic Reaction Suicidal Ideation Thinking Abnormal		Zyban	PS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/20/04ISR Number: 4532029-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0323980A  
Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Unknown		Brain Contusion	Zyban	PS	Glaxosmithkline	ORAL
			Confusional State	Fluanxol	SS		
			Epistaxis				
			Facial Bones Fracture				
			Fall				
			Incoherent				
			Malaise				
			Overdose				
			Pulmonary Oedema				
			Sudden Death				
			Tremor				

Date:12/20/04ISR Number: 4532037-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0357358A  
Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Asthenia	Zyban	PS	Glaxosmithkline	
			Constipation	Lisinopril	C		
			Dry Mouth				
			Ventricular Tachycardia				

Date:12/20/04ISR Number: 4532866-9Report Type:Direct Company Report #CTU 234712  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1 TAB/2 TAB		Adverse Drug Reaction	Vioxx Unknown			
			Arthralgia	Unknown	PS		ORAL
Disability	DAY/1WK/2		Drug Tolerance Increased				
			Fibromyalgia				
WK ORAL			Muscle Spasms	Wellbutrin			

1 TAB DAY Myalgia Unknown Unknown SS ORAL  
Pruritus  
ORAL Urticaria Morphine C  
Antidepressants C

Date:12/20/04ISR Number: 4533070-0Report Type:Direct Company Report #CTU 234699  
Age:46 YR Gender:Male I/FU:I

Outcome PT  
Life-Threatening Abdominal Pain Upper  
Hospitalization - Agitation  
Initial or Prolonged Anger  
Disability Anorexia  
Anxiety  
Asthenia  
Condition Aggravated  
Confusional State  
Depression  
Dizziness  
Drug Ineffective  
Dysgeusia  
Headache  
Heart Rate Increased  
Hostility  
Hyperhidrosis

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Impulsive Behaviour Insomnia Irritability				
		Libido Decreased Marital Problem Myalgia Nervousness Panic Attack Restlessness Suicide Attempt Tinnitus Tremor Vision Blurred Vomiting Weight Decreased	Wellbutrin	150 Mil.	PS	

Date:12/21/04ISR Number: 4533689-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0537281A  
Age: Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
Disability		Arthralgia	Lamictal	PS	Glaxosmithkline	ORAL
2 YR		Chest Discomfort	Wellbutrin	SS	Glaxosmithkline	ORAL
Other		Facial Pain				
200MG Twice		Gait Disturbance				
per day	2 YR	Oedema Peripheral				

Date:12/21/04ISR Number: 4533691-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0537485A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
Other		Abortion Induced Drug Exposure During Pregnancy	Bupropion	PS	Glaxosmithkline	

Date:12/21/04ISR Number: 4533694-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0537602A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hepatic Failure Renal Failure		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:12/21/04ISR Number: 4533696-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0537624A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Colitis		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice per day	3 MON	Somnolence Stool Analysis Abnormal					

Date:12/21/04ISR Number: 4533697-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0537625A

Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL
6 MON				No Concurrent Medication	C		

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Freedom Of Information (FOI) Report

Date:12/21/04ISR Number: 4533704-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0537916A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression Medication Error		Wellbutrin	PS	Glaxosmithkline	

Date:12/21/04ISR Number: 4533723-4Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0361120A  
Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Hypertensive Nephropathy		Zyban	PS	Glaxosmithkline	
UNKNOWN	150MG Twice per day	Nephropathy  Renal Amyloidosis Renal Failure					

Date:12/21/04ISR Number: 4533946-4Report Type:Expedited (15-DaCompany Report #2004S1003059  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety Arthralgia Change Of Bowel Habit	Health Professional	Bupropion Hydrochloride Tablets 75mg Mylan	PS	Mylan	ORAL
Required Intervention to 75 MG TID, Prevent Permanent ORAL Impairment/Damage		Feeling Abnormal  Gastric Disorder Haematochezia Nervousness Pharmaceutical Product Complaint Restlessness Treatment Noncompliance		Antibiotic Indomethacin	C C		

Date:12/21/04ISR Number: 4534057-4Report Type:Direct  
Age:51 YR Gender:Male I/FU:I

Company Report #CTU 234759

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash		Bupropion	PS		
				Gabapentin	C		
				Prazosin Hcl	C		
				Ketoconazole 2%			
				Shampoo	C		
				Ketoconazole 2%			
				Cream	C		
				Hydrocortisone	C		

Date:12/21/04ISR Number: 4534214-7Report Type:Direct Company Report #CTU 234851  
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Nausea		Generic Wellbutrin	PS		
1 QD		Pharmaceutical Product Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/21/04ISR Number: 4537347-4Report Type:Expedited (15-DaCompany Report #MK200412-0181-1  
Age:43 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Multiple Drug Overdose	Literature Health Professional	Restoril Capsules Bupropion	PS SS		

Date:12/21/04ISR Number: 4537782-4Report Type:Expedited (15-DaCompany Report #MK200411-0327-1  
Age:50 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Multiple Drug Overdose	Literature Health Professional	Propoxyphene Napsylate/Apap Bupropion Alprazolam	PS SS SS		

Date:12/22/04ISR Number: 4539754-2Report Type:Expedited (15-DaCompany Report #US-GDB-0411585  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged TOPICAL	1 APP QD TP	Anaphylactic Reaction Drug Hypersensitivity	Consumer	Differin (Adapalene) Cream 0.1% (Dpt)	PS		
600 MG QD PO		Gastrooesophageal Reflux Disease		Wellbutrin Wellbutrin Nuvaring	SS SS C		ORAL

Date:12/23/04ISR Number: 4536145-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0406391A  
Age: Gender:Male I/FU:F

Outcome	PT
Other	Acquired Porphyria Aggression Agitation Akathisia Amnesia

Arthralgia  
Asthenia  
Bipolar Disorder  
Body Temperature  
Fluctuation  
Coordination Abnormal  
Crying  
Depression  
Disability  
Dizziness  
Dysarthria  
Dysphemia  
Dyspnoea  
Electric Shock  
Emotional Disorder  
Encephalopathy  
Facial Palsy  
Fatigue  
Feeling Abnormal  
Headache  
Hyperaesthesia  
Hypersensitivity  
Hypoxia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
42	DAY	Immune System Disorder Injury Insomnia Intentional Self-Injury	Wellbutrin	PS	Glaxosmithkline	ORAL
		Mania	Tranquilizer	SS		
		Mastocytosis	Depakote	SS		
		Metabolic Encephalopathy	Zyprexa	SS		
		Muscle Spasms	Alcohol	C		
		Myalgia				
		Myoclonus				
		Nervous System Disorder				
		Nightmare				
		Overdose				
		Pain Of Skin				
		Petit Mal Epilepsy				
		Suicidal Ideation				
		Suicide Attempt				
		Throat Tightness				
		Tremor				
		Vestibular Disorder				
		Vomiting				

Date:12/23/04ISR Number: 4536189-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0538149A

Age:28 YR Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
Other		Drug Exposure During Pregnancy Premature Rupture Of Membranes Stillbirth	Bupropion	PS	Glaxosmithkline	

Date:12/23/04ISR Number: 4536190-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0538149B

Age: Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
Congenital Anomaly		Chorioamnionitis Drug Exposure During Pregnancy	Bupropion	PS	Glaxosmithkline	

Intra-Uterine Death  
Micrognathia  
Umbilical Cord  
Abnormality

Date:12/23/04ISR Number: 4536210-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0361887A

Age:58 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	1TAB Per day	Chorea		Zyban	PS	Glaxosmithkline	ORAL
UNKNOWN		Dystonia		Procyclidine	SS	Glaxosmithkline	
50MCG Per day		Facial Nerve Disorder		Thyroxine	C	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/23/04ISR Number: 4536228-XReport Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0045476A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged dosage text		Cardiac Disorder		Zyban	PS	Glaxosmithkline	ORAL
		Delusion					
		Dyspnoea					
		Hyperventilation					
		Panic Attack					
		Suicidal Ideation					

Date:12/23/04ISR Number: 4536846-9Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0537702A

Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Anxiety		Zyban	PS	Glaxosmithkline	ORAL
		Crying					
		Depression					
		Homicidal Ideation					
		Insomnia					
		Mood Swings					
		Sleep Disorder					
		Suicidal Ideation					

Date:12/23/04ISR Number: 4536856-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0538325A

Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 8 MON		Cerebrovascular Accident		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Difficulty In Walking		Unknown Blood			
		Drug Ineffective		Pressure Medication	C		
		Visual Disturbance		Diabetes Medication	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Blindness Transient		Zyban	PS	Glaxosmithkline	ORAL
1TAB Variable							
Other		Rash					
dose	11 DAY	Swelling		Diclofenac Sodium	C	Glaxosmithkline	
UNKNOWN	1UNIT	Three					
times per day							
UNKNOWN	1UNIT	Three		Metronidazole	C	Glaxosmithkline	
times per day							
UNKNOWN	2UNIT	Four		Co-Codamol	C		
times per day							
UNKNOWN	.5TAB	At		Co-Careldopa	C		
night							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Feeling Abnormal		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	5 DAY	Vitreous Floaters					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/23/04ISR Number: 4536907-4Report Type:Expedited (15-DaCompany Report #CH-GLAXOSMITHKLINE-B0362601A  
Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	ORAL
57 DAY		Excitability					
		Negative Thoughts					
		Suicidal Ideation					

Date:12/23/04ISR Number: 4537073-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0526859A  
Age:75 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Condition Aggravated		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG See		Depression		Wellbutrin Sr	SS	Glaxosmithkline	
dosage text		Medication Error					

Date:12/23/04ISR Number: 4537074-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0531761A  
Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
600MG Unknown		Convulsion					
		Drooling					
		Eye Rolling					
		Overdose					
		Tremor					

Date:12/23/04ISR Number: 4537078-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0535456A  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Convulsion Wellbutrin Xl PS Glaxosmithkline ORAL  
Medication Error

Date:12/23/04ISR Number: 4537079-2Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0536818A  
Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	200MG Per day	Premature Ejaculation		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:12/23/04ISR Number: 4537080-9Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0538431A  
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	52 DAY	Visual Acuity Reduced		Zyban	PS	Glaxosmithkline	ORAL
				Nefazodone	C		

Date:12/23/04ISR Number: 4537096-2Report Type:Expedited (15-DaCompany Report #DK-GLAXOSMITHKLINE-B0362606A  
Age:85 YR Gender:Female I/FU:F

Outcome	PT
Other	Abdominal Pain Upper Fatigue Feeling Abnormal Insomnia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Mania Tinnitus				
Dose	Duration		Report Source	Product	Role	Manufacturer
6	DAY			Zyban	PS	Glaxosmithkline

Date:12/23/04ISR Number: 4537158-XReport Type:Expedited (15-DaCompany Report #GXKR2004US00593  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose						
Hospitalization - Initial or Prolonged		Cardiac Failure Congestive		Metoprolol	PS	Novartis Sector: Pharma
75 mg, BID		Drug Interaction		Bupropion (Ngx)	SS	
150 mg, BID		Dyspnoea		Diltiazem	C	
480 mg/day		Fatigue Hypotension Metabolic Disorder Oedema Peripheral Pulmonary Hilum Mass Pulmonary Vascular Disorder Rales Sinus Bradycardia				

Date:12/23/04ISR Number: 4539354-4Report Type:Expedited (15-DaCompany Report #B0361967A  
Age:12 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose						
Disability		Difficulty In Walking Pruritus Serum Sickness Systemic Lupus	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS	
100 MG/TWICE		Erythematous Rash				
PER DAY				Methylphenidate Hcl	C	

Date:12/27/04ISR Number: 4539589-0Report Type:Direct  
Age:27 YR Gender:Male I/FU:I

Company Report #CTU 235183

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Generic Wellbutrin	PS		ORAL
1 PO BID							

Date:12/27/04ISR Number: 4542295-XReport Type:Expedited (15-DaCompany Report #US-GDP-0411585  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Pain Anaphylactic Reaction	Consumer	Differin (Adapalene) Cream 0.1% (Dpt)	PS		
1 APP QD TP							
150 MG QD PO		Blood Oestrogen Decreased		Wellbutrin	SS		ORAL
300 MG QD PO		Drug Hypersensitivity		Wellbutrin	SS		ORAL
		Dysthymic Disorder Emotional Distress Eye Irritation Gastrooesophageal Reflux Disease Nausea		Nuvaring	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/28/04ISR Number: 4540558-5Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 235221

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Bupropion 10 Mg	PS		
3DAY		Convulsion Drug Toxicity					

Date:12/28/04ISR Number: 4541299-0Report Type:Expedited (15-DaCompany Report #B0361960A  
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Drug Interaction Haemodynamic Instability Procedural Complication Procedural Hypertension	Literature Health Professional	Wellbutrin (Formulation Unknown) (Bupropion Hydrochloride) Linezolid (Formulation Unknown) (Linezolid) Propofol Suxamethonium Isoflurane Fentanyl Antibiotics	PS    SS C C C C C		

Date:12/29/04ISR Number: 4540891-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0406391A  
Age: Gender:Male I/FU:F

Outcome	PT
Other	Acquired Porphyria Aggression Agitation Akathisia Amnesia Arthralgia Asthenia Bipolar Disorder Body Temperature Fluctuation Coordination Abnormal

Crying  
Depression  
Disability  
Dizziness  
Dysarthria  
Dysphemia  
Dyspnoea  
Electric Shock  
Emotional Disorder  
Encephalopathy  
Facial Palsy  
Fatigue  
Feeling Abnormal  
Headache  
Hyperaesthesia  
Hypersensitivity  
Hypoxia  
Immune System Disorder  
Injury  
Insomnia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
42	DAY	Intentional Self-Injury Mania Mastocytosis Metabolic Encephalopathy		Wellbutrin	PS	Glaxosmithkline	ORAL
		Muscle Spasms Myalgia Myoclonus Nervous System Disorder Nightmare Overdose Pain Of Skin Petit Mal Epilepsy Schizoaffective Disorder Suicidal Ideation Suicide Attempt Throat Tightness Tremor Vestibular Disorder Vomiting		Tranquilizer Depakote Zyprexa Alcohol	SS SS SS C		

Date:12/29/04ISR Number: 4540901-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0538500A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day		Dizziness Fall Loss Of Consciousness Tinnitus Tremor		Wellbutrin Ditropan Paroxetine Lisinopril Atenolol Ranitidine Diuretic Folic Acid Tylenol Arthritis	PS C C C C C C C C	Glaxosmithkline Glaxosmithkline Glaxosmithkline	ORAL

Date:12/29/04ISR Number: 4540902-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0538552A

Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other 150MG Twice	Deafness	Wellbutrin	PS	Glaxosmithkline	ORAL
per day	20 DAY				
		Flovent	C	Glaxosmithkline	
		Serevent	C	Glaxosmithkline	
		Synthroid	C	Glaxosmithkline	

Date:12/29/04ISR Number: 4540933-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0362498A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2TAB Twice		Psychotic Disorder		Zyban	PS	Glaxosmithkline	ORAL
per day	4	WK					

Date:12/29/04ISR Number: 4543654-1Report Type:Direct Company Report #CTU 235322  
 Age:41 YR Gender:Male I/FU:I

Outcome  
 PT  
 Depressive Symptom  
 Drug Ineffective



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pharmaceutical Product  
Complaint

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
100 MG Q DAY			Bupropion	PS		ORAL
PO						

Date:12/30/04ISR Number: 4543388-3Report Type:Direct Company Report #CTU 235434  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage		Stevens-Johnson Syndrome		Welbutrin	PS		

Date:12/30/04ISR Number: 4544193-4Report Type:Expedited (15-DaCompany Report #GXKR2004US00593  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cardiac Failure Congestive Drug Interaction	Literature Health Professional	Metoprolol (Metoprolol Tartrate) Unknown	PS		
75 MG, BID		Hypotension Metabolic Disorder		Bupropion (Ngx) (Bupropion)	SS		
150 MG, BID		Pulmonary Hilum Mass Pulmonary Vascular Disorder Sinus Bradycardia		Diltiazem (Diltiazem)	C		

Date:12/30/04ISR Number: 4544284-8Report Type:Expedited (15-DaCompany Report #K200401961  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability		Chorea Dystonia	Foreign Health Professional Other	Kemadrin (Procyclidine) Tablet, 5mg Zyban (Bupropion Hydrochloride) Tablet, 1 Tab	PS  SS		ORAL
1 TAB, QD,  ORAL				Thyroxine Sodium (Levothyroxine Sodium)	C		

Date:01/03/05ISR Number: 4543808-4Report Type:Expedited (15-DaCompany Report #GXKR2004US00593  
Age:56 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Cardiac Failure Congestive Drug Interaction Fatigue Hypotension Metabolic Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pulmonary Hilum Mass Pulmonary Vascular Disorder					
75 MG, BID		Rales Sinus Bradycardia	Literature Health Professional	Metoprolol (Metoprolol Tartrate)	PS		
150 MG, BID				Bupropion (Ngx) Bupropion)	SS		
				Diltiazem (Diltiazem)	C		
				..	C		

Date:01/04/05ISR Number: 4544140-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0533311A  
Age:62 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Life-Threatening			Arrhythmia Cardiac Failure		Wellbutrin Xl Diovan	PS C	Glaxosmithkline	ORAL
1 YR			Congestive		Norvasc	C		
1 YR					Lipitor	C		
1 YR					Aldactone	C		
1 YR					Coreg	C	Glaxosmithkline	
1 YR					Lasix	C	Glaxosmithkline	
1 YR					Potassium	C		

Date:01/04/05ISR Number: 4544158-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0363099A  
Age:36 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Unknown		Wk	Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL

TRANSDERMAL 68MG See Drug Interaction Implanon SS  
 Pregnancy  
 dosage text Unintended Pregnancy

Date:01/04/05ISR Number: 4545763-XReport Type:Direct Company Report #USP 57021  
 Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Medication Error		Wellbutrin Xl 150 Mg	PS		
XL TAB							
				Wellbutrin Sr 150 Mg	SS		
SR TAB							

Date:01/05/05ISR Number: 4544989-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0539065A  
 Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2 MON	Paranoia					
		Suicide Attempt					

Date:01/05/05ISR Number: 4545007-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0350764A  
 Age:27 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cardiospasm		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	31 DAY						
Initial or Prolonged		Chest Pain		Actiskenan	SS	Glaxosmithkline	ORAL
20MG Per day	1 DAY						
				Efferalgan	C	Glaxosmithkline	ORAL
1 DAY							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/05/05ISR Number: 4545014-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0363078A  
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged dosage text	8 DAY	Arthralgia Joint Swelling Myalgia Pyrexia Rash Serum Sickness		Zyban	PS	Glaxosmithkline	ORAL

Date:01/05/05ISR Number: 4545017-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0363180A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Unknown Initial or Prolonged		Tetany		Zyban	PS	Glaxosmithkline	ORAL

Date:01/05/05ISR Number: 4545020-1Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0363390A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN Initial or Prolonged		Depressed Mood Dizziness Fall Feeling Abnormal Medication Error Overdose		Zyban	PS	Glaxosmithkline	

Date:01/05/05ISR Number: 4548241-7Report Type:Direct Company Report #CTU 235766  
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death	Agitation	Desipramine	PS
50 MG			
	Anger	Wellbutrin (Samples)	SS
	Fear	Lamictal	SS
100 MG	Overdose		

Date:01/06/05ISR Number: 4546485-1Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0506123A  
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG per day		Abortion Spontaneous		Bupropion	PS	Glaxosmithkline	
		Drug Exposure During Pregnancy					
		Foetal Heart Rate Abnormal					
		Ultrasound Antenatal Screen Abnormal					

Date:01/06/05ISR Number: 4546488-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0516229B  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged Congenital Anomaly		Drug Exposure During Pregnancy		Wellbutrin	PS	Glaxosmithkline	
		Pulmonary Hypertension					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/06/05ISR Number: 4546489-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0519165A  
 Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation Anxiety Bedridden Cholelithiasis Completed Suicide Crying Derealisation Diarrhoea Dizziness Feeling Abnormal Gastroenteritis Viral Gun Shot Wound Headache Insomnia Irritability Lethargy Nausea Nervousness Pyrexia Social Avoidant Behaviour Toothache Tremor Vomiting		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Date:01/06/05ISR Number: 4546494-2Report Type:Expedited (15-DaCompany Report #2004-03363  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged		Cardiac Disorder Dermatitis Exfoliative Discomfort Ill-Defined Disorder Injury Medication Error Oedema Peripheral Pain Rash Erythematous Rash Pustular Respiratory Disorder		Bupropion Lamisil Allopurinol	PS C C	Glaxosmithkline Glaxosmithkline Glaxosmithkline	ORAL

Skin Exfoliation  
Skin Lesion  
Skin Reaction  
Sleep Disorder  
Swollen Tongue

Date:01/06/05ISR Number: 4546495-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0533311A  
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Life-Threatening		Cardiac Failure		Diovan	C		
1 YR							
		Congestive		Norvasc	C		
1 YR							
				Lipitor	C		
1 YR							
				Aldactone	C		
1 YR							
				Coreg	C	Glaxosmithkline	
1 YR							
				Lasix	C	Glaxosmithkline	
1 YR							

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Freedom Of Information (FOI) Report

1 YR Potassium C

Date:01/06/05ISR Number: 4546506-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0539079A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion		Wellbutrin	PS	Glaxosmithkline	

Date:01/06/05ISR Number: 4546513-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0539461A  
 Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day Initial or Prolonged		Convulsion Fall Overdose Skin Laceration		Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:01/07/05ISR Number: 4547433-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0406391A  
 Age: Gender:Male I/FU:F

Outcome	PT
Other	Acquired Porphyria Aggression Agitation Akathisia Alcohol Use Amnesia Arthralgia Asthenia Bipolar Disorder Body Temperature Fluctuation Convulsion Coordination Abnormal Crying Depression Depression Suicidal

Disability  
Dizziness  
Dysarthria  
Dysphemia  
Dyspnoea  
Electric Shock  
Emotional Disorder  
Encephalopathy  
Facial Palsy  
Fatigue  
Feeling Abnormal  
Hair Metal Test Abnormal  
Headache  
Hyperaesthesia  
Hypersensitivity  
Hypoxia  
Immune System Disorder  
Injury  
Insomnia  
Intentional Self-Injury  
Mania

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
42	DAY	Mastocytosis Metabolic Encephalopathy Multiple Chemical Sensitivity	Wellbutrin	PS	Glaxosmithkline	ORAL
		Muscle Spasms	Tranquilizer	SS		
		Myalgia	Depakote	SS		
		Myoclonus	Zyprexa	SS		
		Nervous System Disorder	Alcohol	C		
		Neurotoxicity				
		Nightmare				
		Overdose				
		Pain Of Skin				
		Petit Mal Epilepsy				
		Porphyria				
		Schizoaffective Disorder				
		Suicidal Ideation				
		Suicide Attempt				
		Throat Tightness				
		Tremor				
		Vestibular Disorder				
		Vomiting				

Date:01/07/05ISR Number: 4547434-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0516229A

Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Drug Exposure During Pregnancy		Effexor	C		
		Placental Insufficiency					
		Pre-Eclampsia					

Date:01/07/05ISR Number: 4547447-0Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0539707A

Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
Other		Cyanosis					
		Dyspnoea					
		Tremor					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Antidepressant Drug Level Increased	Foreign Study Health	Trevilor (Venlafaxine Hydrochloride)	PS		ORAL
SEE IMAGE	45	DAY					
		Poisoning Supraventricular	Professional Other	Zyban (Amfebutamone Hydrochloride)	SS		ORAL
SEE IMAGE	15	DAY					
		Extrasystoles Tachycardia Ventricular Extrasystoles		Zopiclone (Zopiclone) Tibolone (Tibolone)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/10/05ISR Number: 4548478-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0525434A  
Age:76 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Per day	3	WK		Wellbutrin	PS	Glaxosmithkline	ORAL
			Amnesia				
			Balance Disorder	Prevacid	C		
			Dizziness	Estradiol	C		
			Dry Mouth	Synthroid	C	Glaxosmithkline	
			Dysphemia	Hctz	C		
			Feeling Abnormal				
			Hallucination				
			Tremor				
			Vision Blurred				

Date:01/10/05ISR Number: 4548479-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0532210A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Congenital Anomaly							
			Double Ureter	Bupropion	PS	Glaxosmithkline	
			Drug Exposure During Pregnancy				
			Renal Disorder				
			Vesicoureteric Reflux				

Date:01/10/05ISR Number: 4548482-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0537255A  
Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Per day		WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Vision Blurred				
			Visual Acuity Reduced	Cozaar	C		

Date:01/10/05ISR Number: 4548492-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0539841A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Blood Urine Present Wellbutrin PS Glaxosmithkline ORAL  
150MG Per day 5 DAY  
Vaginal Haemorrhage No Concurrent Medication C

Date:01/10/05ISR Number: 4548527-6Report Type:Expedited (15-DaCompany Report #2005000353  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypotension		Zyban	PS	Glaxosmithkline	ORAL
4500MG Single		Overdose					
dose		Suicide Attempt		Ethanol	SS		ORAL

Date:01/10/05ISR Number: 4553828-1Report Type:Direct Company Report #CTU 235994  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pharmaceutical Product		Bupropion	PS		ORAL
150 MGM BID		Complaint					
PO		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/10/05ISR Number: 4566200-5Report Type:Direct  
Age:68 YR Gender:Male I/FU:I

Company Report #CTU 236069

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Insomnia		Zyban (Generic) 150 Mg Bid	PS		
150 MG BID		Tachycardia Tremor					

Date:01/10/05ISR Number: 4567643-6Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 236060

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Affect Lability		Wellbutrin Sr 150 Mg	PS		
Hospitalization - THESE TWICE A Initial or Prolonged DAY		Drug Ineffective  Pharmaceutical Product Complaint					

Date:01/11/05ISR Number: 4549644-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0539903A  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539903A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Valtrex	PS	Glaxosmithkline	ORAL
Other 500MG Per day  300MG Unknown		Drug Interaction		Wellbutrin Xl	SS	Glaxosmithkline	ORAL

Date:01/11/05ISR Number: 4549645-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0540033A  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540033A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression		Wellbutrin	PS	Glaxosmithkline	ORAL
Other 100MG Twice							

per day 7 MON Drug Ineffective  
Irritability

Viagra

C

Date:01/11/05ISR Number: 4549646-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0540040A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health	Wellbutrin	PS	Glaxosmithkline	ORAL
Life-Threatening		Convulsion	Professional	Alcohol	SS		
Hospitalization - Initial or Prolonged Disability Other		Overdose					

Date:01/11/05ISR Number: 4549657-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0363375A  
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Periorbital Oedema		Zyban	PS	Glaxosmithkline	ORAL
150MG per day	8 DAY	Urticaria		Lansoprazole	C		ORAL
30MG per day							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/12/05ISR Number: 4550424-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0540137A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/12/05ISR Number: 4553858-XReport Type:Direct Company Report #CTU 236375  
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS		
UNKNOWN DOSE		Impulsive Behaviour					
ONCE DAILY				Sominex	C		

Date:01/12/05ISR Number: 4563722-8Report Type:Direct Company Report #CTU 236486  
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Paranoia		Bupropion	PS		ORAL
150 MG BID		Pharmaceutical Product					
ORAL		Complaint					
		Thinking Abnormal					

Date:01/13/05ISR Number: 4551752-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0533112A  
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Psoriasis		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	9 MON			Ssri	C		

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Brain Damage Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 21 DAY		Arthralgia		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged UNKNOWN		C-Reactive Protein Increased Coagulation Factor V Level Decreased Coagulation Factor Vii Level Decreased Fibrin D Dimer Increased Leukocytosis Lymphopenia Proteinuria Prothrombin Level Decreased Pyrexia Urticaria Generalised		Adepal	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/13/05ISR Number: 4553485-4Report Type:Expedited (15-DaCompany Report #D0039102A  
 Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	5 DAY	Disturbance In Attention Drug Abuser Hallucinations, Mixed	Foreign Literature Health	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL		Insomnia Psychotic Disorder Schizophreniform Disorder Suicidal Ideation Vision Blurred	Professional				

Date:01/13/05ISR Number: 4553883-9Report Type:Direct Company Report #CTU 236672  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG P.O.	2 YR	Aphthous Stomatitis		Wellbutrin Xl 150 Mg	PS		ORAL

Date:01/13/05ISR Number: 4553884-0Report Type:Direct Company Report #CTU 236669  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 BID P.O.		Aphthous Stomatitis		Wellbutrin Sr 150 Mg	PS		ORAL

Date:01/13/05ISR Number: 4554060-8Report Type:Direct Company Report #CTU 236614  
 Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Gun Shot Wound Insomnia Intentional Self-Injury Mania		Wellbutrin Lithium Lamictal Wellbutrin	PS C C C		

Date:01/14/05ISR Number: 4552332-4Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496788A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Imitrex	PS	Glaxosmithkline	
SUBCUTANEOUS	6MG Four	Fall					
times per day				Wellbutrin Sr	SS	Glaxosmithkline	
				Lotronex	SS	Glaxosmithkline	
				Fioricet	SS		
				Prozac	SS		
				Strattera	SS		

Date:01/14/05ISR Number: 4553150-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0320004A  
Age:33 YR Gender:Female I/FU:F

Outcome	PT
Other	Anxiety
	Aptyalism
	Asthenia
	Cytolytic Hepatitis
	Dermatitis Exfoliative
	Hypersensitivity
	Hypotension

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	22 DAY	Inflammation Insomnia Leukopenia	Zyban	PS	Glaxosmithkline	ORAL
3 DAY		Lymphadenopathy Nausea	Tiaprofenic Acid	SS		ORAL
3 DAY		Painful Respiration	Aspegic	SS		ORAL
		Pruritus Purpura Rash Maculo-Papular Rash Morbilliform Rash Scarlatiniform Thrombocytopenia Tremor				

Date:01/14/05ISR Number: 4553159-XReport Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0364019A

Age: Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose Other UNKNOWN		PT Hyperthyroidism	Zyban	PS	Glaxosmithkline	
		Personality Change Visual Acuity Reduced Visual Field Defect				

Date:01/14/05ISR Number: 4553163-1Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0364304A

Age: Gender:Male I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1TAB See Initial or Prolonged dosage text		PT Blood Glucose Increased Dizziness	Zyban	PS	Glaxosmithkline	ORAL
		Electrocardiogram Abnormal Feeling Abnormal Hypoaesthesia Visual Disturbance				



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/05ISR Number: 4567668-0Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 236837

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other BID PO		Drug Effect Decreased		Bupropion Er 15 Mg	PS		ORAL
		Pharmaceutical Product Complaint					

Date:01/15/05ISR Number: 4605804-8Report Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #A03200400600

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abnormal Behaviour - Chest Pain Confusional State	Consumer	Ambien - Zolpidem Tartrate - Tablet - Other	PS		ORAL
5 MS HS - ORAL		Depression					
10 MG HS - ORAL		Dry Mouth Headache		Ambien - Zolpidem Tartrate - Tablet	SS		ORAL
		Heart Rate Increased					
150 MG QD - ORAL		Libido Decreased Mood Swings Suicide Attempt		Wellbutrin - Bupropion Hydrochloride - Tablet - 150 Mg	SS		ORAL
				Paroxetine Hydrochloride	C		

Date:01/16/05ISR Number: 4606009-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #HQWYE882211FEB04

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Health Professional	Effexor (Venlafaxine Hydrochloride,			

ORAL				Unspec)	PS		ORAL
				Wellbutrin - Slow Release (Amfebutamone Hydrochloride)			
ORAL					SS		ORAL

Date:01/18/05ISR Number: 4554288-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0537119A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser Medication Error		Wellbutrin	PS	Glaxosmithkline	NASAL

Date:01/18/05ISR Number: 4554291-7Report Type:Expedited (15-DaCompany Report #2004108860  
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dementia Weight Increased		Zyban Nicorette Gum	PS SS	Glaxosmithkline Glaxosmithkline	ORAL
4	DAY			Nicorette	SS	Glaxosmithkline	
4	DAY			Nicorette Patch	SS	Glaxosmithkline	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/18/05ISR Number: 4554736-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0494832A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Paxil	PS	Glaxosmithkline	ORAL
Other		Anxiety		Zyban	SS	Glaxosmithkline	ORAL
		Convulsion					
		Drug Dependence					
		Drug Withdrawal Syndrome					
		Insomnia					
		Intentional Misuse					
		Laceration					
		Overdose					
		Suicide Attempt					

Date:01/18/05ISR Number: 4554742-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0519822A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Paxil	PS	Glaxosmithkline	ORAL
Hospitalization -		Depression					
30MG Per day							
Initial or Prolonged		Dizziness		Wellbutrin	SS	Glaxosmithkline	
		Drug Withdrawal Syndrome		Flexeril	SS		
		Feeling Abnormal		Skelaxin	SS		
		Intentional Misuse		Antiinflammatory	SS		
		Suicidal Ideation		Aspirin	SS	Glaxosmithkline	
		Suicide Attempt		Theophylline	SS		
				Risperdal	SS		
				Cold Medicine	SS		

Date:01/18/05ISR Number: 4554744-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0536055A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Life-Threatening		Dermatitis					
300MG In the							
Hospitalization -		Skin Exfoliation					
morning	18 DAY						
Initial or Prolonged		Stevens-Johnson Syndrome		Seroquel	C		
Disability							
Other							

Date:01/18/05ISR Number: 4554752-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0540816A  
Age:23 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 20MG Per day Initial or Prolonged	Agitation Anger Crying Drug Withdrawal Syndrome Hyperhidrosis Hyperventilation Hypoaesthesia Irritability Mood Swings Nausea Pain Suicide Attempt		Paxil Wellbutrin	PS SS	Glaxosmithkline Glaxosmithkline	ORAL



ORAL

Multivitamins

SS

ORAL

PRN; ORAL

Glucosamine

SS

ORAL

PRN; ORAL

Compazine

SS

ORAL

10 MG PRN;

ORAL

Date:01/18/05ISR Number: 4557138-8Report Type:Expedited (15-DaCompany Report #B0273496A

Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthralgia Coagulation Factor V Level Decreased Coagulation Factor Vii	Foreign Literature Health Professional	Bupropion Hydrochloride Tablet -Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL		Level Decreased Factor Ii Deficiency Inflammation Lymphocyte Count Decreased Neutrophil Count Increased Proteinuria Pyrexia Urticaria Generalised White Blood Cell Count Increased		Adepal	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/19/05ISR Number: 4556030-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0537277A  
Age:16 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
21 DAY			Grand Mal Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
			Loss Of Consciousness				

Date:01/19/05ISR Number: 4556032-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0540290A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
100MG Unknown			Hallucination	Wellbutrin	PS	Glaxosmithkline	ORAL
			Medication Error	No Concurrent			
			Panic Attack	Medication	C		

Date:01/19/05ISR Number: 4556044-2Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0346847A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
Hospitalization - Initial or Prolonged			Agitation	Zyban	PS	Glaxosmithkline	ORAL
			Alcohol Poisoning				
			Confusional State				
			Dehydration				
			Extrapyramidal Disorder				
			Overdose				
			Suicide Attempt				

Date:01/19/05ISR Number: 4556843-7Report Type:Direct Company Report #CTU 237176  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
100 MG BID			Feeling Abnormal	Wellbutrin Sr 100 Mg Bid	PS		

Date:01/19/05ISR Number: 4559125-2Report Type:Expedited (15-DaCompany Report #2004AL001522  
Age:27 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 20 MG; X1; PO	Arteriospasm Coronary	Foreign	Kadian	PS		ORAL
Initial or Prolonged Required PO	Self-Medication	Health Professional	Bupropion Hydrochloride	SS		ORAL
Intervention to Prevent Permanent Impairment/Damage		Other	Efferalgan	C		

Date:01/20/05ISR Number: 4557056-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0406391A  
Age: Gender:Male I/FU:F

Outcome	PT
Other	Acquired Porphyria Aggression Agitation Akathisia Alcohol Use Amnesia Arthralgia Asthenia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
42	DAY	Bipolar Disorder Body Temperature Fluctuation	Wellbutrin	PS	Glaxosmithkline	ORAL
		Convulsion	Tranquilizer	SS		
		Coordination Abnormal	Depakote	SS		
		Crying	Zyprexa	SS		
		Depression	Alcohol	C		
		Depression Suicidal				
		Disability				
		Dizziness				
		Dysarthria				
		Dysphemia				
		Dyspnoea				
		Electric Shock				
		Emotional Disorder				
		Encephalopathy				
		Facial Palsy				
		Fatigue				
		Feeling Abnormal				
		Hair Metal Test Abnormal				
		Head Injury				
		Headache				
		Hyperaesthesia				
		Hypersensitivity				
		Hypoxia				
		Immune System Disorder				
		Injury				
		Insomnia				
		Intentional Self-Injury				
		Mania				
		Mastocytosis				
		Medication Error				
		Metabolic Encephalopathy				
		Muscle Spasms				
		Myalgia				
		Myoclonus				
		Nervous System Disorder				
		Nightmare				
		Overdose				
		Pain Of Skin				
		Petit Mal Epilepsy				
		Psychotic Disorder				
		Schizoaffective Disorder				
		Suicidal Ideation				
		Suicide Attempt				

Throat Tightness  
Tremor  
Vestibular Disorder  
Vomiting

Date:01/20/05ISR Number: 4557059-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0535570A

Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Excoriation		No Concurrent			
		Grand Mal Convulsion		Medication	C		
		Loss Of Consciousness					
		Syncope					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/20/05ISR Number: 4557060-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0538024A  
 Age:23 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Per day Initial or Prolonged 30 DAY	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
	Incontinence		Symbyax	SS		
	Loss Of Consciousness Sexual Dysfunction					

Date:01/20/05ISR Number: 4557064-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0540443A  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300MG See Initial or Prolonged dosage text	Blood Pressure Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
	Coma					
	Drug Ineffective		Lexapro	SS		
	Drug Interaction		Atenolol	C		
	Medication Error		Avandia	C	Glaxosmithkline	
			Nexium	C		
			Aspirin	C	Glaxosmithkline	
			Aricept	C		
			Neurontin	C		
			Centrum Silver	C		
			Magnesium	C		
			Potassium	C		
			Duragesic	C		
			B12	C	Glaxosmithkline	
			Hydrocodone	C		
			Zocor	C		
			Diazepam	C		
			Isosorbide	C		

Date:01/20/05ISR Number: 4557065-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0540945A  
 Age:44 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Hospitalization - 300MG Per day Initial or Prolonged	Convulsion Fall Joint Injury	Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	ORAL
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Date:01/20/05ISR Number: 4557066-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0540971A  
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 450MG In the morning		Grand Mal Convulsion Memory Impairment Speech Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:01/20/05ISR Number: 4557077-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0364668A  
Age: Gender:Male I/FU:I

Outcome Other	PT Adverse Event Aggression Anxiety Coordination Abnormal Feeling Of Despair Irritability
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Panic Attack				
		Sensory Disturbance				
		Suicidal Ideation	Report Source	Product	Role	Manufacturer
Dose	Duration					Route
6	WK	Thinking Abnormal		Zyban	PS	Glaxosmithkline
						ORAL

Date:01/20/05ISR Number: 4557078-4Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0364698A  
 Age: Gender:Female I/FU:F

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose							
Hospitalization -		Cerebral Artery Embolism		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Cerebrovascular Accident					
		Coma					
		Convulsion					
		Feeling Abnormal					
		Myocardial Infarction					

Date:01/20/05ISR Number: 4557081-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0364815A  
 Age:49 YR Gender:Female I/FU:F

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose							
Other		Renal Colic		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							
				Celecoxib	C		ORAL
200MG Per day				Paracetamol	C	Glaxosmithkline	
UNKNOWN				Co-Proxamol	C		
UNKNOWN				Lisinopril	C		
UNKNOWN							

Date:01/20/05ISR Number: 4557086-3Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0364857A  
 Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Valve Disease		Zyntabac	PS	Glaxosmithkline	
		Hemiparesis		Nicotinell	C	Glaxosmithkline	
		Pulmonary Oedema					

Date:01/20/05ISR Number: 4557087-5Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0364991A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Zyban	PS	Glaxosmithkline	
150MG Twice							
per day							

Date:01/20/05ISR Number: 4560094-XReport Type:Direct Company Report #CTU 237485  
 Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Sr	PS		
150 MG ONE QD				Neurontin	C		

Date:01/20/05ISR Number: 4560160-9Report Type:Direct Company Report #CTU 237286  
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT
		Depressive Symptom
		Pharmaceutical Product

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Complaint

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
			Wellbutrin Sr	PS		

Date:01/20/05ISR Number: 4560207-XReport Type:Direct Company Report #CTU 237476  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropion Ssr 150 Mg	PS		
Other		Depression					
150 MG		Disease Recurrence Drug Ineffective Pharmaceutical Product Complaint					

Date:01/21/05ISR Number: 4558783-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541566A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Life-Threatening Hospitalization - Initial or Prolonged		Convulsion Overdose					

Date:01/21/05ISR Number: 4558789-7Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0348497A  
 Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
Other		Drug Interaction					
1TAB Twice per day	1 MON	Weight Increased		Reductil	SS		ORAL
15MG per day	2 MON						

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness Feeling Abnormal Hallucination, Tactile Nervousness Palpitations		Zyban	PS	Glaxosmithkline	ORAL

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -		Grand Mal Convulsion		Tylenol	SS	Glaxosmithkline	
Initial or Prolonged		Intentional Misuse		Zoloft	C		
Other		Refusal Of Treatment By Patient Suicide Attempt					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/24/05ISR Number: 4559531-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0540508A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Per day				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	MON			Wellbutrin	SS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:01/24/05ISR Number: 4559532-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0540547A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
4 WK				Wellbutrin	PS	Glaxosmithkline	ORAL
				Synthroid	C	Glaxosmithkline	
				Loss Of Consciousness			

Date:01/24/05ISR Number: 4559555-9Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0045685A

Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
				Zyban	PS	Glaxosmithkline	ORAL

Date:01/24/05ISR Number: 4559761-3Report Type:Expedited (15-DaCompany Report #US-MERCK-0501USA02727

Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
				Zocor	PS	Merck & Co., Inc	ORAL
				Intron A	SS		
INTRAVENOUS							
							DRIP
37 DAY							
				Wellbutrin	SS		ORAL
				Allopurinol	SS		ORAL

Lethargy  
Nausea

Clonazepam	SS	ORAL
Trazodone		
Hydrochloride	SS	ORAL
Remeron	SS	ORAL
Vitamins		
(Unspecified)	SS	ORAL
Glucosamine	SS	ORAL
Compazine	SS	ORAL

Date:01/24/05ISR Number: 4560653-4Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 237702

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypertension Pharmaceutical Product Complaint		Bupropion 75 Mg Two Tabs Po Bid	PS		

Date:01/24/05ISR Number: 4560654-6Report Type:Direct  
Age:39 YR Gender:Male I/FU:I

Company Report #CTU 237703

Outcome	PT
	Dizziness Drug Ineffective Nausea Pharmaceutical Product



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Complaint

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
ONE TAB TWICE DAILY			Bupropion Sr 150 Mg	PS		

Date:01/24/05ISR Number: 4560675-3Report Type:Direct Company Report #CTU 237720  
 Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dyspepsia				Bupropion Hydrochloride Extended Release Tabs	PS	Eon Labs	ORAL
TWICE DAILY	1 WK			Seroquel	C		
				Adalat Cc	C		
				Aspirin	C		
				Doxazosin	C		
				Hctz	C		
				Zantac	C		

Date:01/24/05ISR Number: 4560775-8Report Type:Direct Company Report #CTU 237723  
 Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dyspepsia				Bupropion Hydrochloride Extended Release Tabs 150 Mg Tab	PS	Eon Labs	ORAL
TWICE DAILY	1 WK			Eon Labs			
				Lithium	C		
				Seroquel	C		

Date:01/24/05ISR Number: 4560777-1Report Type:Direct  
Age:41 YR Gender:Male I/FU:I

Company Report #CTU 237722

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dyspepsia		Bupropion Hydrochloride Extended Release Tabs 150mg Tab Eon Labs	PS	Eon Labs	ORAL
TWICE DAILY	1	WK		Topiramate	C		

Date:01/24/05ISR Number: 4560778-3Report Type:Direct  
Age:26 YR Gender:Male I/FU:I

Company Report #CTU 237721

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dyspepsia		Bupropion Hydrochloride Extended Release Tabs 150 Mg Tab Eon Labs	PS	Eon Labs	ORAL
TWICE DAILY	1	WK		Ibuprofen	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/24/05ISR Number: 4563691-0Report Type:Expedited (15-DaCompany Report #MK200501-0123-1

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Bronchiolitis Drug Interaction	Foreign Health Professional Other	Anafranil Wellbutrin	PS SS		

Date:01/25/05ISR Number: 4560533-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541125A

Age:83 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100MG Twice per day	2 WK	Hallucination Medication Error Paranoia Pharmaceutical Product Complaint		Wellbutrin Lithium Ativan	PS C C	Glaxosmithkline Glaxosmithkline	ORAL

Date:01/25/05ISR Number: 4560534-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541132A

Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - Initial or Prolonged		Haemorrhage Overdose		Wellbutrin Unknown	PS C	Glaxosmithkline	ORAL

Date:01/25/05ISR Number: 4560550-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0365184A

Age:37 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG per day	1 DAY	Dyspnoea Migraine		Zyban Glucosamine	PS C	Glaxosmithkline	ORAL
UNKNOWN							

Oropharyngeal Swelling

Co Q10 Enzyme

C

ORAL

Date:01/25/05ISR Number: 4560552-8Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0365187A

Age:21 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthma		Zyban	PS	Glaxosmithkline	ORAL
12 DAY		Bronchospasm		Salbutamol	C	Glaxosmithkline	
UNKNOWN							

Date:01/26/05ISR Number: 4561595-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0527550A

Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG At		Oedema Peripheral					
night	2 DAY	Palmar Erythema					
		Pruritus					
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/26/05ISR Number: 4561604-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541103A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day 4 MON	Aggression		Wellbutrin	PS	Glaxosmithkline	ORAL
		Anger		Synthroid	C	Glaxosmithkline	
		Antisocial Behaviour		Premarin	C		
		Irritability		Klonopin	C		
		Negative Thoughts					

Date:01/26/05ISR Number: 4561605-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541137A

Age:22 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Pancreatitis		Wellbutrin	PS	Glaxosmithkline	ORAL
				Lexapro	C		

Date:01/26/05ISR Number: 4561606-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541138A

Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day 3 WK	Joint Stiffness		Wellbutrin	PS	Glaxosmithkline	ORAL
		Musculoskeletal Stiffness		Lipitor	C		
		Oedema Peripheral		Aspirin	C	Glaxosmithkline	
		Rash					
		Serum Sickness					

Date:01/26/05ISR Number: 4561607-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541141A

Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day 3 WK	Erythema		Wellbutrin	PS	Glaxosmithkline	ORAL
		Musculoskeletal Stiffness		Micronor	C		
		Oedema Peripheral					

Orbital Oedema  
Rash  
Serum Sickness  
Urticaria

Date:01/26/05ISR Number: 4561616-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541965A  
Age:16 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 4 Initial or Prolonged	YR	Epistaxis Erythema Intentional Misuse Overdose		Wellbutrin Xl Vitamin B Lamictal Seroquel Prozac Strattera Marijuana	PS SS C C C C C	Glaxosmithkline Glaxosmithkline	ORAL

Date:01/26/05ISR Number: 4561643-8Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0365434A  
Age:45 YR Gender:Female I/FU:F

Outcome	PT
Other	Dysarthria Hypoaesthesia Oral

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Paraesthesia  
Tongue Paralysis

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1 DAY		Zyban	PS	Glaxosmithkline	ORAL
100MG per day			Albyl Minor	C	Glaxosmithkline	
100MG per day			Seloken Zoc	C		
			Nobligan Retard	C		
			Celebra	C		

Date:01/26/05ISR Number: 4563027-5Report Type:Expedited (15-DaCompany Report #GXBR2005US00444  
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Asthenia Confusional State Decreased Appetite	Health Professional	Allopurinol Tablets Usp (Ngx) (Allopurinol) Tablet	PS		ORAL
100 MG, QHS, ORAL		Dehydration		Bupropion Hcl Tablets, (Ngx) (Bupropion) Tablet	SS		ORAL
300 MG, BID, ORAL		Drug Interaction Lethargy Nausea		Clonazepam (Ngx) (Clonazepam) Tablet	SS		ORAL
0.5 MG, QHS, ORAL				Mirtazapine Tablets (Ngx) (Mirtazapine) Tablet	SS		ORAL
45 MG, QHS, ORAL				Prochlorperazine Maleate Tablets Usp (Ngx) (Prochlorperazine)			

10 MG, PRN,	Tablets	SS		ORAL
ORAL				
	Trazodone Hcl Tablets Usp (Ngx) (Trazodone) Tablet	SS		ORAL
50 MG, QHS,				
ORAL				
	Intron A "Schering-Plough" (Interferon Alfa 2-B) Injection	SS	Schering-Plough	ORAL
44 MIU, QW5,				
ORAL				
	Zocor "Merck" (Simvastatin)	SS	Merck	ORAL
10 MG, QD,				
ORAL				
	Multivitamins (Ascorbic Acid, Ergocalciferol, Folic Acid, Panthenol...	SS		ORAL
PRN, ORAL				
	Glucosamine (Glucosamine)	SS		ORAL
ORAL				



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/26/05ISR Number: 4566337-0Report Type:Direct  
Age:53 YR Gender:Male I/FU:I

Company Report #CTU 238109

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Myalgia		Bupropion	PS		

Date:01/26/05ISR Number: 4566367-9Report Type:Direct  
Age:45 YR Gender:Male I/FU:I

Company Report #CTU 238104

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation Irritability		Bupropion	PS		
				Gabapentin	C		
				Divalproex Ec (Delayed Release)	C		
				Fluoxetine Hcl	C		
				Hydromorphone Hcl	C		
				Morphine	C		
				Ranitidine Hcl	C		
				Bupropion Hcl	C		
				Vitamin B Complex/ Vitamin C Cap	C		
				Prazosin Hcl	C		
				Capsaicin	C		

Date:01/27/05ISR Number: 4570261-7Report Type:Direct  
Age:20 YR Gender:Female I/FU:I

Company Report #CTU 238334

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 150 DAILY		Generalised Oedema Hypotension		Wellbutrin Sr 150 Mg	PS		ORAL
ORAL		Rash Wheezing					

Date:01/27/05ISR Number: 4576966-6Report Type:Direct  
Age:54 YR Gender:Female I/FU:I

Company Report #CTU 238422

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	100 MG	1	Asthenia		Welbutrin Sr	PS		
			Cerebrovascular Accident					
DAILY			Confusional State		Welbutrin Sr	SS		
	150 MG	1	Dysgeusia					
DAILY			Feeling Abnormal					
			Partial Seizures					
			Sensory Disturbance					
			Visual Disturbance					

Date:01/28/05ISR Number: 4563282-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0042892A  
Age: Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	75MG	Twice	Chills		Wellbutrin	PS	Glaxosmithkline	ORAL
			Drug Ineffective					
	per day	4 DAY	Feeling Cold		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
	100MG	Unknown						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563283-3Report Type:Periodic  
Age:17 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0126084A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	2 WK	Arthralgia Joint Swelling Oedema Peripheral Pain In Extremity Rash Pruritic Serum Sickness Urticaria Generalised		Wellbutrin  Prozac	PS  C	Glaxosmithkline	ORAL  ORAL

Date:01/28/05ISR Number: 4563286-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0431982A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Chest Pain Increased Appetite Nicotine Dependence		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563287-0Report Type:Periodic  
Age:16 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0432445A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Three times per day		Drug Abuser Hallucination, Visual Intentional Misuse		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563288-2Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441460A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Weight Decreased		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563289-4Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441484A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Suicidal Ideation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563290-0Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441661A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dysgeusia Parosmia		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563291-2Report Type:Periodic  
Age:43 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441767A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Haematuria Hepatic Enzyme Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563292-4Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0441795A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		Insomnia		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563293-6Report Type:Periodic  
 Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0442009A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563294-8Report Type:Periodic  
 Age:66 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0442011A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice		Diplopia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	2 MON	Dizziness					
300MG Per day		Feeling Jittery		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
20MG Twice				Vasotec	C		ORAL
per day				Hctz	C		ORAL
25MG Per day				Tenormin	C		ORAL
50MG Twice							
per day							

Date:01/28/05ISR Number: 4563295-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0442046A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sexual Dysfunction		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563296-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442414A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Anger Irritability		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563297-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442538A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	

UNKNOWN

Date:01/28/05ISR Number: 4563298-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0442616A  
Age: YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alcohol Poisoning		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563299-7Report Type:Periodic  
 Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0442977A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Psychotic Disorder		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563300-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443006A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL
1	DAY						

Date:01/28/05ISR Number: 4563301-2Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443033A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563302-4Report Type:Periodic  
 Age:42 YR Gender:Female I/FU:I

Company Report #S03-USA-05003-01

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	11 DAY	Contusion		Wellbutrin	PS	Glaxosmithkline	
150MG Per day							
Initial or Prolonged		Convulsion		Celexa	SS		ORAL
20MG Per day							
Other		Disorientation					
		Fall					
		Parosmia					
		Tooth Fracture					
		Weight Increased					

Date:01/28/05ISR Number: 4563303-6Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0443270A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice		Seasonal Allergy		Wellbutrin	PS	Glaxosmithkline	ORAL
per day							

Date:01/28/05ISR Number: 4563304-8Report Type:Periodic Company Report #S03-USA-04967-01  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Per day		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
10MG Per day				Lexapro	SS		ORAL

Date:01/28/05ISR Number: 4563305-XReport Type:Periodic Company Report #S03-USA-04869-01  
 Age:53 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
6 MON		Alopecia Effluvium		Wellbutrin	PS	Glaxosmithkline	ORAL
10MG Per day	1 YR			Lexapro	SS		ORAL
				Risperdal	C		
				Klonopin	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563306-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443498A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Meniere'S Disease		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563308-5Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443915A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anhedonia		Wellbutrin	PS	Glaxosmithkline	ORAL
5 WK		Depression		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Hypersomnia		Lexapro	C		
		Oedema		Hydrochlorothiazide	C		

Date:01/28/05ISR Number: 4563309-7Report Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443922A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - 150MG Twice Initial or Prolonged per day	1 MON	Hallucination					
		Heart Rate Increased		Lexapro	C		
		Tremor					

Date:01/28/05ISR Number: 4563310-3Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0443930A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563311-5Report Type:Periodic  
Age:16 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0443936A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dystonia Swollen Tongue		Wellbutrin Geodon Lexapro	PS C C	Glaxosmithkline	ORAL

UNKNOWN

Date:01/28/05ISR Number: 4563312-7Report Type:Periodic  
Age:13 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0444037A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75MG Per day	6 DAY	Back Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
75MG Per day	6 DAY	Diarrhoea		Bupropion	SS	Glaxosmithkline	ORAL
		Eye Pain Hypomania Increased Appetite Nightmare Pain In Jaw Pharyngolaryngeal Pain Pyrexia Swelling Face					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563313-9Report Type:Periodic  
Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0444111A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation		Wellbutrin Prozac	PS C	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563314-0Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0490789A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression		Bupropion	PS	Glaxosmithkline	ORAL
3 DAY		Drug Ineffective		Zyban	SS	Glaxosmithkline	ORAL
150MG Twice per day	5 DAY			Monopril Paxil	C C	Glaxosmithkline	

Date:01/28/05ISR Number: 4563315-2Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491309A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypertension		Wellbutrin	PS	Glaxosmithkline	
1 YR							

Date:01/28/05ISR Number: 4563316-4Report Type:Periodic  
Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491318A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hyponatraemia		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563317-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0491566A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563318-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0491714A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day							

Date:01/28/05ISR Number: 4563319-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0491779A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus Generalised		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Unknown							

Date:01/28/05ISR Number: 4563320-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0491973A  
Age: Gender:Female I/FU:I

Outcome	PT
	Agitation
	Anorexia
	Asthenia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
4	WK	Drug Ineffective Insomnia Paranoia Tremor Weight Decreased	Bupropion	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563321-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0492259A  
Age:6 YR Gender: I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day		PT Grand Mal Convulsion	Bupropion	PS	Glaxosmithkline	ORAL
			Adderall Zyprexa	C C		

Date:01/28/05ISR Number: 4563322-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0492908A  
Age: YR Gender: I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT Renal Disorder	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563323-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0493260A  
Age: YR Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose UNKNOWN		PT Hyperhidrosis	Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563324-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0493314A  
Age:24 YR Gender:Male I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Pain Of Skin					
Date:01/28/05ISR Number: 4563325-5Report Type:Periodic			Company Report #US-GLAXOSMITHKLINE-A0493962A				
Age: YR	Gender:Female	I/FU:F					
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Hypersensitivity		Wellbutrin	PS	Glaxosmithkline	ORAL
Date:01/28/05ISR Number: 4563326-7Report Type:Periodic			Company Report #US-GLAXOSMITHKLINE-A0494331A				
Age:	Gender:Female	I/FU:I					
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
Date:01/28/05ISR Number: 4563327-9Report Type:Periodic			Company Report #US-GLAXOSMITHKLINE-A0494686A				
Age:36 YR	Gender:Female	I/FU:I					
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day	3 WK			Zomig	C		
				Fioricet	C		
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563328-0Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495030A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amenorrhoea		Wellbutrin	PS	Glaxosmithkline	
				Oxycontin	C		
				Lexapro	C		

Date:01/28/05ISR Number: 4563329-2Report Type:Periodic  
 Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0495040A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nightmare		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563330-9Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495043A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Irritability		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563331-0Report Type:Periodic  
 Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495250A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	
Other							

Date:01/28/05ISR Number: 4563332-2Report Type:Periodic  
 Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495288A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Bupropion	PS	Glaxosmithkline	
100MG Single		Drug Exposure During					
dose							

Pregnancy  
Feeling Abnormal

Date:01/28/05ISR Number: 4563333-4Report Type:Periodic Company Report #S03-USA-05410-01  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction		Wellbutrin	PS	Glaxosmithkline	ORAL
		Feeling Abnormal		Lexapro	SS		
UNKNOWN		Impaired Driving Ability					

Date:01/28/05ISR Number: 4563334-6Report Type:Periodic Company Report #S03-USA-05399-01  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Prolactin Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
				Celexa	SS		
UNKNOWN							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563335-8Report Type:Periodic  
Age: YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0495338A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563336-XReport Type:Periodic  
Age:80 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495339A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Depression Weight Decreased		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
225MG Unknown	5 MON			Lorazepam Unknown Sleeping Pill	C C		

Date:01/28/05ISR Number: 4563337-1Report Type:Periodic  
Age:8 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495378A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Stomach Discomfort		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Concerta	C		

Date:01/28/05ISR Number: 4563338-3Report Type:Periodic  
Age:91 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0495948A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin Hydrochlorothiazide	PS SS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563339-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496005A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin Celexa	PS SS	Glaxosmithkline	ORAL

UNKNOWN

Date:01/28/05ISR Number: 4563340-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0496059A  
 Age: YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperacusis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563341-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0496499A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Depression Suicidal Ideation		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563342-5Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496666A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Screen Positive		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563343-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496739A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Libido Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
				Celexa	SS		

UNKNOWN

Date:01/28/05ISR Number: 4563344-9Report Type:Periodic  
 Age:25 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496748A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Screen Positive		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563345-0Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0496772A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Cognitive Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563346-2Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0496776A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
				Luvox	C		

Date:01/28/05ISR Number: 4563347-4Report Type:Periodic Company Report #USA040156776  
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gastric Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563348-6Report Type:Periodic Company Report #03US08760  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gastrooesophageal Reflux Disease		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563349-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0496955A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Premenstrual Syndrome		Wellbutrin	PS	Glaxosmithkline	ORAL

10 MON



Somnolence

Wellbutrin

PS

Glaxosmithkline

ORAL

Date:01/28/05ISR Number: 4563355-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498028A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Libido Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563356-5Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498337A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							

Date:01/28/05ISR Number: 4563357-7Report Type:Periodic  
Age:68 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0498745A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Night Sweats		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
75MG Twice							
per day	18 MON			Avandia Zocor	C C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Naproxen C  
 Prevacid C

Date:01/28/05ISR Number: 4563358-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498763A  
 Age: YR Gender: I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Medication Error		Wellbutrin	PS	Glaxosmithkline	NASAL

Date:01/28/05ISR Number: 4563359-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0498774A  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
	Psychomotor Hyperactivity		Ssri	C		

Date:01/28/05ISR Number: 4563360-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499052A  
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Halitosis		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG	Unknown					

Date:01/28/05ISR Number: 4563361-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499219A  
 Age:48 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
2	WK					

Date:01/28/05ISR Number: 4563362-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499609A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG	Per day			Lithium	C	Glaxosmithkline	

Date:01/28/05ISR Number: 4563363-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0499639A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Parosmia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563364-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0500572A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG	Twice	Tremor		Wellbutrin	PS	Glaxosmithkline	
	per day						



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563365-6Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501021A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563366-8Report Type:Periodic  
 Age: YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501036A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563367-XReport Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0501097A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Unknown		Nightmare		Wellbutrin	PS	Glaxosmithkline	ORAL
				Seroquel	C		

Date:01/28/05ISR Number: 4563368-1Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501437A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Skin Discolouration		Wellbutrin	PS	Glaxosmithkline	ORAL
				Lexapro	C		

Date:01/28/05ISR Number: 4563369-3Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0501604A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice		Abdominal Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	12 YR	Drug Interaction					
UNKNOWN		Ear Congestion		Percocet	SS		
UNKNOWN		Insomnia		Decadron	SS		
UNKNOWN		Nausea		Kytril	SS	Glaxosmithkline	
UNKNOWN		Pharyngolaryngeal Pain		Compazine	SS	Glaxosmithkline	
		Tinnitus		Ativan	SS		
		Tremor		Benadryl	SS	Glaxosmithkline	
		Vomiting					
		White Blood Cell Count Decreased					

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
7 DAY		Rash Pruritic		Wellbutrin	PS	Glaxosmithkline	ORAL
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563372-3Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502147A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN							

Date:01/28/05ISR Number: 4563373-5Report Type:Periodic  
 Age:14 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502152A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
2	WK			Neurontin	C		

Date:01/28/05ISR Number: 4563374-7Report Type:Periodic  
 Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502178A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Thrombocytopenia		Wellbutrin	PS	Glaxosmithkline	
Other							

Date:01/28/05ISR Number: 4563375-9Report Type:Periodic  
 Age:80 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502186A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Feeling Abnormal		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563376-0Report Type:Periodic  
 Age:38 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502990A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 2 MON							

Lethargy  
Myalgia  
Nausea  
Pyrexia

Date:01/28/05ISR Number: 4563377-2Report Type:Periodic  
Age: YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0502991A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Erythema Multiforme		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563378-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503190A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 DAY		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
				Axid	C		
				Lotensin	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563379-6Report Type:Periodic  
 Age: YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503425A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563380-2Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503446A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563381-4Report Type:Periodic  
 Age:64 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503459A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Three		Euphoric Mood					
times per day	6	Nausea		Xanax	C		
				Ativan	C		

Date:01/28/05ISR Number: 4563382-6Report Type:Periodic  
 Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503531A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
75MG Twice							
per day	8	DAY		Klonopin	C		
				Prometrium	C		
				Synthroid	C	Glaxosmithkline	
				Voltaren	C	Glaxosmithkline	
				Allegra	C		
				Prevacid	C		

Date:01/28/05ISR Number: 4563383-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0503806A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
100MG Unknown	2 WK			Prozac	SS		
UNKNOWN							

Date:01/28/05ISR Number: 4563384-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504065A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
75MG Twice							
per day	3 MON						

Date:01/28/05ISR Number: 4563385-1Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504097A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Twitching		Wellbutrin	PS	Glaxosmithkline	ORAL
1 YR				Diuretic	C		
				Hormone Replacement	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563386-3Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504215A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Burning Sensation Dermatitis Exfoliative Erythema		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563387-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504275A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:01/28/05ISR Number: 4563388-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504323A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Three							
times per day							
2 WK				Wellbutrin Xl	SS	Glaxosmithkline	ORAL
				Zocor	C		
				Klonopin	C		
				Risperdal	C		
				Ditropan Xl	C		
				Ultram	C		
				Trazodone	C		

Date:01/28/05ISR Number: 4563389-9Report Type:Periodic  
 Age:18 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504329A

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Alanine Aminotransferase Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563390-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504332A  
Age: YR Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Mental Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563391-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504592A  
Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Anxiety Drug Ineffective		Wellbutrin Prozac	PS C	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563392-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0504656A  
Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Depression Suicidal Ideation		Wellbutrin	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563393-0Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504864A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	4 DAY	Rash		Adderall	C		

Date:01/28/05ISR Number: 4563394-2Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504867A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression		Wellbutrin	PS	Glaxosmithkline	ORAL
5 YR		Hostility		Ambien	C		
		Insomnia		Xanax	C		

Date:01/28/05ISR Number: 4563395-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504868A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -				No Concurrent Medication	C		
400MG Unknown	3 MON						
Initial or Prolonged							

Date:01/28/05ISR Number: 4563396-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505132A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Three							
times per day	4 WK						

Date:01/28/05ISR Number: 4563397-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505235A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthma Cough Dysgeusia Dyspepsia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563398-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505626A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563399-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505629A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	
9 DAY		Sleep Disorder Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563400-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0505934A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 WK		Vision Blurred		Wellbutrin	PS	Glaxosmithkline	ORAL
				Nortriptyline	C		

Date:01/28/05ISR Number: 4563401-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506447A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
5 DAY		Rash Generalised		Wellbutrin	PS	Glaxosmithkline	
5 MON				Lexapro	C		

Date:01/28/05ISR Number: 4563402-9Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0506507A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
				Alcohol	SS		

Date:01/28/05ISR Number: 4563403-0Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0506510A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hepatic Enzyme Increased Tachycardia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563404-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #2004S1000747

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Flatulence		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563405-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506580A  
Age: Gender: I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
		Malabsorption		Lamictal	SS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563406-6Report Type:Periodic Company Report #2004S1000814  
Age:53 YR Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Anorexia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Dry Mouth					

Date:01/28/05ISR Number: 4563407-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506917A  
Age:41 YR Gender:Female I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	6 MON			Trazodone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563408-XReport Type:Periodic  
Age:25 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506924A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 75MG Twice per day	2 WK	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
				Alcohol	C		
				Marijuana	C		

Date:01/28/05ISR Number: 4563409-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507091A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 150MG Per day		Adverse Event		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563410-8Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507155A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 150MG Three times per day		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
				Celexa	C		
				Trazodone	C		

Date:01/28/05ISR Number: 4563411-XReport Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507281A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 50MG Twice per day	4 MON	Headache		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563412-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507422A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Angina Pectoris		Wellbutrin	PS	Glaxosmithkline	ORAL
4	DAY						

Date:01/28/05ISR Number: 4563413-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507534A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
2	WK						

Date:01/28/05ISR Number: 4563414-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507536A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	
8	MON	Anxiety Insomnia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563415-7Report Type:Periodic  
 Age:85 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0507885A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lethargy		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Three		Medication Error					
times per day	11	DAY	Pharmaceutical Product	Aspirin	C	Glaxosmithkline	
UNKNOWN		Complaint		Duragesic	C		
UNKNOWN				Synthroid	C	Glaxosmithkline	
UNKNOWN				Prozac	C		
UNKNOWN				Megace	C		
UNKNOWN				Ativan	C		
UNKNOWN				Neurotin	C		
UNKNOWN				Zantac	C	Glaxosmithkline	
UNKNOWN				Ativan	C		

Date:01/28/05ISR Number: 4563416-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0507935A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Restless Legs Syndrome		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563417-0Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508188A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Suicidal Ideation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:01/28/05ISR Number: 4563418-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508356A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	12 MON	Alopecia	Wellbutrin	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:01/28/05ISR Number: 4563419-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508403A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Dysphemia	Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563420-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508420A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Hospitalization - Initial or Prolonged		Depressed Level Of Consciousness	Wellbutrin	PS	Glaxosmithkline	
Other			Loss Of Consciousness	Unknown Medication	SS		
			Overdose				



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563421-2Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508446A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	6	MON		Wellbutrin	PS	Glaxosmithkline	ORAL
				Pepcid	C		

Date:01/28/05ISR Number: 4563422-4Report Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508470A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	3	YR		Wellbutrin	PS	Glaxosmithkline	ORAL
				Cellcept	C		
				Plaquenil	C		
				Vioxx	C		

Date:01/28/05ISR Number: 4563423-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508669A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563424-8Report Type:Periodic  
Age:10 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508737A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
37.5MG Twice per day	13	DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
				Risperdal	C		

Date:01/28/05ISR Number: 4563425-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0508954A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
		Weight Increased					

Date:01/28/05ISR Number: 4563426-1Report Type:Periodic  
Age:37 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0509072A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	
Other				Synthroid	C	Glaxosmithkline	
1 MON				Lamictal	C	Glaxosmithkline	
				Zyprexa	C		
				Prozac	C		
				Unspecified Inhaler	C		

RESPIRATORY

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563427-3Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509208A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	18 DAY	Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	ORAL
		Anorexia Myalgia		Trazodone	C		

Date:01/28/05ISR Number: 4563428-5Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509211A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Three		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
times per day	4 WK			Birth Control Pills	C		

Date:01/28/05ISR Number: 4563429-7Report Type:Periodic  
Age:45 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0509429A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 MON		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	
		Feeling Abnormal Insomnia					

Date:01/28/05ISR Number: 4563430-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509442A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 DAY		Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Interferon	C		

Date:01/28/05ISR Number: 4563431-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509670A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Haematochezia		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563432-7Report Type:Periodic  
Age: Gender:I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509848A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error Overdose		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563433-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510193A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Accidental Overdose Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563434-0Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510329A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 MON		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563435-2Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510401A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	17 DAY	Crying Disorientation Dizziness		Wellbutrin Effexor Blood Pressure Medication	PS C C	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563436-4Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510510A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Feeling Jittery		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563437-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510552A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 MON		Headache Tachycardia Ventricular Extrasystoles		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563438-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510657A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Weight Decreased					
per day	1 YR						

Date:01/28/05ISR Number: 4563439-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510913A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563440-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0510925A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL
				Lithium	SS	Glaxosmithkline	
				Bupropion			
				Hydrochloride	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563441-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511232A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Disorientation		Wellbutrin	PS	Glaxosmithkline	
		Drug Interaction		Gabitril	SS		ORAL
		Euphoric Mood		Estrogen	SS		
				Xanax	SS		
				Thyroid Medication	SS		
				Pain Medication	SS		

Date:01/28/05ISR Number: 4563442-XReport Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511445A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 YR		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
				Nicotine Patch	C	Glaxosmithkline	

Date:01/28/05ISR Number: 4563443-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511455A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 DAY	Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
		Insomnia		Neurontin	C		
		Night Sweats		Lamictal	C	Glaxosmithkline	
		Palpitations		Doxepin	C		
				Estrogen	C		
				Xanax	C		

Date:01/28/05ISR Number: 4563444-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511478A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
7 DAY		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563445-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511615A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563446-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0511888A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Screen Positive		Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563447-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512026A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Twice							
per day	2	DAY					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563448-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512089A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Sexual Dysfunction		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563449-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512220A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75MG Three times per day	6 YR	Acne Pharmaceutical Product Complaint		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563450-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0512243A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563451-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512245A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Twice per day	2 YR	Lethargy Muscle Twitching		Wellbutrin	PS	Glaxosmithkline	ORAL
				Trazodone	C		

Date:01/28/05ISR Number: 4563452-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0512335A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin	PS	Glaxosmithkline	
2 WK		Vomiting					

Date:01/28/05ISR Number: 4563453-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0512390A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	
				Zyban	SS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563454-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0512606A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	
Other							

Date:01/28/05ISR Number: 4563455-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0513171A  
 Age:45 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Asthenia					
		Blood Pressure Increased					
		Disturbance In Attention					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Drug Interaction Heart Rate Increased Vision Blurred	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day				Provigil	SS		ORAL
100MG Per day							

Date:01/28/05ISR Number: 4563456-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0513272A  
 Age:40 YR Gender:Female I/FU:F

		PT Hepatic Enzyme Increased	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration			Wellbutrin	PS	Glaxosmithkline	ORAL
Dose				No Concurrent Medication	C		
Other							

Date:01/28/05ISR Number: 4563457-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0513583A  
 Age: Gender:Female I/FU:F

		PT Insomnia	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration			Wellbutrin	PS	Glaxosmithkline	ORAL
Dose							
150MG Unknown		Parosmia Pharmaceutical Product Complaint					

Date:01/28/05ISR Number: 4563458-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0513634A  
 Age:23 YR Gender:Female I/FU:I

		PT Agitation	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration			Wellbutrin	PS	Glaxosmithkline	ORAL
Dose				No Concurrent Medication	C		
75MG Per day	2 WK	Feeling Jittery					

Date:01/28/05ISR Number: 4563459-5Report Type:Periodic  
Age:12 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0513647A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
6 WK		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563460-1Report Type:Periodic  
Age:25 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0513649A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Twice		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	29 DAY	Weight Increased					
YR				Zyprexa	C		ORAL
400MG Twice				Neurotin	C		ORAL
per day	YR						
YR				Cogentin	C		ORAL

Date:01/28/05ISR Number: 4563461-3Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0513830A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563462-5Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513952A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN		Loss Of Libido Orgasm Abnormal					

Date:01/28/05ISR Number: 4563463-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0513992A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gingivitis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563464-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514086A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypotrichosis		Wellbutrin	PS	Glaxosmithkline	
2	MON						

Date:01/28/05ISR Number: 4563465-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514280A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Asthenia Depressed Mood Drug Ineffective Emotional Disorder Mood Altered Sleep Disorder					

Date:01/28/05ISR Number: 4563466-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514287A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563467-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514326A  
Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Sodium Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL
				Xanax	C		
				Klonopin	C		

Date:01/28/05ISR Number: 4563468-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514616A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sexual Dysfunction		Wellbutrin	PS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563469-8Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514619A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	
				Pegasys	SS		
				Copegus	SS		

Date:01/28/05ISR Number: 4563470-4Report Type:Periodic  
 Age:10 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514684A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Paranoia		Wellbutrin	PS	Glaxosmithkline	ORAL
37.5MG Twice							
per day	2	MON		Risperdal	C		

Date:01/28/05ISR Number: 4563471-6Report Type:Periodic  
 Age: Gender:I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514695A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bruxism		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563472-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0514800A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563473-XReport Type:Periodic  
 Age:43 YR Gender:Male I/FU:I

Company Report #04US03216

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other  
Disturbance In Attention  
Insomnia  
Mania  
Wellbutrin PS Glaxosmithkline ORAL

Date:01/28/05ISR Number: 4563474-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514902A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bruxism		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563475-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0514905A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				No Concurrent Medication	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563476-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515037A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 WK		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
		Libido Decreased Motion Sickness Vomiting					

Date:01/28/05ISR Number: 4563477-7Report Type:Periodic  
 Age:29 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515131A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 11 DAY		Oral Mucosal Blistering		Wellbutrin	PS	Glaxosmithkline	ORAL
		Rash Generalised Stevens-Johnson Syndrome					

Date:01/28/05ISR Number: 4563478-9Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515135A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 MON		Prostatic Specific		Wellbutrin	PS	Glaxosmithkline	ORAL
		Antigen Increased					

Date:01/28/05ISR Number: 4563479-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515329A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anger Mental Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563480-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0515374A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	
				Multiple Medications	C		

Date:01/28/05ISR Number: 4563481-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0515566A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Increased Appetite		Wellbutrin	PS	Glaxosmithkline	ORAL
18 MON		Weight Increased					

Date:01/28/05ISR Number: 4563482-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0515706A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	
Other							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563483-2Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0515913A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erythema Multiforme		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563484-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516108A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG	Unknown			No Concurrent Medication	C		

Date:01/28/05ISR Number: 4563488-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516393A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dry Mouth		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563489-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516725A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
2 MON		Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563490-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516769A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Somnolence		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563491-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516775A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563492-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516908A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563493-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517076A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia Myalgia		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563494-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0517271A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		Tinnitus		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563495-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518180A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563496-0Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0518211A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563497-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518339A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 YR		Tinnitus		Wellbutrin	PS	Glaxosmithkline	ORAL
		White Blood Cell Count Decreased					

Date:01/28/05ISR Number: 4563498-4Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518512A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Per day	3 DAY	Abnormal Dreams		Bupropion	PS	Glaxosmithkline	ORAL

Insomnia

No Concurrent

Medication

C

Date:01/28/05ISR Number: 4563499-6Report Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0518530A

Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Screen Positive		Wellbutrin	PS	Glaxosmithkline	ORAL
				Zoloft	C		
				Ativan	C		
				Neurontin	C		
				Duragesic	C		
				Lortab	C		
				Restoril	C		
				Naprosyn	C		
				Ambien	C		
				Xanax	C		
				Tylenol Pm	C		

Date:01/28/05ISR Number: 4563500-XReport Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0518533A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
		Urticaria		Wellbutrin	SS	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563501-1Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0518557A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563502-3Report Type:Periodic  
Age: Gender:I I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518986A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
	8MG Per day			Warfarin	C	Glaxosmithkline	

Date:01/28/05ISR Number: 4563503-5Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #20040304012

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Idiopathic		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Thrombocytopenic Purpura					
200MG Twice		Platelet Count Decreased		Topamax	SS		
per day		300MG per day 317 DAY					
UNKNOWN				Lexapro	SS		
	20MG Per day						

Date:01/28/05ISR Number: 4563504-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519182A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563505-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0519370A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	OTHER
		Gastrooesophageal Reflux Disease					

Date:01/28/05ISR Number: 4563506-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0519402A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Interaction Tachycardia		Effexor	C		
				Strattera	C		

Date:01/28/05ISR Number: 4563507-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0519699A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563508-4Report Type:Periodic  
Age:26 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0519712A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 MON		Crying		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
75MG Per day	22 DAY	Depression		Wellbutrin	SS	Glaxosmithkline	ORAL
112.5MG per day		Diarrhoea		Effexor Xr	C		ORAL
	33 DAY	Mood Swings					
		Stool Analysis Abnormal		Trazodone	C		ORAL
5MG As required				Diazepam	C		ORAL
5MG As required	35 DAY			Diazepam	C		ORAL

Date:01/28/05ISR Number: 4563509-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519717A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
				Effexor	C		

Date:01/28/05ISR Number: 4563510-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519741A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563511-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0519842A

Outcome Dose Other 7 YR  
Duration  
PT Convulsion  
Report Source  
Product Wellbutrin  
Role PS  
Manufacturer Glaxosmithkline  
Route ORAL

Date:01/28/05ISR Number: 4563512-6Report Type:Periodic Company Report #S04-USA-04203-01  
Age: Gender: I/FU:I

Outcome Dose Other UNKNOWN  
Duration  
PT Mania Weight Increased  
Report Source  
Product Wellbutrin Lexapro  
Role PS SS  
Manufacturer Glaxosmithkline  
Route ORAL

Date:01/28/05ISR Number: 4563513-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0520110A  
Age: Gender:Male I/FU:I

Outcome Dose  
Duration  
PT Drug Ineffective Erectle Dysfunction Libido Decreased  
Report Source  
Product Wellbutrin Unknown Medication  
Role PS C  
Manufacturer Glaxosmithkline  
Route ORAL

Date:01/28/05ISR Number: 4563514-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0520365A  
Age:72 YR Gender:Female I/FU:I

Outcome Dose 300MG Unknown 6 MON  
Duration  
PT Agitation Anxiety  
Report Source  
Product Wellbutrin Nexium Ambien Provigil  
Role PS C C C  
Manufacturer Glaxosmithkline  
Route ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563515-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520404A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563516-3Report Type:Periodic  
Age:55 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0520765A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Unknown	YR	Ejaculation Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
		Sexual Dysfunction		Celexa	C		
				Unknown Medication	C		

Date:01/28/05ISR Number: 4563517-5Report Type:Periodic  
Age:50 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0521063A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	5 MON	Abdominal Pain Upper		Wellbutrin Xr	PS	Glaxosmithkline	ORAL
75MG Per day		Confusional State					
		Constipation		Effexor Xr	C		
		Dyspnoea					
		Erythema					
		Flatulence					
		Hyperhidrosis					
		Pharmaceutical Product Complaint					
		Stomach Discomfort					

Date:01/28/05ISR Number: 4563518-7Report Type:Periodic  
Age:77 YR Gender:Female I/FU:I

Company Report #DSA\_24592\_2004

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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		Dermatitis Exfoliative	Wellbutrin	PS	Glaxosmithkline	ORAL
		Disease Recurrence	Cardizem	SS	Glaxosmithkline	
UNKNOWN	240MG Per day	21 DAY				
		Hypersensitivity	Advair	C	Glaxosmithkline	
RESPIRATORY						
		Sensory Loss				
(INHALATION)		31 DAY				
		Skin Discolouration	Xopenex	C		
31 DAY						
		Skin Hypertrophy	Unknown Medication	C		
		Weight Decreased	Prednisone	C		
36 DAY						

Date:01/28/05ISR Number: 4563519-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0521404A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563520-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0521423A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Drug Ineffective					
		Drug Tolerance Decreased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563521-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521551A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563522-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521668A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		Amnesia YR Bradyphrenia Mental Impairment		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563523-0Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0521815A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2 WK		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563524-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521828A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 WK		Dyspnoea		Wellbutrin	PS	Glaxosmithkline	ORAL
				Hyzaar	C		
				Zocor	C		
				Norvasc	C		
				Flomax	C		

Date:01/28/05ISR Number: 4563525-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521839A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Liver Function Test		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN		Abnormal		No Concurrent Medication	C		

Date:01/28/05ISR Number: 4563526-6Report Type:Periodic Company Report #USA040567498  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563527-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0522170A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
450MG Per day		Diarrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Ineffective					
		Stool Analysis Abnormal					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563528-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #04US05065

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563529-1Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522638A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Wellbutrin	PS	Glaxosmithkline	ORAL
6	MON	Nervousness					

Date:01/28/05ISR Number: 4563530-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522658A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3	WK					

Date:01/28/05ISR Number: 4563531-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522662A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Burning Sensation					
		Pain In Extremity					

Date:01/28/05ISR Number: 4563532-1Report Type:Periodic  
 Age:29 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523513A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Somnolence		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563533-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0523769A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	
Other							

Date:01/28/05ISR Number: 4563534-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0523829A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lethargy		Wellbutrin	PS	Glaxosmithkline	ORAL
.5TAB Per day		Somnolence		Adderall	C		
		Thinking Abnormal		Valium	C		
		Throat Irritation		Vicodin	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563535-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524013A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
1 WK		Panic Attack Tremor					

Date:01/28/05ISR Number: 4563536-9Report Type:Periodic  
Age:66 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524060A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Unknown		Head Discomfort		Ativan	C		
				Potassium	C		
				Vitamins	C		

Date:01/28/05ISR Number: 4563537-0Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0524198A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Psychomotor Hyperactivity Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL
				Zantac 75	C	Glaxosmithkline	

Date:01/28/05ISR Number: 4563538-2Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524532A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Acne Weight Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563539-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524559A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Euphoric Mood		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563540-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0524675A  
Age: Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563541-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0524682A  
Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 WK		Anxiety Compulsions		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563542-4Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0524876A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Euphoric Mood		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563543-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525036A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Unknown	1	MON				

Date:01/28/05ISR Number: 4563545-XReport Type:Periodic  
Age:14 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525076A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Increased Tendency To		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG	Per day	1	YR				
		Bruise		Naprosyn	C		
				Naprosyn	C		
UNKNOWN							

Date:01/28/05ISR Number: 4563546-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525570A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
2	WK						
		Skin Disorder					
		Urticaria					

Date:01/28/05ISR Number: 4563547-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #S2004US11201

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased Feeling Abnormal Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563548-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525719A  
 Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Nicoderm Cq Clear 21mg	PS SS	Glaxosmithkline Glaxosmithkline	ORAL
TRANSDERMAL							

Date:01/28/05ISR Number: 4563549-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525815A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Macular		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563550-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525824A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG Three			Drug Ineffective	Wellbutrin	PS	Glaxosmithkline	ORAL
			Pharmaceutical Product				
times per day 1	1	YR					
150MG Twice			Complaint	Wellbutrin	SS	Glaxosmithkline	ORAL
per day	2	YR					
				Xanax	C		

Date:01/28/05ISR Number: 4563551-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526132A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Hormone Level Abnormal	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563552-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526433A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
				Paxil	C	Glaxosmithkline	
				Pamelor	C		

Date:01/28/05ISR Number: 4563553-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0526660A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Rash	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563554-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0526679A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563555-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0526703A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Jittery		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN				Zoloft	SS		
5	YR						

Date:01/28/05ISR Number: 4563556-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0526857A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Breast Discharge		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563557-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0527100A  
Age:43 YR Gender:Female I/FU:I

Outcome  
PT  
Feeling Jittery  
Heart Rate Increased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Insomnia Sleep Disorder				
Dose	Duration		Report Source	Product	Role	Manufacturer
75MG Twice				Wellbutrin	PS	Glaxosmithkline
per day	2 WK			Valtrex Vitamins	C C	Glaxosmithkline
						Route
						ORAL

Date:01/28/05ISR Number: 4563558-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0527280A  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline
75MG Twice		Nausea				Route
per day	2 WK	Palpitations		Valtrex	C	Glaxosmithkline

Date:01/28/05ISR Number: 4563559-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0527285A  
 Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline
3 MON				Wellbutrin Sr	SS	Glaxosmithkline
150MG Twice						Route
per day	3 MON			Wellbutrin Xl	SS	Glaxosmithkline
150MG Twice						ORAL
per day	3 MON			No Concurrent Medication	C	

Date:01/28/05ISR Number: 4563560-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0527363A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Polymenorrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
4	MON	Weight Decreased					

Date:01/28/05ISR Number: 4563561-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0527581A

Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
1	MON	Pyrexia		Trazodone	C		
				Lithium	C	Glaxosmithkline	
				Levoxyl	C	Glaxosmithkline	

Date:01/28/05ISR Number: 4563562-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0527622A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Liver Function Test		Wellbutrin	PS	Glaxosmithkline	ORAL
3	YR	Abnormal		Strattera	C		
				Adderall	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563563-1Report Type:Periodic  
Age:65 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0527971A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Wellbutrin	PS	Glaxosmithkline	ORAL
9	MON			Toprol	C		
				Celebrex	C		
				Zocor	C		
				Diuretic	C		

Date:01/28/05ISR Number: 4563564-3Report Type:Periodic  
Age:19 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528261A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563565-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528263A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Feeling Abnormal Stomach Discomfort		Bupropion	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563566-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528271A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anger		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG	Unknown	Depression Insomnia					

Date:01/28/05ISR Number: 4563567-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528393A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyskinesia		Wellbutrin Dilantin	PS C	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563568-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0528994A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Disturbance In Attention Nervousness		Wellbutrin Sudafed	PS SS	Glaxosmithkline Glaxosmithkline	ORAL ORAL

Date:01/28/05ISR Number: 4563569-2Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0528996A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563570-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529112A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Haemolytic Anaemia		Wellbutrin	PS	Glaxosmithkline	ORAL
				Neurontin	C		
				Oxycontin	C		

Date:01/28/05ISR Number: 4563571-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #04US02766 TEAM B

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563572-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529476A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563573-4Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0529502A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563574-6Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0529516A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
UNKNOWN		Dissociation		Wellbutrin	PS	Glaxosmithkline	
		Sensory Disturbance		Ethanol	SS		
				No Concurrent			

Medication

C

Date:01/28/05ISR Number: 4563575-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530024A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563576-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530046A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563577-1Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530107A

Outcome	PT
	Heart Rate Increased Insomnia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
			Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563578-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0530112A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Increased Appetite		Wellbutrin Zoloft	PS C	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563579-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0530113A  
 Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Therapeutic Response		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day	2 MON	Unexpected		Birth Control Pills	C		ORAL

Date:01/28/05ISR Number: 4563580-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0530116A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Rash Macular					

Date:01/28/05ISR Number: 4563581-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0530220A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

0 DAY Pain Bupropion PS Glaxosmithkline ORAL  
Luvox C

Date:01/28/05ISR Number: 4563582-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0530608A  
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menstrual Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	1 WK			No Concurrent Medication	C		

Date:01/28/05ISR Number: 4563583-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0530651A  
Age:22 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice							
per day	5 YR			Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day	2 MON			Wellbutrin	SS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563584-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530828A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
75MG Twice			Gastrointestinal Disorder				
per day	1	MON					

Date:01/28/05ISR Number: 4563585-0Report Type:Periodic  
Age: Gender:I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530903A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
			Stool Analysis Abnormal				

Date:01/28/05ISR Number: 4563586-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531074A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
2 DAY			Dizziness				
			Nausea				

Date:01/28/05ISR Number: 4563587-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531228A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
Other			Convulsion	Dilantin	C		

Date:01/28/05ISR Number: 4563588-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531307A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

1 WK Feeling Jittery Wellbutrin PS Glaxosmithkline ORAL  
1 WK Stratterra SS

Date:01/28/05ISR Number: 4563589-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0531446A  
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563590-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0531733A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Zyprexa	C		
300MG	Unknown						

Date:01/28/05ISR Number: 4563591-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0531777A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Stool Analysis Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563592-8Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531959A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563593-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532025A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Skin Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563594-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532033A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563595-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532361A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Pain Upper		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563596-5Report Type:Periodic  
Age:18 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532540A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL

3 WK

Vicodin	C	
Augmentin	C	Glaxosmithkline
Meridia	C	

Date:01/28/05ISR Number: 4563597-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532833A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563598-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532910A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Depression Formication Mental Disorder		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563599-0Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532916A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
1 MON		Amnesia		Wellbutrin	PS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563600-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533106A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563601-6Report Type:Periodic  
Age:75 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533249A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Unknown	3 WK	Fall		Aricept	C		
				Namenda	C		

Date:01/28/05ISR Number: 4563602-8Report Type:Periodic  
Age:43 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0533267A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Hepatic Enzyme Increased		No Concurrent			
600MG Unknown	4 MON	Jaundice		Medication	C		
		Mental Impairment					
		Panic Attack					
		Restlessness					

Date:01/28/05ISR Number: 4563603-XReport Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0533280A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mouth Ulceration		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563604-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533445A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
5 YR		Testicular Swelling		Wellbutrin	PS	Glaxosmithkline	ORAL
				Citalopram	C		

Date:01/28/05ISR Number: 4563605-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0533656A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	6 MON	Drug Ineffective		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Unknown	8 WK	Drug Interaction		Bupropion	SS	Glaxosmithkline	ORAL
300MG Per day		Therapeutic Response		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
UNKNOWN	875MG Unknown 14 DAY	Unexpected		Amoxicillin	SS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563606-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0533999A  
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aggression Agitation Insomnia		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563607-7Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534128A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75MG Twice		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
per day	7 MON			No Concurrent Medication	C		

Date:01/28/05ISR Number: 4563608-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534630A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG Three times per day		Bruxism Hair Plucking		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563609-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534675A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 MON		Drug Ineffective		Wellbutrin Wellbutrin Xl	PS C	Glaxosmithkline Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563610-7Report Type:Periodic  
 Age:49 YR Gender:Male I/FU:I

Company Report #04US06388

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Tinnitus Visual Acuity Reduced		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563611-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0534888A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	
4	DAY	Fatigue Insomnia Memory Impairment Nervousness Tremor					

Date:01/28/05ISR Number: 4563612-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0534895A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Tinnitus		Wellbutrin	PS	Glaxosmithkline	

Date:01/28/05ISR Number: 4563613-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0534903A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mood Swings		Wellbutrin Lithium Klonopin	PS C C	Glaxosmithkline Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4563614-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0535046A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Paraesthesia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563615-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0535147A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Screen Positive		Wellbutrin	PS	Glaxosmithkline	ORAL
2 YR				No Concurrent Medication	C		

Date:01/28/05ISR Number: 4563616-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0535399A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN							

Date:01/28/05ISR Number: 4563617-XReport Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0535403A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth Dry Throat Gingival Swelling Swollen Tongue		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563618-1Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0535728A

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563619-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535729A  
Age: Gender:Male I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Energy Increased Psychomotor Hyperactivity		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:01/28/05ISR Number: 4563620-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535741A  
Age: Gender: I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Convulsion		Wellbutrin	PS	Glaxosmithkline	
Other							

Date:01/28/05ISR Number: 4564339-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541320A  
Age: Gender:Male I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Photopsia		Effexor	SS		
450MG Per day							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4564340-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541324A  
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2	MON					
		Coprolalia		Testosterone Cream	C	Glaxosmithkline	
		Irritability					
		Judgement Impaired					
		Personality Change					

Date:01/28/05ISR Number: 4564341-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541340A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2	MON					
		Crying		Adderall Xr	C		
		Depression					
		Hallucination					
		Pharyngolaryngeal Pain					
		Sleep Disorder					
		Urticaria					

Date:01/28/05ISR Number: 4564343-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541534A  
Age:12 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	18	MON					
		Anger					
		Crying		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Per day	1	WK					
		Depression					
		Drug Ineffective					
		Tremor					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Feeling Abnormal Feeling Of Despair Medication Error Panic Attack Suicidal Ideation		Xanax	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300MG Per day		Atrioventricular Block Decreased Appetite		Lamictal Wellbutrin	PS SS	Glaxosmithkline Glaxosmithkline	ORAL ORAL
		Dyspepsia Nausea		No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4565967-XReport Type:Expedited (15-DaCompany Report #B0133705A  
 Age:34 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Required SEE DOSAGE	Arthralgia Blood Pressure Decreased Chills	Foreign Literature Health	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
Intervention to TEXT / ORAL Prevent Permanent Impairment/Damage	Diabetes Mellitus Inadequate Control Inflammation Oedema Peripheral	Professional	Insulin Lispro Injection (Insulin Lispro)	SS		
SUBCUTANEOUS	SUBCUTANEOUS Pyrexia Serum Sickness Urticaria Generalised Vomiting					

Date:01/28/05ISR Number: 4575333-9Report Type:Direct Company Report #CTU 238582  
 Age:50 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100 MG QAM	Agitation		Bupropion Sr 100g	PS		
	Disturbance In Attention Dyskinesia Eye Disorder Feeling Abnormal Pharmaceutical Product Complaint		Marinol Bextra Flovent Zelnorm Aciphex Serevent Hydrocodone/Apap Xanax	C C C C C C C		

Date:01/28/05ISR Number: 4575477-1Report Type:Direct Company Report #CTU 238549  
 Age:53 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 MG BID ,	Depression		Bupropion	PS		ORAL

PO

Hallucination, Auditory

Multiple Sclerosis

Benzotropine	C
Fluphenazine	C
Zyprexa	C
Lorazepam	C

Date:01/28/05ISR Number: 4575503-XReport Type:Direct  
Age:53 YR Gender:Male I/FU:I

Company Report #CTU 238593

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pharmaceutical Product		Bupropion	PS		ORAL
150 MG BID PO		Complaint					
		Tremor					

Date:01/28/05ISR Number: 4575510-7Report Type:Direct  
Age:33 YR Gender:Female I/FU:I

Company Report #CTU 238597

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Migraine		Bupropion Sr	150mg PS		ORAL
150MG QD		Pharmaceutical Product					
ORAL		Complaint		Neurontin	C		
				Alprazolam	C		
				Trileptal	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/31/05ISR Number: 4565496-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541550A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion Death Overdose		Wellbutrin	PS	Glaxosmithkline	

Date:01/31/05ISR Number: 4565502-6Report Type:Expedited (15-DaCompany Report #BPC-ZN-05-048

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Blood Cholesterol Increased Diabetes Mellitus Non-Insulin-Dependent Epilepsy Hypertension		Zyban	PS	Glaxosmithkline	ORAL

Date:01/31/05ISR Number: 4565514-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0363099A

Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG See dosage text	42 DAY	Abortion Induced Drug Exposure During Pregnancy		Zyban	PS	Glaxosmithkline	ORAL
TRANSDERMAL dosage text	68MG See	Drug Ineffective Drug Interaction Pregnancy Unintended Pregnancy		Implanon	SS		

Date:01/31/05ISR Number: 4565515-4Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0364300A

Age:68 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		2U per day	17 DAY	Abdominal Pain Upper	Zyban	PS	Glaxosmithkline	ORAL
UNKNOWN			1MG Unknown	Cholecystitis	Orap	C		
UNKNOWN			25MG Unknown	Hepatic Enzyme Increased	Anafranil	C		
UNKNOWN			600MG Unknown	Insomnia	Glucophage	C		
UNKNOWN				Pyrexia	Minidiab	C		

Date:01/31/05ISR Number: 4565516-6Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0364369A  
Age:58 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		per day	34 DAY	Abnormal Behaviour	Zyban	PS	Glaxosmithkline	
UNKNOWN			150MG Twice	Affect Lability				
				Delusion				
				Personality Change				
				Psychotic Disorder				

Date:01/31/05ISR Number: 4565521-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0365371A  
Age:53 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Twice	per day		Depression	Zyban	PS	Glaxosmithkline	ORAL
				Lethargy				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

UNKNOWN	80MG Per day	Propranolol	C
UNKNOWN	10MG Per day	Escitalopram	C

Date:01/31/05ISR Number: 4565529-4Report Type:Expedited (15-DaCompany Report #NA-GLAXOSMITHKLINE-B0366028A  
 Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Visual Disturbance		Zyban	PS	Glaxosmithkline	
UNKNOWN		8 DAY		No Concurrent Medication	C		

Date:01/31/05ISR Number: 4565532-4Report Type:Expedited (15-DaCompany Report #2005000353  
 Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Diarrhoea		Zyban	PS	Glaxosmithkline	ORAL
4500MG Single							
Initial or Prolonged		Hypotension					
dose							
Other		Intentional Misuse		Ethanol	SS		ORAL
		Overdose					
		Suicide Attempt					
		Tachycardia					

Date:01/31/05ISR Number: 4567958-1Report Type:Expedited (15-DaCompany Report #B0365424A  
 Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Aggression	Foreign	Wellbutrin Sr Tablet			
Initial or Prolonged		Depression	Literature	- Controlled Release			
		Irritability	Health	(Bupropion			
		Multiple Drug Overdose	Professional	Hydrochloride)	PS		ORAL
SEE DOSAGE		Paranoia					

TEXT / ORAL

Psychomotor Agitation  
 Psychotic Disorder  
 Midazolam Tablet  
 (Midazolam) SS ORAL

14 TABLET /  
 SINGLE DOSE /  
 ORAL

Respiratory Rate  
 Increased  
 Somnolence  
 Suicide Attempt

Date:01/31/05ISR Number: 4569930-4Report Type:Expedited (15-DaCompany Report #2005-124455-NL  
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia	Health	Remeron /Net/	PS		ORAL
Other		Confusional State	Professional	Intron A	SS	Schering-Plough	
45 MG QD ORAL		Decreased Appetite		"Schering-Plough"			
INTRAVENOUS	44 MIU	Dehydration					
INTRAVENOUS		Depression					
(NOS)		Drug Interaction		Wellbutrin	SS		ORAL
300 MG BID		Lethargy					
ORAL		Malignant Melanoma Stage		Allopurinol	SS		ORAL
100 MG QD		Iii					
ORAL		Nausea		Clonazepam	SS		ORAL
0.5 MG QD		Oral Intake Reduced					
ORAL				Trazodone	SS		ORAL
50 MG QD ORAL				Zocor	SS		ORAL
10 MG QD ORAL				Glucosamine	SS		
DF PRN ORAL				Multivitamins	SS		ORAL
DF PRN ORAL							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

10 MG PRN  
 ORAL  
 Compazine SS ORAL

Date:02/01/05ISR Number: 4566620-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541516A  
 Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged TRANSDERMAL		Application Site Vesicles Chest Pain		Nicoderm Cq Clear 21mg	PS	Glaxosmithkline	
150MG per day		Electrocardiogram Change		Wellbutrin	SS	Glaxosmithkline	ORAL
		Heart Rate Decreased Nicotine Dependence Ventricular Extrasystoles		Ativan	C		ORAL

Date:02/01/05ISR Number: 4567164-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0534891A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation Anger Belligerence Delusional Disorder, Persecutory Type Emotional Disorder Muscle Spasms Psychotic Disorder Tension		Wellbutrin	PS	Glaxosmithkline	

Date:02/01/05ISR Number: 4567169-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0536385A  
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 450MG Per day		Complex Partial Seizures		Wellbutrin XL	PS	Glaxosmithkline	ORAL

Initial or Prolonged Hemiparesis  
Ill-Defined Disorder  
Medication Error  
Overdose

Date:02/01/05ISR Number: 4567181-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0542715A  
Age:87 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Fall					
		Hip Fracture					

Date:02/01/05ISR Number: 4567199-8Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0364711A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Zyban	PS	Glaxosmithkline	ORAL
1TAB Variable		Feeling Abnormal					
dose	6	DAY					
		Formication		Keflex	C	Glaxosmithkline	ORAL
500MG Four		Hallucination, Tactile					
times per day	6	DAY					
		Palpitations					
		Tremor					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/05ISR Number: 4567213-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0369201A  
 Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged dosage text	9 WK	Inflammation  Myositis  Pain In Extremity Therapeutic Response Decreased Vasculitis		Zyban   Diane 35	PS   C	Glaxosmithkline	ORAL   ORAL

Date:02/01/05ISR Number: 4578191-1Report Type:Direct Company Report #CTU 238840  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1 PO DAILY		Nausea  Vomiting		Wellbutrin Xl 150 Mg	PS		ORAL

Date:02/02/05ISR Number: 4568287-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541948A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 10 MON		Hepatic Enzyme Increased  Hepatic Neoplasm Malignant Hepatocellular Damage Lung Neoplasm Malignant		Wellbutrin  Lipitor Biaxin	PS  C C	Glaxosmithkline	ORAL

Date:02/02/05ISR Number: 4568291-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0542699A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Suicidal Ideation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/02/05ISR Number: 4568296-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0363857A  
 Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthralgia Asthenia Back Pain		Zyban Sodium Stibogluconate	PS  SS	Glaxosmithkline  Glaxosmithkline	
INTRAVENOUS day	1391MG 15	Per Cerebrovascular Accident					
500MG Three times per day	16	Chest Pain Dizziness		Keflex	C	Glaxosmithkline	ORAL
200MG Per day	34	Fatigue		Fluconazole	C		ORAL
		Flank Pain Headache Limb Discomfort Muscle Spasms Myalgia Nausea Renal Failure Acute Rhabdomyolysis Vomiting		Rifampin Ketoconazole	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/02/05ISR Number: 4578096-6Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 238908

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Zyban	PS		ORAL
150 MG PO 1							
PO QD X3D							
THEN 2 / D							

Date:02/02/05ISR Number: 4578112-1Report Type:Direct  
Age:26 YR Gender:Male I/FU:I

Company Report #CTU 238899

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Hypersensitivity Stevens-Johnson Syndrome		Wellbutrin Xl 150 Mg Tab	PS		ORAL
1 TAB PO QD							
Vomiting							

Date:02/02/05ISR Number: 4578364-8Report Type:Direct  
Age:48 YR Gender:Female I/FU:I

Company Report #CTU 238928

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Chest Pain Palpitations		Wellbutrin (Bupropion)	PS		
Depakote C							
Atenolol C							
Vitamins C							

Date:02/03/05ISR Number: 4569330-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0540290A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Unknown							
Medication Error							
No Concurrent							

Panic Attack

Medication

C

Date:02/03/05ISR Number: 4569331-9Report Type:Expedited (15-DaCompany Report #BPC-ZN-05-048

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day		Blood Cholesterol	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged Disability			Increased Diabetes Mellitus Non-Insulin-Dependent Epilepsy Hypertension				

Date:02/03/05ISR Number: 4569332-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0542637A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	UNKNOWN		Suicide Attempt	Wellbutrin	PS	Glaxosmithkline	
Initial or Prolonged Disability	UNKNOWN			Amitriptyline	SS		
	UNKNOWN			Fluvoxamine	SS		
	UNKNOWN			Unknown Medication	SS		
	UNKNOWN			Phenazopyridine	SS		
	UNKNOWN			Pyridium	SS		
	UNKNOWN			Unknown Medication	SS		
	UNKNOWN			Zyprexa	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Alcohol

SS

Date:02/03/05ISR Number: 4569333-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0542840A  
Age:50 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Coma		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged	Hallucination		Zoloft	C		
	Renal Failure		Neurontin	C		
	Somnolence		Zyrtec	C	Glaxosmithkline	
			Synthroid	C	Glaxosmithkline	

Date:02/03/05ISR Number: 4569344-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0363180A  
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Tetany		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown						
Initial or Prolonged						

Date:02/03/05ISR Number: 4569354-XReport Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0045476A  
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Anxiety		Zyban	PS	Glaxosmithkline	ORAL
150MG See						
Initial or Prolonged	Cardiac Disorder					
dosage text	Delusion					
	Dyspnoea					
	Hyperventilation					
	Panic Attack					
	Suicidal Ideation					

Date:02/03/05ISR Number: 4570536-1Report Type:Direct  
Age:35 YR Gender:Female I/FU:I

Company Report #CTU 239213

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
3 TABS Q DAY		Drug Ineffective		Bupropion Hcl 100	PS		
		Pharmaceutical Product Complaint					

Date:02/04/05ISR Number: 4570585-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0534857A  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3 DAY		Anger		Tagamet	C	Glaxosmithkline	
Initial or Prolonged		Anorexia		Vicodin	C		
		Bipolar Disorder					
		Coordination Abnormal					
		Diarrhoea					
		Disturbance In Attention					
		Fatigue					
		Nausea					



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Freedom Of Information (FOI) Report

Date:02/04/05ISR Number: 4570588-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0540443A

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300MG See Initial or Prolonged dosage text	Blood Pressure Decreased  Coma		Wellbutrin	PS	Glaxosmithkline	ORAL
	Drug Ineffective		Lexapro	SS		
	Drug Interaction		Atenolol	C		
	Medication Error		Avandia	C	Glaxosmithkline	
			Nexium	C		
			Aspirin	C	Glaxosmithkline	
			Aricept	C		
			Neurontin	C		
			Centrum Silver	C		
			Magnesium	C		
			Potassium	C		
			Duragesic	C		
			B12	C	Glaxosmithkline	
			Hydrocodone	C		
			Zocor	C		
			Diazepam	C		
			Isosorbide	C		

Date:02/04/05ISR Number: 4570592-0Report Type:Expedited (15-DaCompany Report #S05-USA-00304-01

Age:22 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged UNKNOWN	Disease Recurrence Pancreatitis		Wellbutrin Lexapro	PS SS	Glaxosmithkline	ORAL

Date:02/04/05ISR Number: 4570608-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0365867A

Age:56 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG per day 30 DAY	Atrial Fibrillation		Zyban	PS	Glaxosmithkline	ORAL

Initial or Prolonged	Dyspnoea	Vagifem	C	
UNKNOWN				
	Joint Swelling	Combivent	C	
UNKNOWN				
	Tachycardia	Warfarin Sodium	C	Glaxosmithkline
UNKNOWN				
		Estriol	C	
UNKNOWN				

Date:02/04/05ISR Number: 4570614-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0369124A

Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	300MG per day 9 DAY	Bradycardia		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Creatinine Renal		Vaccine	SS	Glaxosmithkline	
SUBCUTANEOUS	2UNIT Single	Clearance Decreased					
dose	1 DAY	Cytolytic Hepatitis		Lescol	C		ORAL
1UNIT per day		Dyspnoea Exertional					
		Oedema					
		Oedema Peripheral					

Date:02/04/05ISR Number: 4574444-1Report Type:Expedited (15-DaCompany Report #2004111620

Age:73 YR Gender:Female I/FU:F

Outcome  
Hospitalization -  
Initial or Prolonged

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Freedom Of Information (FOI) Report

Disability Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 200 MG (200 MG, 1 AS NECESSARY), ORAL		Abdominal Pain Lower	Health	Celebrex (Celecoxib)	PS		ORAL
		Abdominal Tenderness	Professional				
		Asthenia					
		Cerebrovascular Accident					
SEE IMAGE		Constipation		Bupropion			
		Diastolic Dysfunction		Hydrochloride			
		Disorientation		(Bupropion			
		Fall		Hydrochloride)	SS		
		Hyperreflexia		Biselect (Bisoprolol			
		Meningioma		Fumarate,			
		Peripheral Coldness		Hydrochlorothiazide)	C		
		Petechiae		Nifedipine			
		Pyuria		(Nifedipine)	C		
		Rash		Diazepam (Diazepam)	C		
		Somnolence		Propacet			
		Treatment Noncompliance		(Dextropropoxyphene			
		Urinary Tract Infection		Napsilate,			
		Vasculitis		Paracetamol)	C		

Date:02/04/05ISR Number: 4578288-6Report Type:Direct Company Report #CTU 239311  
Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Q DAY X 3 DAYS THEN BID [~1 WEEK ]	1 WK	Rash Generalised		Zyban 150 Mg	PS		

Date:02/04/05ISR Number: 4578297-7Report Type:Direct Company Report #CTU 239304  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Q DAY X 3 Initial or Prolonged DAYS THEN BID		Rash		Zyban 150 Mg	PS		
[~1 WEEK]	1	WK					

Date:02/07/05ISR Number: 4572411-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0543321A  
Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100MG Twice per day		Liver Function Test Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice per day		Protein Total Increased		Lamictal	C	Glaxosmithkline	

Date:02/07/05ISR Number: 4572430-9Report Type:Expedited (15-DaCompany Report #DK-GLAXOSMITHKLINE-B0369735A  
Age:69 YR Gender:I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN	150MG	Mouth Ulceration Unknown 25 DAY		Zyban	PS	Glaxosmithkline	
UNKNOWN		Stevens-Johnson Syndrome		Centyl + Kcl	C		
UNKNOWN	80MG	Unknown		Micardis	C	Glaxosmithkline	
UNKNOWN	100MG	Unknown		Selo-Zok	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/07/05ISR Number: 4575893-8Report Type:Expedited (15-DaCompany Report #004003

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Consumer	Fluoxetine(Fluoxetine)	PS		ORAL
20 MG, QD,		Disorientation	Health	e) Capsule, 20mg			
ORAL		Drug Interaction	Professional				
		Loss Of Consciousness		Wellbutrin - Slow			
		Nausea		Release (Bupropion			
				Hydrochloride) 150			
150 MG, QD,				Mg	SS		ORAL
ORAL							

Date:02/08/05ISR Number: 4572631-XReport Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0531757A

Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
37.5MG Per		Abdominal Pain		Paxil Cr	PS	Glaxosmithkline	ORAL
day	16 MON	Abnormal Dreams					
		Diarrhoea		Wellbutrin	SS	Glaxosmithkline	ORAL
		Disorientation		Clonazepam	C		
		Headache					
		Nausea					
		Paraesthesia					
		Vision Blurred					

Date:02/08/05ISR Number: 4572649-7Report Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0532717A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination		Paxil	PS	Glaxosmithkline	ORAL
37.5MG Per							
day							

300MG Per day				Wellbutrin	SS	Glaxosmithkline	ORAL
				Geodon	C		

Date:02/08/05ISR Number: 4572657-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0533119A  
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
37.5MG Per			Abdominal Distension	Paxil Cr	PS	Glaxosmithkline	ORAL
day	8	MON	Affect Lability				
100MG Twice			Nausea	Wellbutrin Sr	SS	Glaxosmithkline	ORAL
per day	6	WK	Nightmare				
			Sleep Disorder	No Concurrent			
			Vertigo	Medication	C		
			Weight Increased				

Date:02/08/05ISR Number: 4572672-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0534324A  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
25MG Per day			Dizziness	Paxil Cr	PS	Glaxosmithkline	ORAL
150MG Per day	7	DAY	Hot Flush	Wellbutrin Xl	SS	Glaxosmithkline	ORAL
			Hyperhidrosis				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/08/05ISR Number: 4573207-0Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0519822A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 30MG Per day		Dizziness		Paxil	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Drug Withdrawal Syndrome		Wellbutrin	SS	Glaxosmithkline	
		Feeling Abnormal		Flexeril	SS		
		Intentional Misuse		Skelaxin	SS		
		Suicidal Ideation		Antiinflammatory	SS		
		Suicide Attempt		Aspirin	SS	Glaxosmithkline	
				Theophylline	SS		
				Risperdal	SS		
				Cold Medicine	SS		

Date:02/08/05ISR Number: 4573474-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523468A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
20MG Per day	4 MON	Confusional State		Paxil	PS	Glaxosmithkline	ORAL
3 DAY		Food Craving		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Somnolence					
		Weight Increased					

Date:02/08/05ISR Number: 4573502-5Report Type:Periodic  
Age:76 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524232A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
50MG Per day	5 YR	Diarrhoea		Paxil	PS	Glaxosmithkline	ORAL
		Drug Ineffective		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
				Zocor	C		

Date:02/08/05ISR Number: 4573505-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524392A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety		Paxil	PS	Glaxosmithkline	ORAL
		Drug Exposure During Pregnancy		Wellbutrin	SS	Glaxosmithkline	ORAL

Date:02/08/05ISR Number: 4573522-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0525070A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety		Paxil	PS	Glaxosmithkline	ORAL
		Libido Decreased		Wellbutrin	SS	Glaxosmithkline	ORAL
		Nervousness					
		Tremor					

Date:02/08/05ISR Number: 4573606-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0527300A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT
		Blood Creatinine Increased
		Drug Ineffective
		Fatigue
		Lethargy



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Malaise Weight Decreased Weight Increased	Report Source	Product	Role	Manufacturer	Route
40MG Per day	10 YR			Paxil	PS	Glaxosmithkline	ORAL
300MG Per day	4 MON			Wellbutrin	SS	Glaxosmithkline	ORAL
				Benecol	C		
				Aspirin	C	Glaxosmithkline	
				Maxide	C		
				Ranitidine	C	Glaxosmithkline	

Date:02/08/05ISR Number: 4574107-2Report Type:Periodic  
Age:72 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530234A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
20MG Per day	6 WK	Blood Glucose Increased		Paxil	PS	Glaxosmithkline	ORAL
150MG Per day	1 DAY	Drug Ineffective		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
				Insulin Lantus	C		
				Novolog	C		
				Plavix	C		
				Hytrin	C		
				Accupril	C		
				Cordarone	C		
				Lanoxin	C	Glaxosmithkline	
				Coreg	C	Glaxosmithkline	
				Methotrexate	C		
				Crestor	C		
				Triamterene + Hydrochlorothiazide	C	Glaxosmithkline	
				Centrum Silver	C		
				Baby Aspirin	C	Glaxosmithkline	
				Calcium	C		
				Folic Acid	C		

Date:02/08/05ISR Number: 4574113-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530368A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anorexia		Paxil	PS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	ORAL

Date:02/08/05ISR Number: 4574117-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0530648A  
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anorexia		Paxil	PS	Glaxosmithkline	ORAL
20MG Per day		Depression		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Per day	1 MON	Mental Impairment		Wellbutrin Xl	C	Glaxosmithkline	
		Somnolence		Atrovent	C	Glaxosmithkline	
		Vision Blurred		Advair	C	Glaxosmithkline	
				Hytrin	C		

Date:02/08/05ISR Number: 4574119-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0530823A  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness		Paxil	PS	Glaxosmithkline	ORAL
20MG Per day	18 MON	Dysgeusia		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
300MG Per day							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

No Concurrent Medication C

Date:02/08/05ISR Number: 4574149-7Report Type:Periodic  
Age:51 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0531838A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
40MG Per day	2 YR	Apathy		Paxil	PS	Glaxosmithkline	ORAL
300MG Per day		Decreased Interest		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Depression					
		Lethargy					
		Sleep Disorder					
		Therapeutic Response					
		Unexpected					

Date:02/08/05ISR Number: 4574173-4Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533261A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
20MG Per day	10 YR	Drug Ineffective		Paroxetine	PS	Glaxosmithkline	ORAL
		Hyperhidrosis		Paxil	SS	Glaxosmithkline	ORAL
12.5MG Per day	2 MON	Insomnia		Paxil	SS	Glaxosmithkline	ORAL
300MG Per day	6 YR			Wellbutrin	SS	Glaxosmithkline	ORAL
300MG In the morning	2 MON			Wellbutrin	SS	Glaxosmithkline	ORAL
				Trazodone	C		
				Multivitamin	C		
				Fibercon	C		

Date:02/08/05ISR Number: 4574608-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0515151A  
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorgasmia		Paxil	PS	Glaxosmithkline	ORAL
3 MON		Drug Ineffective		Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day	6 MON	Insomnia		No Concurrent Medication	C		

Date:02/08/05ISR Number: 4574623-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0515534A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Affect Lability		Paxil	PS	Glaxosmithkline	ORAL
5 YR		Drug Ineffective		Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day		Libido Decreased Sexual Dysfunction Weight Increased					

Date:02/08/05ISR Number: 4574684-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0516755A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
		Nausea		Paxil	SS	Glaxosmithkline	ORAL
				Flonase	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Allegra C

Date:02/08/05ISR Number: 4574811-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0517089A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
12.5MG Per day	8 DAY	Drug Ineffective		Paxil Cr	PS	Glaxosmithkline	ORAL
150MG Twice per day	1 MON	Fatigue Increased Appetite		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:02/08/05ISR Number: 4575065-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544253A  
 Age: Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/08/05ISR Number: 4575078-5Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0364991A  
 Age: Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 150MG Twice per day		Completed Suicide Gun Shot Wound		Zyban	PS	Glaxosmithkline	

Date:02/08/05ISR Number: 4575086-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0369079A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Suicidal Ideation		Zyban	PS	Glaxosmithkline	ORAL

Date:02/09/05ISR Number: 4575616-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0349816A  
 Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardiac Arrest		Bupropion	PS	Glaxosmithkline	ORAL
		Completed Suicide		Aspirin	SS	Glaxosmithkline	ORAL
		Convulsion		Acetaminophen	SS	Glaxosmithkline	ORAL
		Haematemesis		Ethanol	SS		ORAL
		Overdose					

Date:02/09/05ISR Number: 4575617-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0349817A  
 Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardio-Respiratory Arrest		Bupropion	PS	Glaxosmithkline	ORAL
		Completed Suicide		Topiramate	SS		ORAL
		Intentional Misuse		Venlafaxine	SS		ORAL
		Overdose		Olanzapine	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/09/05ISR Number: 4575637-XReport Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0370110A  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG per day	DAY	Disability	Zyban	PS	Glaxosmithkline	ORAL
			Dyspnoea				
			Exercise Tolerance				
			Decreased				
			Heart Rate Increased				
			Malaise				
			Palpitations				
			Panic Attack				
			Visual Disturbance				

Date:02/09/05ISR Number: 4575660-5Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0045685A  
 Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Twice		Neutropenia	Zyban	PS	Glaxosmithkline	ORAL
	per day			Betabion	C		ORAL
	100MG Twice			Diazepam	C		
	per day			Vitamin B 12	C	Glaxosmithkline	ORAL
	UNKNOWN						
	1000UG Per						
	day						

Date:02/09/05ISR Number: 4577254-4Report Type:Expedited (15-DaCompany Report #2004AL000696  
 Age:15 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Aggression	Children'S Apap			
			Agitation	Elixir			
			Literature				
			Health				

PO	Anion Gap Abnormal Atrioventricular Block	Professional	(Acetaminophen) (Alpharma)	PS	Alpharma	ORAL
PO	Base Excess Increased Bradycardia Cardiac Arrest		Infants' Apap Drops (Acetaminophen) (Alpharma)	SS	Alpharma	ORAL
PO	Completed Suicide		Bupropion	SS		ORAL
PO	Convulsion		Asa	SS		ORAL
PO	Grand Mal Convulsion Multiple Drug Overdose Ph Body Fluid Decreased		Fluoxetine Capsules, 20 Mg (Siegfried Ltd.)	SS	Siegfried Ltd	ORAL
PO	Pyrexia Ventricular Extrasystoles		Fluoxetine Oral Solution Usp, 20mg/5ml (Alpharma)	SS	Alpharma	ORAL
PO			Caffeine	SS		ORAL
PO			Valproic Acid	SS		ORAL
PO			Sertraline	SS		ORAL
PO			Aspirin/Acetaminophe n/Caffeine	SS		ORAL

Date:02/09/05ISR Number: 4578279-5Report Type:Expedited (15-DaCompany Report #BASF-05-001  
Age:61 YR Gender:Male I/FU:I

Outcome PT  
Other Asthenia  
Confusional State  
Decreased Appetite  
Dehydration

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Freedom Of Information (FOI) Report

Dose	Duration	Drug Interaction Lethargy Nausea Oral Intake Reduced	Report Source	Product	Role	Manufacturer	Route
INTRAVENOUS	44 MU 5X		Health Professional	Intron A	PS		
WEEKLY							
INTRAVENOUS				Wellbutrin	SS		ORAL
300 MG BID							
ORAL							

Date:02/09/05ISR Number: 4578400-9Report Type:Expedited (15-DaCompany Report #S05-USA-00523-01  
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Catheter Related Infection	Consumer	Lexapro (Escitalopram)	PS		ORAL
20 MG QD PO		Coma Drug Interaction Hypotension Sepsis		Lexapro (Escitalopram)	SS		
300 MG QD PO				Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
150 MG QD PO				Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
150 MCG BID				Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
PO				Atenolol	C		
				Isosorbide	C		
				Potassium	C		
				Zocor	C		
				Aspirin	C		
				Avandia			

(Rosiglitazone	
Maleate)	C
Nexium	
(Esomeprazole)	C
Aricept (Donepezil	
Hydrochloride)	C
Neurontin	
(Gabapentin)	C
Duragesic (Fentanyl)	C
Hydrocodone	C
Centrum Silver	C
Magnesium	C
Vitamin B12	C
Diazepam	C

Date:02/09/05ISR Number: 4579248-1Report Type:Direct  
 Age: Gender: I/FU:I

Company Report #CTU 239876

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error Pharmaceutical Product Complaint		Bupropion Er 150mg Er Eon Labs	PS	Eon Labs	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/10/05ISR Number: 4577336-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0543706A

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Contusion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 3 DAY						
Initial or Prolonged	Convulsion		Iron	C		
	Impaired Driving Ability					
	Tongue Biting					

Date:02/10/05ISR Number: 4577344-6Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0544312A

Age:22 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Abnormal Dreams		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice						
Initial or Prolonged	Foaming At Mouth					
per day						
	Grand Mal Convulsion		Diane	C		
	Insomnia					
	Urinary Incontinence					

Date:02/10/05ISR Number: 4577345-8Report Type:Expedited (15-DaCompany Report #CHPA2005US00542

Age:11 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Suicidal Ideation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/10/05ISR Number: 4579610-7Report Type:Expedited (15-DaCompany Report #2004AL000507

Age:51 YR Gender:Female I/FU:F

Outcome	PT
Death	Aggression
	Alanine Aminotransferase
	Increased
	Aspartate
	Aminotransferase
	Increased

Atrioventricular Block  
First Degree  
Base Excess Negative  
Bilirubin Conjugated  
Increased  
Blood Albumin Decreased  
Blood Alkaline  
Phosphatase Increased  
Blood Bicarbonate  
Decreased  
Blood Bilirubin Increased  
Blood Chloride Increased  
Blood Creatine  
Phosphokinase Increased  
Blood Glucose Increased  
Blood Lactate  
Dehydrogenase Increased  
Blood Ph Decreased  
Blood Pressure Decreased  
Bundle Branch Block Right  
Completed Suicide  
Electrocardiogram Pr  
Prolongation  
Electrocardiogram Qrs

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Complex Prolonged Generalised Oedema Grand Mal Convulsion Hepatic Steatosis					
		Multiple Drug Overdose Oxygen Saturation Decreased Pco2 Decreased Pco2 Increased	Literature Health Professional	Propoxyphene Napsylate And Acetaminophen Tablets Usp, 100 Mg/650 Mg (Purepac)	PS	Purepac	ORAL
PO		Po2 Decreased Pyrexia		Alprazolam Tablets Usp, 2 Mg (Purepac)	SS	Purepac	ORAL
PO		Qrs Axis Abnormal		Zolpidem	SS		ORAL
PO		Splenomegaly		Ethanol	SS		ORAL
PO		Urine Output Decreased		Bupropion	SS		ORAL
PO				Quetiapine	SS		ORAL

Date:02/10/05ISR Number: 4579916-1Report Type:Expedited (15-DaCompany Report #2004AL000680  
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alanine Aminotransferase Increased	Literature Health	Alprazolam Tablets Usp, 2 Mg (Purepac)	PS	Purepac	ORAL
PO		Aspartate Aminotransferase Increased Blood Creatine Phosphokinase Increased	Professional	Propoxyphene Napsylate And Acetaminophen Tablets Usp, 100 Mg/650 Mg (Purepac)	SS	Purepac	ORAL
PO		Blood Urea Increased		Bupropion	SS		ORAL
PO		Coma		Quetiapine	SS		ORAL
PO		Completed Suicide Drug Ineffective Electrocardiogram St Segment Elevation Hepatic Steatosis					

Multiple Drug Overdose  
Pulmonary Oedema  
Pupillary Reflex Impaired  
Splenomegaly

Date:02/11/05ISR Number: 4578715-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544234A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3	MON					
		Lethargy		Seroquel	C		
		Sedation		Asthma Medication	C		

Date:02/11/05ISR Number: 4578717-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544500A  
Age:11 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardio-Respiratory Arrest		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	11	MON					
Life-Threatening				Strattera	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/11/05ISR Number: 4578718-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544527A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Overdose		Wellbutrin Adderall	PS C	Glaxosmithkline	ORAL

Date:02/11/05ISR Number: 4578726-9Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0365820A

Age:64 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Ageusia		Zyban	PS	Glaxosmithkline	
UNKNOWN	150MG	Twice					
		Anosmia					
per day							
UNKNOWN	2CAP	Dysgeusia		Multivitamin	C		
		per day					
		Hypertension					
		Rhinorrhoea					

Date:02/11/05ISR Number: 4578736-1Report Type:Expedited (15-DaCompany Report #CH-GLAXOSMITHKLINE-B0370275A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Zyban	PS	Glaxosmithkline	ORAL
		Drug Exposure During					
		Pregnancy					

Date:02/14/05ISR Number: 4580227-9Report Type:Expedited (15-DaCompany Report #004003

Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Sensation In Eye		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day							
		Anxiety		Prozac	SS		
20MG Per day							
		Confusional State					

Disorientation  
Hypoaesthesia Oral  
Loss Of Consciousness  
Nausea  
Tremor

Date:02/14/05ISR Number: 4580235-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544531A  
Age:6 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Hallucination		Metadate Cd	C		
		Insomnia		Clonidine	C		
		Medication Error					
		Pharmaceutical Product					
		Complaint					

Date:02/14/05ISR Number: 4580260-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0370023A  
Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Discomfort					
per day	18 DAY	Hypersensitivity					
		Scar					

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Freedom Of Information (FOI) Report

Date:02/14/05ISR Number: 4580274-7Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0045476A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG See Initial or Prolonged dosage text		Anxiety		Zyban	PS	Glaxosmithkline	ORAL
25MG Unknown		Cardiac Disorder Delusion Dysphagia Dyspnoea Hallucination, Visual Hyperventilation Hypoaesthesia Panic Attack Suicidal Ideation Throat Tightness		Saroten	C	Glaxosmithkline	ORAL

Date:02/14/05ISR Number: 4580951-8Report Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0516772A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Lamictal Wellbutrin	PS SS	Glaxosmithkline Glaxosmithkline	ORAL ORAL

Date:02/14/05ISR Number: 4580986-5Report Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0518076A

Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
25MG Per day	4 DAY	Cognitive Disorder		Lamictal	PS	Glaxosmithkline	ORAL
150MG In the morning	3 YR	Confusional State Coordination Abnormal Speech Disorder		Wellbutrin Xl	SS	Glaxosmithkline	ORAL

Date:02/14/05ISR Number: 4581081-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0520555A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Lamictal	PS	Glaxosmithkline	ORAL
100MG Per day				Paxil	SS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	ORAL
				Provigil	C		

Date:02/14/05ISR Number: 4581089-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0520760A  
Age:19 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion		Lamictal	PS	Glaxosmithkline	ORAL
Other				Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day							

Date:02/14/05ISR Number: 4581363-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0528567A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Lamictal	PS	Glaxosmithkline	ORAL
				Eskalith	SS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	ORAL
				Parnate	SS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/05ISR Number: 4581546-2Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533960A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
25MG Twice		Drug Ineffective		Lamictal	PS	Glaxosmithkline	ORAL
per day	21 DAY	Lethargy					
		Weight Increased		Wellbutrin	SS	Glaxosmithkline	ORAL
				Lanoxin	C	Glaxosmithkline	
				Lisinopril	C		
				Claritin	C		

Date:02/14/05ISR Number: 4581856-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504104A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
225MG Per day	5 MON	Cough		Lamictal	PS	Glaxosmithkline	ORAL
200MG Per day		Dry Eye		Wellbutrin	SS	Glaxosmithkline	ORAL
		Influenza		Trazodone	C		
		Muscle Twitching					
		Photosensitivity Reaction					

Date:02/14/05ISR Number: 4581861-2Report Type:Periodic  
Age:57 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0504308A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
25MG Four		Nervousness		Lamictal	PS	Glaxosmithkline	ORAL
times per day	2 MON	Tremor					
300MG Per day	2 WK			Wellbutrin	SS	Glaxosmithkline	ORAL
				Klonopin	C		
				Depakote	C		

Date:02/14/05ISR Number: 4581891-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505135A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Lamictal	PS	Glaxosmithkline	ORAL
100MG Per day	6	MON		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
100MG Twice							
per day				Wellbutrin	C	Glaxosmithkline	

Date:02/14/05ISR Number: 4581909-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0505666A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Pruritic		Lamictal	PS	Glaxosmithkline	ORAL
50MG Per day	16	DAY		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
150MG Per day	1	DAY		Lexapro	C		

Date:02/14/05ISR Number: 4581922-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0506127A  
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anaemia		Lamictal	PS	Glaxosmithkline	ORAL
Hospitalization -							
100MG Three							
Initial or Prolonged							
times per day							
Other				Wellbutrin Sr	SS	Glaxosmithkline	ORAL
450MG per day				Topamax	C		
				Levothroid	C	Glaxosmithkline	

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Freedom Of Information (FOI) Report

Reglan	C	Glaxosmithkline
Singulair	C	
Cytomel	C	Glaxosmithkline
Insulin	C	
Coreg	C	Glaxosmithkline
Allegra D	C	
Iron	C	

Date:02/14/05ISR Number: 4581925-3Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0506219A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Pruritus		Lamictal	PS	Glaxosmithkline	ORAL
50MG Twice		Rash Papular					
per day	3 WK			Wellbutrin Xl	SS	Glaxosmithkline	ORAL
				Trileptal	C		

Date:02/14/05ISR Number: 4581938-1Report Type:Periodic  
Age:41 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0513406A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Lamictal	PS	Glaxosmithkline	ORAL
2 WK		Feeling Abnormal		Wellbutrin	SS	Glaxosmithkline	ORAL
		Headache		Herb Drug	SS		
		Hunger		Ultram	SS		
		Nausea		Fioricet	SS		
				Allegra	C		
				Ambien	C		
				Synthroid	C	Glaxosmithkline	
				Midodrin	C		

Date:02/15/05ISR Number: 4582687-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541516A  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged TRANSDERMAL 150MG per day	Application Site Vesicles Chest Pain  Electrocardiogram Change  Heart Rate Decreased Nicotine Dependence Ventricular Extrasystoles	Nicoderm Cq Clear 21mg  Wellbutrin  Ativan	PS  SS  C	Glaxosmithkline  Glaxosmithkline	ORAL  ORAL
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Date:02/15/05ISR Number: 4582769-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544511A  
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day	6	MON	Transient Ischaemic Attack	Wellbutrin  Paxil No Concurrent Medication	PS  SS  C	Glaxosmithkline  Glaxosmithkline	ORAL  ORAL

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Freedom Of Information (FOI) Report

Date:02/15/05ISR Number: 4582771-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544769A

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Convulsion		Wellbutrin	PS	Glaxosmithkline	NASAL
Initial or Prolonged	Medication Error		Robitussin	C		
	Overdose					

Date:02/15/05ISR Number: 4582788-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0369124A

Age:54 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Bradycardia		Zyban	PS	Glaxosmithkline	ORAL
300MG per day 9 DAY						
Initial or Prolonged	Creatinine Renal		Vaccine	SS	Glaxosmithkline	
SUBCUTANEOUS 2UNIT Single	Clearance Decreased					
dose 1 DAY	Cytolytic Hepatitis		Lescol	C		ORAL
1UNIT per day	Dyspnoea Exertional					
	Haemoglobin Decreased					
	Oedema					
	Oedema Peripheral					
	Weight Increased					

Date:02/15/05ISR Number: 4588414-0Report Type:Direct

Company Report #CTU 240438

Age:37 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Abnormal Behaviour		Wellbutrin Xl 300			
Hospitalization -	Anger		Mg Glaxo Smith			
Initial or Prolonged	Balance Disorder		Kline	PS	Glaxo Smith Kline	ORAL
ONCE A DAY						
Required	Bipolar Disorder					
ORAL						
Intervention to	Condition Aggravated					
Prevent Permanent	Depression					
Impairment/Damage	Impaired Work Ability					
	Memory Impairment					

Date:02/16/05ISR Number: 4584195-5Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0545331A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Constipation Faeces Hard Rectal Haemorrhage		Zyban	PS	Glaxosmithkline	ORAL

Date:02/16/05ISR Number: 4584200-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0364304A  
Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 1TAB See Initial or Prolonged dosage text		Blood Glucose Increased  Dizziness  Dry Mouth		Zyban	PS	Glaxosmithkline	ORAL
2.5MG Per day		Electrocardiogram Abnormal Feeling Abnormal Hypoaesthesia Somnolence Visual Disturbance		Tritace	C		ORAL





ORAL				Topiramate (Topiramate)	SS		ORAL
ORAL				Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	SS		ORAL
ORAL				Olanzapine (Olanzapine)	SS		ORAL

Date:02/16/05ISR Number: 4589364-6Report Type:Expedited (15-DaCompany Report #B0349818A  
Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Acidosis Blood Ph Increased Brain Oedema Coma	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL		Completed Suicide Convulsion		Topiramate (Topiramate)	SS		ORAL
ORAL		Heart Rate Decreased Hypotension		Clonidine (Clonidine)	SS		ORAL
ORAL		Multiple Drug Overdose Tachycardia		Valproic Acid (Valproic Acid)	C		ORAL

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Freedom Of Information (FOI) Report

Date:02/16/05ISR Number: 4589365-8Report Type:Expedited (15-DaCompany Report #B0349825A

Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Completed Suicide Drug Toxicity Toxicologic Test Abnormal	Literature Health Professional	Wellbutrin Sr Tablet - Controlled Release (Bipropion Hydrochloride)	PS		ORAL
ORAL				Quetiapine (Formulation Unknown) (Quetiapine)	SS		ORAL
ORAL				Diazepam (Formulation Unknown) (Diazepam)	SS		ORAL

Date:02/16/05ISR Number: 4590024-6Report Type:Expedited (15-DaCompany Report #B0349819A

Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Creatine Phosphokinase Increased Blood Creatinine Increased	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL				Alprazolam (Alprazolam)	SS		ORAL
ORAL		Completed Suicide Electrocardiogram St		Paracetamol (Acetaminophen)	SS		ORAL
ORAL		Segment Elevation Hepatic Steatosis		Dextropropoxyphene (Dextropropoxyphene)	SS		ORAL
ORAL		Multiple Drug Overdose Pulmonary Oedema		Quetiapine (Quetiapine)	SS		ORAL
ORAL		Splenomegaly					

Date:02/16/05ISR Number: 4590025-8Report Type:Expedited (15-DaCompany Report #B0349826A  
Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Acute Respiratory Distress Syndrome Completed Suicide Grand Mal Convulsion	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL		Overdose Paralysis		Temazepam (Temazepam)	SS		ORAL
ORAL		Pneumonia Pneumonia Aspiration Pneumothorax Somnolence					

Date:02/16/05ISR Number: 4590026-XReport Type:Expedited (15-DaCompany Report #B0349823A  
Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anoxic Encephalopathy Cardiac Arrest Coma Convulsion	Literature Health Professional	Wellbutrin Sr Tablet - Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL		Pyrexia		Clonazepam (Clonazepam)	SS		ORAL
ORAL							

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ORAL Alprazolam (Alprazolam) SS ORAL

Date:02/16/05ISR Number: 4590027-1Report Type:Expedited (15-DaCompany Report #B0349824A  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Pressure Decreased Cardiac Arrest Completed Suicide Convulsion	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
150 MG/ ORAL		Ejection Fraction Decreased		Paxil (Paroxetine Hydrochloride)	SS		ORAL
20 MG/ORAL		Intentional Misuse Mental Impairment		Citalopram (Citalopram)	SS		ORAL
20 MG / ORAL		Mental Status Changes Nausea		Clonazepam (Clonazepam)	SS		ORAL
.5 MG / ORAL		Ventricular Dysfunction					

Date:02/16/05ISR Number: 4590028-3Report Type:Expedited (15-DaCompany Report #B0349820A  
Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Ethanol Increased Blood Ph Decreased Blood Pressure Decreased Completed Suicide	Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		ORAL
ORAL		Convulsion		Ethanol (Alcohol)	SS		ORAL
ORAL		Drug Level Above Therapeutic		Sertraline (Sertraline)	SS		ORAL
ORAL		Drug Toxicity Heart Rate Increased Hypoxic Encephalopathy					

Intentional Self-Injury  
Myoclonus

Date:02/16/05ISR Number: 4590029-5Report Type:Expedited (15-DaCompany Report #B0349821A  
Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest Completed Suicide Convulsion	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL

ORAL

Date:02/16/05ISR Number: 4590030-1Report Type:Expedited (15-DaCompany Report #B0349822A  
Age:15 YR Gender:Male I/FU:F

Outcome	PT
Death	Aggression Agitation Atrioventricular Block Blood Chloride Decreased Blood Creatine Phosphokinase Increased Blood Creatine Phosphokinase Mb

FDA - Adverse Event Reporting System (AERS)

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Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
		Increased Blood Glucose Increased Blood Potassium Decreased Body Temperature					
		Increased Bradycardia Cardiac Arrest Completed Suicide	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)			
ORAL					PS		ORAL
		Convulsion		Aspirin (Aspirin)	SS		ORAL
ORAL							
		Incoherent Muscle Rigidity		Paracetamol (Acetaminophen)	SS		ORAL
ORAL							
		Toxicologic Test Abnormal		Caffeine (Caffeine)	SS		ORAL
ORAL							
		Ventricular Extrasystoles		Fluoxetine (Fluoxetine)	SS		ORAL
ORAL							
				Valproic Acid (Valproic Acid)	SS		ORAL
ORAL							
				Sertraline (Sertraline)	SS		ORAL
ORAL							

Date:02/17/05ISR Number: 4584846-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0443202A  
 Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
1	YR						
		Insomnia Therapeutic Response Unexpected		Provigil Betaseron	C C		

Date:02/17/05ISR Number: 4584847-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0495725A  
 Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

300MG Per day 6 MON	Drug Ineffective	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Sinemet	C		
		Comtan	C		
		Mirapex	C		
		Flomax	C		
		Lasix	C	Glaxosmithkline	
		Synthroid	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4584848-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0500360A  
 Age:82 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200MG Per day 2 YR		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Blood Pressure Increased		Lexapro	SS		ORAL
		Decreased Appetite		Aspirin	C	Glaxosmithkline	
		Drug Ineffective		Zocor	C		
		Energy Increased		Prilosec	C	Glaxosmithkline	
		Stress		Klonopin	C		ORAL
		Weight Decreased					

Date:02/17/05ISR Number: 4584849-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0515731A  
 Age:46 YR Gender:Female I/FU:F

Outcome	PT
	Dizziness
	Drug Ineffective



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		Nausea Pain					
Dose	Duration		Report Source	Product	Role	Manufacturer	Route
150MG Per day	7 MON			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Synthroid	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4584850-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0517689A  
 Age:39 YR Gender:Female I/FU:F

Outcome		PT					
Dose	Duration		Report Source	Product	Role	Manufacturer	Route
300MG Per day	2 MON	Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Synthroid	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4584851-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0518540A  
 Age: Gender:Female I/FU:F

Outcome		PT					
Dose	Duration		Report Source	Product	Role	Manufacturer	Route
150MG Twice		Metrorrhagia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
per day	7 MON			Provera	C		
				Atenolol	C		
				Hormone	C		

Date:02/17/05ISR Number: 4584852-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0528459A  
 Age: Gender:Male I/FU:F

Outcome		PT					
Dose	Duration		Report Source	Product	Role	Manufacturer	Route
Other		Agitation Hypertension Hypertensive Crisis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584853-2Report Type:Periodic  
Age:28 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0530922A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
7 DAY							
Initial or Prolonged							

Date:02/17/05ISR Number: 4584854-4Report Type:Periodic  
Age:35 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0531288A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	4 WK	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
20MG Unknown		Dizziness		Paxil	C	Glaxosmithkline	ORAL
		Dry Mouth		Aspirin	C	Glaxosmithkline	
		Headache		Cold Medication	C		
		Malaise					
		Nausea					

Date:02/17/05ISR Number: 4584855-6Report Type:Periodic  
Age:44 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0531413A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 MON			Lamictal	C	Glaxosmithkline	ORAL

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UNKNOWN				Lorazepam	C		
UNKNOWN				Keppra	C		
				Unknown Medication	C		

Date:02/17/05ISR Number: 4584856-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0531548A  
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
119 DAY							

Date:02/17/05ISR Number: 4584857-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532020A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4584858-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532055A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
UNKNOWN		Therapeutic Response		Wellbutrin Xl	PS	Glaxosmithkline	
		Decreased		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4584859-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532064A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Per day	Dizziness	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	Feeling Hot And Cold	Lexapro	C		
	Rash	Lithium	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4584860-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532065A  
 Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	12 WK	Convulsion		Aciphex	C		
		Grand Mal Convulsion		Nasonex	C		
				Hydrocodone	C		
				Ultram	C		
				Zomig	C		

Date:02/17/05ISR Number: 4584861-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532189A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
1 WK		Irritability		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Lexapro	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584862-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532195A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 DAY		Dizziness Heart Rate Increased Somnolence		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584863-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532196A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Glucose Increased Drug Interaction		Wellbutrin Xl Glipizide Metformin	PS SS SS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584864-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532197A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness Headache Hot Flush Nausea		Wellbutrin Xl Lexapro	PS C	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584865-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532363A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 YR		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584866-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532508A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584867-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532509A  
Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fatigue		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584868-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532511A  
Age: Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

150MG In the

morning 1 MON

Celexa C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584869-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532562A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	2 MON	Drug Ineffective	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Zetia	C		
				Allegra	C		

Date:02/17/05ISR Number: 4584870-2Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532572A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Stool Analysis Abnormal	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584871-4Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532573A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Unknown	4 MON	Alopecia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584872-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532681A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	5 DAY		Headache	Wellbutrin	PS	Glaxosmithkline	ORAL
			Nausea	Celexa	C		
			Stomach Discomfort				
			Tinnitus				

Date:02/17/05ISR Number: 4584873-8Report Type:Periodic  
 Age:33 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532702A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3	WK	Anxiety	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Drug Ineffective	Klonopin	C		
				Risperdal	C		
				Metoprolol	C		

Date:02/17/05ISR Number: 4584874-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532712A  
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3	WK	Blood Pressure Increased	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4584875-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532716A  
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day			Diplopia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Prednisone	C		
				Coumadin	C	Glaxosmithkline	
				Codeine	C		
				Methotrexate	C		
				Furosemide	C	Glaxosmithkline	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Celebrex	C	
Vitamin Complex	C	
Folic Acid	C	
Vit D	C	
Vit C	C	Glaxosmithkline

Date:02/17/05ISR Number: 4584876-3Report Type:Periodic  
 Age:15 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532720A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6	MON		Ambien	C		

Date:02/17/05ISR Number: 4584877-5Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532721A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Somnolence		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2	WK		Prozac	C		

Date:02/17/05ISR Number: 4584878-7Report Type:Periodic  
 Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532730A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Exposure During		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Pregnancy		Claritin	C		
		Headache		Valium	C		

Date:02/17/05ISR Number: 4584879-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532733A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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300MG Per day 12 MON Headache Wellbutrin Xl PS Glaxosmithkline ORAL  
Paraesthesia Hormone Replacement C

Date:02/17/05ISR Number: 4584880-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532830A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584881-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532831A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 5 MON				Atenolol	C		
				Klonopin	C		
				Xanax	C		
				Luvox	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584882-9Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532837A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	1TAB Per day	3 MON	Erectile Dysfunction	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Zyprexa	C		ORAL

Date:02/17/05ISR Number: 4584883-0Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0532840A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Convulsion	Wellbutrin	PS	Glaxosmithkline	
Other							

Date:02/17/05ISR Number: 4584884-2Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532850A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Amnesia	Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584885-4Report Type:Periodic  
 Age:11 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532851A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	3 WK	Nightmare	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Depakote	C		
				Risperdal	C		
				Ritalin	C		

Date:02/17/05ISR Number: 4584886-6Report Type:Periodic  
 Age:21 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532861A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR						

Date:02/17/05ISR Number: 4584887-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532866A  
Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Alkaline		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	5 MON	Phosphatase Increased					

Date:02/17/05ISR Number: 4584888-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532868A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
450MG Per day							

Date:02/17/05ISR Number: 4584889-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532874A  
Age:30 YR Gender:Female I/FU:I

Outcome	PT
	Accidental Overdose
	Dizziness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Feeling Abnormal

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300MG Per day			Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584890-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532918A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sluggishness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2 WK				Paxil	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4584891-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0532919A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584892-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0533057A  
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Eructation Paraesthesia		Zetia	C		

Date:02/17/05ISR Number: 4584893-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0533066A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584894-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533070A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Breath Odour		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
5 YR		Libido Decreased					

Date:02/17/05ISR Number: 4584895-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0533079A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Twice		Nausea					
per day	14 DAY						

Date:02/17/05ISR Number: 4584896-9Report Type:Periodic  
Age:46 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0533091A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 MON			Celebrex	C		
				Prednisone	C		
UNKNOWN	10MG Variable						

dose

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Date:02/17/05ISR Number: 4584901-XReport Type:Periodic  
Age:41 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533248A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	7 DAY	Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Bruxism		Ambien	C		
		Dehydration					
		Depression					
		Heart Rate Increased					
		Hyperhidrosis					
		Insomnia					
		Pollakiuria					

Date:02/17/05ISR Number: 4584902-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533255A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584903-3Report Type:Periodic  
Age:42 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0533286A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	239 DAY						

Date:02/17/05ISR Number: 4584904-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533288A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584905-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533407A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anxiety Heart Rate Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584906-9Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533423A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	5 DAY	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Anger Tremor		Zelnorm	C		

Date:02/17/05ISR Number: 4584907-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533436A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Arthralgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584908-2Report Type:Periodic  
Age:39 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533437A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	6	WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4584909-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533444A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	4	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Effexor	C		
				Restoril	C		
				Ativan	C		
				Ibuprofen	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584910-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533448A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Leukopenia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Transaminases Increased		Paxil	SS	Glaxosmithkline	
				Plaquenil	C		
				Coumadin	C	Glaxosmithkline	
				Avinza	C	Glaxosmithkline	
				Topamax	C		
				Valium	C		
				Lasix	C	Glaxosmithkline	
				Benadryl	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4584911-2Report Type:Periodic  
Age:16 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533450A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4584912-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533451A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Unknown		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584913-6Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533454A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 WK	Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hot Flush		Meclizine Hcl	C		

Hyperhidrosis  
Nausea

Diclofenac

C

Date:02/17/05ISR Number: 4584914-8Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0533457A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	29 DAY			Lipitor	C		
				Lorazepam	C		

Date:02/17/05ISR Number: 4584915-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533581A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584916-1Report Type:Periodic  
Age:40 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0533582A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 WK		Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Fatigue		Nexium	C		
		Headache		Claritin	C		
		Pruritus					
		Somnolence					

Date:02/17/05ISR Number: 4584917-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533603A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	4 WK	Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Cardizem	C	Glaxosmithkline	
				Dyazide	C	Glaxosmithkline	
				Kdur	C	Glaxosmithkline	
				Glyburide	C		
				Toprol Xl	C		
				Pravachol	C		
				Xanax	C		

Date:02/17/05ISR Number: 4584918-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533620A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oedema		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Prozac	SS		

Date:02/17/05ISR Number: 4584919-7Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533745A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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150MG Twice	Dyspepsia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
per day	3 MON				
150MG Twice		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
per day					
		Synthroid	C	Glaxosmithkline	
		Prevacid	C		

Date:02/17/05ISR Number: 4584920-3Report Type:Periodic  
 Age:75 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533746A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Balance Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Contusion		Namenda	C		
		Depressive Symptom		Folic Acid	C		
		Drug Ineffective		B1 (Vitamin)	C		
		Dry Mouth		Multivitamin	C		
		Fall		Aricept	C		
				Vytone	C		
				Diovan	C		
				B12	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584921-5Report Type:Periodic  
Age:59 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533757A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	6 DAY	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Anorexia		Aspirin	C	Glaxosmithkline	
		Anxiety					
		Heart Rate Increased					
		Hyperhidrosis					
		Nausea					
		Tremor					

Date:02/17/05ISR Number: 4584922-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533763A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 MON		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Visual Disturbance					

Date:02/17/05ISR Number: 4584923-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533821A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1 WK	Abnormal Dreams		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584924-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533822A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584925-2Report Type:Periodic  
Age:74 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533943A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2	WK		Effexor Xr	C		
				Prevacid	C		
				Verapamil	C		
				Atenolol	C		

Date:02/17/05ISR Number: 4584926-4Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533946A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1	YR		Clonazepam	C		
		Chest Pain		Flonase	C	Glaxosmithkline	
		Dry Eye		Ranitidine	C	Glaxosmithkline	
		Gastrooesophageal Reflux		Allegra-D	C		
		Disease		Hydrocone	C		
		Stomatitis					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584927-6Report Type:Periodic  
Age:16 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533949A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Photosensitivity Reaction		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 YR			No Concurrent Medication	C		

Date:02/17/05ISR Number: 4584928-8Report Type:Periodic  
Age:55 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533951A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Sensation Of Foreign Body		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	6 WK	Sinusitis		Celexa	C		
				Trazodone	C		

Date:02/17/05ISR Number: 4584929-XReport Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533963A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Dizziness		Cymbalta	C		
UNKNOWN	40MG Per day	Orthostatic Hypotension					

Date:02/17/05ISR Number: 4584930-6Report Type:Periodic  
Age:20 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533965A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 DAY	Respiration Abnormal		Yasmin	C		
		Tinnitus		Vitamin	C		
				Kondremul	C		

Date:02/17/05ISR Number: 4584931-8Report Type:Periodic  
Age:19 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0533967A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other		Grand Mal Convulsion		Depakote	SS		ORAL
500MG Twice							
per day				Phenergan	C	Glaxosmithkline	ORAL
				Reglan	C	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584932-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534019A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584933-1Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534117A

Outcome PT  
Dyspnoea  
Hypertension

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Insomnia					
		Tachycardia					
		Tinnitus	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Weight Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	9 WK			Allegra	C		
				Nexium	C		

Date:02/17/05ISR Number: 4584934-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0534132A  
 Age: Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Loss Of Libido		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Dose				Unspecified Medications	C		
300MG Per day	2 WK						

Date:02/17/05ISR Number: 4584935-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0534137A  
 Age:54 YR Gender:Female I/FU:F

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Dose		Stool Analysis Abnormal		Zoloft	C		ORAL
450MG Per day	5 MON			Valium	C		
5MG Four							
times per day				Trazolan	C		
.5MG Four							
times per day				Aromasin	C		
400MG At							
night							

Date:02/17/05ISR Number: 4584936-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0534138A  
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oesophageal Rupture		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:02/17/05ISR Number: 4584937-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0534140A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day							
				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4584938-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0534143A  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 2 MON							
				Birth Control Pills	C		
				Nasocort	C		
				Astelin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584939-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534145A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK						

Date:02/17/05ISR Number: 4584940-9Report Type:Periodic  
Age:25 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534150A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	17 DAY	Rash		Oral Contraceptives	C		
				Multivitamin	C		

Date:02/17/05ISR Number: 4584941-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534186A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
7 MON							

Date:02/17/05ISR Number: 4584942-2Report Type:Periodic  
Age: Gender:I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0534314A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:02/17/05ISR Number: 4584943-4Report Type:Periodic  
Age:36 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0534316A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization -	Pruritus	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 20	DAY				
Initial or Prolonged	Rash	Effexor Xr	C		ORAL
1	YR				
Other	Urticaria	Vitamin E	C		ORAL
400IU Three					
times per day					
20MG Three		Inderal	C		ORAL
times per day 1	YR				
80MG Twice		Geodon	C		ORAL
per day 1	YR				
2MG Three		Artane	C		ORAL
times per day 1	YR				

Date:02/17/05ISR Number: 4584944-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0534321A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Effexor	C		

Date:02/17/05ISR Number: 4584945-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0534356A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2	DAY						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584946-XReport Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534475A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6 WK						

Date:02/17/05ISR Number: 4584947-1Report Type:Periodic  
 Age:62 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534484A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	5 WK	Drug Ineffective		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4584948-3Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534494A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK			No Concurrent Medication	C		

Date:02/17/05ISR Number: 4584949-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534496A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	5 MON	Drug Ineffective		Xanax	C		
Initial or Prolonged		Insomnia		Inderal	C		
		Mood Swings		Prilosec	C	Glaxosmithkline	
		Nervousness		Percocet	C		
		Panic Attack					
		Weight Decreased					

Date:02/17/05ISR Number: 4584950-1Report Type:Periodic  
Age:34 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0534502A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day	2 WK	Anxiety	Wellbutrin	PS	Glaxosmithkline	ORAL
	100MG Per day		Insomnia	Wellbutrin	SS	Glaxosmithkline	
			Somnolence				

Date:02/17/05ISR Number: 4584951-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534633A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	2 WK		Crying	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Insomnia				
			Nervousness				

Date:02/17/05ISR Number: 4584952-5Report Type:Periodic  
Age:15 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0534649A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Unknown		Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584953-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0534650A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Unknown	1 YR						

Date:02/17/05ISR Number: 4584954-9Report Type:Periodic  
Age:59 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534656A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Weight Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 MON			Hormone Replacement Therapy	C		

Date:02/17/05ISR Number: 4584955-0Report Type:Periodic  
Age: Gender:I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0534658A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				Viagra	C		

Date:02/17/05ISR Number: 4584956-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534666A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2 WK				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4584957-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534668A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	11 DAY	Ex-Smoker		Albuterol	C	Glaxosmithkline	
		Tinnitus					

Date:02/17/05ISR Number: 4584959-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0534669A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Muscle Twitching		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584960-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0534674A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Palpitations		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Wellbutrin Sr	C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584961-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534694A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Memory Impairment		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Unknown	14 DAY						

Date:02/17/05ISR Number: 4584962-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534695A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blepharospasm		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584963-XReport Type:Periodic  
Age:62 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0534704A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Pain		Nabumetone	SS	Glaxosmithkline	
				Wellbutrin	SS	Glaxosmithkline	ORAL
				Wellbutrin Sr	C	Glaxosmithkline	ORAL
150MG Twice							
per day				Vitamin	C		
				Minerals	C		

Date:02/17/05ISR Number: 4584964-1Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534834A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day		Weight Decreased		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4584965-3Report Type:Periodic  
Age:57 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534841A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
37.5MG Per		Headache					
day	1 WK	Tremor					

Date:02/17/05ISR Number: 4584966-5Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0534850A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Photopsia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584967-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534866A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	20 DAY						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584968-9Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0534880A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	8 MON	Tinnitus	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Simply Sleep	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4584969-0Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0534906A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Anorgasmia Libido Decreased	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584970-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0535005A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	1 YR	Mood Swings Suicidal Ideation	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584971-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0535021A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	1 YR	Headache Nausea	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Paxil Cr	C	Glaxosmithkline	
				Birth Control Pills	C		

Date:02/17/05ISR Number: 4584972-0Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0535028A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK			Yasmin	C		

Date:02/17/05ISR Number: 4584973-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535047A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2 YR							

Date:02/17/05ISR Number: 4584974-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535149A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Renal Atrophy		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR	Renal Disorder		Flomax	C		

Date:02/17/05ISR Number: 4584975-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535160A  
 Age:22 YR Gender:Male I/FU:I

Outcome PT  
 Agitation  
 Energy Increased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Thinking Abnormal

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3 WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Lithium	C	Glaxosmithkline	
			Seroquel	C		
			Protonix	C		

Date:02/17/05ISR Number: 4584976-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535366A  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3 WK	Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dizziness		No Concurrent Medication	C		
		Pruritus					
		Rash					

Date:02/17/05ISR Number: 4584977-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535385A  
 Age:42 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1 WK	Eating Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hyperacusis		Ativan	C		
		Insomnia					

Date:02/17/05ISR Number: 4584978-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535391A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	6 WK	Migraine		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584979-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535395A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Stool Analysis Abnormal		Wellbutrin Xl No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584980-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535432A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xr	PS	Glaxosmithkline	ORAL
10	DAY						

Date:02/17/05ISR Number: 4584981-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535433A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG In the							
morning	17	DAY		Valium	SS		
5MG Four							
times per day				Celexa	C		
4	YR						

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584982-3Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0535560A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gynaecomastia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4584983-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0535563A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK			No Concurrent Medication	C		

Date:02/17/05ISR Number: 4584984-7Report Type:Periodic  
Age:41 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0535566A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	30 DAY			No Concurrent Medication	C		

Date:02/17/05ISR Number: 4584985-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0535568A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK						

Date:02/17/05ISR Number: 4584986-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0535577A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice							
per day				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4584987-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535587A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 WK	Nausea					

Date:02/17/05ISR Number: 4584988-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535589A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
125MG In the							
morning	1 WK						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584989-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0535611A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Intolerance		Wellbutrin Xl Alcohol	PS C	Glaxosmithkline	ORAL ORAL

Date:02/17/05ISR Number: 4584990-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #56965

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin Xl Wellbutrin Sr Bupropion	PS SS SS	Glaxosmithkline Glaxosmithkline Glaxosmithkline	ORAL ORAL ORAL

Date:02/17/05ISR Number: 4584991-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0535628A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3 WK		Nausea					

Date:02/17/05ISR Number: 4584992-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0535715A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	8 DAY	Palpitations		Furosemide	C	Glaxosmithkline	
		Paraesthesia		Atenolol	C		

Date:02/17/05ISR Number: 4584993-8Report Type:Periodic  
Age:33 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0535736A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3	WK	Drug Ineffective	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Fatigue	No Concurrent Medication	C		
			Hunger				
			Loss Of Libido				

Date:02/17/05ISR Number: 4584994-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535740A  
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3	WK	Gastrooesophageal Reflux Disease	Wellbutrin	PS	Glaxosmithkline	ORAL
				Ritalin	C		

Date:02/17/05ISR Number: 4584995-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535766A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Convulsion	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4584996-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0535870A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3	WK					

Date:02/17/05ISR Number: 4584997-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0535872A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	4	WK		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4584998-7Report Type:Periodic  
Age:43 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0535876A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other				Adderall Xr	C		
300MG Per day				Prozac	C		
60MG Per day				Alprazolam	C		
60MG Per day				Ambien	C		
1TAB Twice							
per day							
10MG Per day							

Date:02/17/05ISR Number: 4584999-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0535877A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

300MG Per day                      Dyskinesia                      Wellbutrin Xl                      PS                      Glaxosmithkline                      ORAL

Date:02/17/05ISR Number: 4585000-3Report Type:Periodic                      Company Report #US-GLAXOSMITHKLINE-A0535884A  
 Age:58 YR    Gender:Female                      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2    DAY	Urticaria		Effexor Xr	C		
				Prosom	C		
				Lexapro	C		
				Benadryl	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585001-5Report Type:Periodic                      Company Report #US-GLAXOSMITHKLINE-A0535885A  
 Age:                      Gender:Female                      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Benzodiazepine	C		
450MG Per day							

Date:02/17/05ISR Number: 4585002-7Report Type:Periodic                      Company Report #US-GLAXOSMITHKLINE-A0535909A  
 Age:10 YR    Gender:Female                      I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fear		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3    WK	Personality Change					



Alopecia

Wellbutrin Xl

PS

Glaxosmithkline

ORAL

Klonopin

C

Date:02/17/05ISR Number: 4585008-8Report Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0536023A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fall		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	7 DAY	Headache		Topamax	C		
				Levothyroxine	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585010-6Report Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0536041A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Psychomotor Hyperactivity		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Prozac	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585011-8Report Type:Periodic  
Age:41 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536042A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypertension		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1 DAY			Atenolol	C		
				Xanax	C		
				Anti Inflammatory	C		

Date:02/17/05ISR Number: 4585012-XReport Type:Periodic  
Age:80 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536043A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sensory Disturbance		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6 DAY			Digoxin	C	Glaxosmithkline	
				Lanoxin	C	Glaxosmithkline	
				Actos	C		
				Glyburide	C		

Date:02/17/05ISR Number: 4585013-1Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0536175A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585014-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536220A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585015-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536240A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
450MG Unknown		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Unknown	3 MON			Wellbutrin Sr	SS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585016-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0536507A  
 Age:52 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG In the morning	3 WK	Energy Increased Insomnia Psychomotor Hyperactivity		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Temazepam	C		

Date:02/17/05ISR Number: 4585017-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0536511A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Twice per day	4 DAY	Overdose		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585018-0Report Type:Periodic  
Age:68 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536516A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK			Effexor	C		
				Glucovance	C		
				Lipitor	C		
				Lisinopril	C		
				Morphine	C		
				Nexium	C		

Date:02/17/05ISR Number: 4585019-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536531A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	4 DAY			Thyroid	C		
		Hot Flush					
		Insomnia					

Date:02/17/05ISR Number: 4585020-9Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536544A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Eructation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK						
		Retching					

Date:02/17/05ISR Number: 4585021-0Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536591A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hiccups		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK			No Concurrent			

Medication

C

Date:02/17/05ISR Number: 4585022-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536610A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anger		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585023-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536623A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 DAY			Effexor	C		
				Estrogen	C		

Date:02/17/05ISR Number: 4585024-6Report Type:Periodic  
Age:16 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536624A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
1TAB Per day	4 WK			Protonix	C		
				Melatonin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585025-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536639A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		
				Skin Burning Sensation			

Date:02/17/05ISR Number: 4585026-XReport Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536640A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Unknown Medication	C		
				Effexor	C		
				Muscle Twitching			

Date:02/17/05ISR Number: 4585027-1Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0536667A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other				Adderall	SS		
				Hypnagogic Hallucination			
				UNKNOWN			

Date:02/17/05ISR Number: 4585028-3Report Type:Periodic  
 Age:19 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0536692A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
Other				Grand Mal Convulsion			

Date:02/17/05ISR Number: 4585029-5Report Type:Periodic  
 Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536771A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

300MG Per day	2	MON	Nausea	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Panic Attack	Synthroid	C	Glaxosmithkline	
			Restlessness	Diazid	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585030-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0536773A  
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1	DAY	Agitation	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Disturbance In Attention	Vitamins	C		
			Dizziness	Synthroid	C	Glaxosmithkline	
			Dry Mouth	Tums	C	Glaxosmithkline	
			Nausea				

Date:02/17/05ISR Number: 4585031-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0536777A  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day			Dyspepsia	Wellbutrin	PS	Glaxosmithkline	ORAL
			Gastric Ulcer	Lipitor	C		
				Ibuprofen	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585032-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536778A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585033-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536782A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Jittery		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	9 MON	Insomnia		Effexor	C		
		Somnolence		Synthroid	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585034-9Report Type:Periodic  
Age:48 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0536786A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Discomfort		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day		Hot Flush		Buspar	C		
		Premature Menopause		Glucophage Xr	C		
				Avandia	C	Glaxosmithkline	
				Prevacid	C		

Date:02/17/05ISR Number: 4585035-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536789A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Affect Lability		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 YR			No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585036-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536793A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hypothalamo-Pituitary		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3	YR	Disorders					

Date:02/17/05ISR Number: 4585037-4Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536798A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	9 MON	Tinnitus		Propecia	C		
				Valium	C		
				Allegra	C		

Date:02/17/05ISR Number: 4585038-6Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0536842A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Risperdal	C		
				Lithium	C	Glaxosmithkline	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585039-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536937A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Psychotic Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585040-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0536959A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Strattera	C		
				Effexor Xr	C		

Date:02/17/05ISR Number: 4585041-6Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0537068A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hypertension		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 WK						

Date:02/17/05ISR Number: 4585042-8Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0537074A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dyspnoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Nausea		Doxycycline	SS		
		Palpitations					
		Vomiting					

Date:02/17/05ISR Number: 4585043-XReport Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0537106A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2	DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Anorexia					
		Insomnia		Lexapro	C		
		Nausea		Klonopin	C		
		Somnolence					

Date:02/17/05ISR Number: 4585044-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0537116A  
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1	DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Abdominal Pain					
		Anxiety		No Concurrent			
		Depression		Medication	C		
		Hypertension					
		Lethargy					
		Oropharyngeal Swelling					
		Pain					
		Pyrexia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585045-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0537125A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin Xl	PS	Glaxosmithkline	
6	MON						

Date:02/17/05ISR Number: 4585046-5Report Type:Periodic  
Age:17 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0537229A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Grand Mal Convulsion		Allergy Shots	C		
12	DAY	Incontinence		Keflex	C	Glaxosmithkline	
250MG	Four						
	times per day						

Date:02/17/05ISR Number: 4585047-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0537234A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gastrointestinal Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG	Per day						

Date:02/17/05ISR Number: 4585048-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0537242A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ear Discomfort		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG	Per day			Wellbutrin Sr	SS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585049-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0537250A  
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day				Blood Pressure Medication	C		

Date:02/17/05ISR Number: 4585050-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0537269A  
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 2 MON		Ill-Defined Disorder		Albuterol	C	Glaxosmithkline	
		Migraine		Entex	C		
		Weight Decreased		Serzone	C		
				Hrt	C		
				Flonase	C	Glaxosmithkline	
				Imitrex	C	Glaxosmithkline	
				Vicodin	C		

Date:02/17/05ISR Number: 4585051-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0537284A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Prozac	C		
				Antihistamines	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Percocet C  
Valium C

Date:02/17/05ISR Number: 4585052-0Report Type:Periodic  
Age:26 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0537286A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Somnolence		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Effexor	SS		ORAL

Date:02/17/05ISR Number: 4585053-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0537420A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Stomach Discomfort		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585054-4Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0537422A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 6 DAY				Benadryl	C	Glaxosmithkline	
UNKNOWN				Insulin	C		
				Estrogen Patch	C		
				Aspirin	C	Glaxosmithkline	
				Methyltestosterone	C		

Date:02/17/05ISR Number: 4585055-6Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0537432A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	300MG Per day 6 MON	Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Feeling Abnormal Tremor		Augmentin	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585056-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0537444A  
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day	Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
				Prozac	C		
				Clonopin	C		
				Seroquel	C		

Date:02/17/05ISR Number: 4585057-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0537467A  
 Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG Per day 3 WK	Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Palpitations		Synthroid	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585058-1Report Type:Periodic  
Age:26 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0537470A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	26 DAY						

Date:02/17/05ISR Number: 4585060-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0537471A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
WK		Nausea					

Date:02/17/05ISR Number: 4585061-1Report Type:Periodic  
Age:62 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0537595A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK	Agitation		Prozac	C		
		Insomnia		Crestor	C		
		Pollakiuria		Diovan	C		
				Nexium	C		
				Zantac	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585062-3Report Type:Periodic  
Age:35 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0537619A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3 MON	Insomnia		Lipitor	C		
		Weight Decreased		Avandia	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585063-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0537644A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anger		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585064-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0537758A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accidental Overdose		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Lexapro	C		

Date:02/17/05ISR Number: 4585065-9Report Type:Periodic  
Age:25 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0537762A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 10 DAY		Constipation Flatulence		No Concurrent Medication	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585067-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0537837A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Wellbutrin Sr	C	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585068-4Report Type:Periodic  
Age:39 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0537914A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Hospitalization - 36 DAY				Cymbalta	C		
Initial or Prolonged 1 WK				Remeron	C		
Disability				Zyprexa	C		

Date:02/17/05ISR Number: 4585069-6Report Type:Periodic  
Age: Gender:I/FU:F

Company Report #WLBADROO

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Vomiting					

Date:02/17/05ISR Number: 4585070-2Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538014A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 1 DAY				Paxil	SS	Glaxosmithkline	ORAL
1 DAY				Zetia	C		
				Ventolin Hfa	C	Glaxosmithkline	
				Prevacid	C		

Celebrex C  
Avandia C Glaxosmithkline  
Premarin C

Date:02/17/05ISR Number: 4585071-4Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538025A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK	Headache		Diamox	C		

Date:02/17/05ISR Number: 4585072-6Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538033A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Seroquel	C		
				Depakote	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585073-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538043A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	Dyskinesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hyperacusis		Klonopin	C		
				Trazodone	C		

Date:02/17/05ISR Number: 4585074-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538056A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585075-1Report Type:Periodic  
Age:66 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538235A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day 1 DAY	Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Casodex	C		
				Amiodarone	C		
				Lasix	C	Glaxosmithkline	
				Carvedilol	C	Glaxosmithkline	
				Lipitor	C		
				Synthroid	C	Glaxosmithkline	
				Magnesium Oxide	C		
				Plavix	C		
				Avapro	C		
				Vitamin B-12	C	Glaxosmithkline	
				Spiroinolactone	C		
				Zetia	C		
				Prozac	C		
				Glipizide	C		
				Megestrol	C		
				Isosorbide			
				Mononitrate	C		

Date:02/17/05ISR Number: 4585076-3Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0538241A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Depression					
		Dizziness					
		Irritability					

Date:02/17/05ISR Number: 4585077-5Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538242A

Outcome	PT
Hospitalization - Initial or Prolonged	Anxiety Blood Glucose Decreased Chest Pain Dry Mouth Dyspnoea

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Heart Rate Increased Muscular Weakness Nausea	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3 DAY	Nervousness		Wellbutrin	PS	Glaxosmithkline	ORAL
		Rash		Nexium	C		
		Thinking Abnormal		Estrace	C		
		Tinnitus					
		Weight Decreased					

Date:02/17/05ISR Number: 4585078-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0538301A  
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	9 DAY	Local Swelling		Yasmin	C		

Date:02/17/05ISR Number: 4585079-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0538312A  
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	Psychomotor Hyperactivity		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	1 YR			Lipitor	C		
				Xanax	C		

Date:02/17/05ISR Number: 4585080-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0538324A  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	450MG Per day	Vision Blurred		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	10 MON			Synthroid	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585082-9Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538498A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Heart Rate Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Insomnia		Sudafed	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585083-0Report Type:Periodic  
Age:27 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538509A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Affect Lability		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Drug Ineffective Overdose Tremor		Clomid	C		

Date:02/17/05ISR Number: 4585084-2Report Type:Periodic  
Age:71 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538511A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 YR	Anxiety Tinnitus		Altace Resperidol	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585085-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538524A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	1 WK	Nasopharyngitis	Wellbutrin	PS	Glaxosmithkline	ORAL
				Lisinopril	C		
				Depakote Er	C		
				Trazodone	C		

Date:02/17/05ISR Number: 4585086-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538543A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	1 DAY		Nausea	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585087-8Report Type:Periodic  
 Age:56 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538556A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	15 MON	Muscle Spasms	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Muscle Tightness	Effexor	C		
				Xanax	C		
				Atacand	C		

Date:02/17/05ISR Number: 4585088-XReport Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538581A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	1TAB Per day	11 DAY	Crying	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Dizziness				

Date:02/17/05ISR Number: 4585089-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0538677A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Electric Shock		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

WK

Date:02/17/05ISR Number: 4585090-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0538680A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Skin Odour Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

6 WK

Date:02/17/05ISR Number: 4585091-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0538684A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

150MG Per day

Dry Eye  
Dry Mouth



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585092-1Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0538685A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia		Wellbutrin Xr	PS	Glaxosmithkline	ORAL
150MG Twice		Nasopharyngeal Disorder					
per day	MON			Nexium	C		
				Magnesium	C		
				Calcium D	C		
1200MG Per							
day				Nasal Spray	C		NASAL

Date:02/17/05ISR Number: 4585093-3Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538807A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Night Sweats		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585094-5Report Type:Periodic  
Age:35 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0538809A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR	Stool Analysis Abnormal		Multivitamin	C		
				Wellbutrin Sr	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585095-7Report Type:Periodic  
Age:51 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538810A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Palpitations		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	6	MON		Lipitor	C		
				Accupril	C		
				Hydrochlorothiazide	C		
				Propecia	C		
				Nortriptyline	C		

Date:02/17/05ISR Number: 4585096-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0538817A  
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	9	DAY		Claritin	C		
		Headache		Premarin	C		
		Nausea					

Date:02/17/05ISR Number: 4585097-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0538818A  
 Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	5	WK		Ativan	C		
		Restlessness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585098-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538822A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	1TAB Per day	12 DAY	Off Label Use	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585099-4Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538829A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day		Diarrhoea	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Dizziness	Methotrexate	C		
			Nausea	Folic Acid	C		
			Paraesthesia	Plaquenil	C		
			Paraesthesia Oral	Naproxen	C		
			Vomiting				

Date:02/17/05ISR Number: 4585100-8Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0538843A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Three times per day		Medication Error	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585101-XReport Type:Periodic  
Age:70 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538849A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Per day 4 DAY Pharynx Discomfort

Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Glucophage	C		
Flomax	C		
Timolol Maleate	C		
Trusopt	C		
Coreg	C	Glaxosmithkline	
Hctz	C		
Neurontin	C		
Cozaar	C		
Zocor	C		
Lantus	C		
Novolog	C		
Plavix	C		
Robaxin	C		
Tylenol No. 3	C		
Lasix	C	Glaxosmithkline	
Ambien	C		

Date:02/17/05ISR Number: 4585103-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538850A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585104-5Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0538915A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585105-7Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539064A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Affect Lability		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	10 MON	Drug Ineffective		Glucophage	C		
		Sleep Disorder		Oral Contraceptives	C		

Date:02/17/05ISR Number: 4585106-9Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539070A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	6 DAY	Pharyngolaryngeal Pain		Synthroid	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585107-0Report Type:Periodic  
Age:70 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539075A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	6 WK	Pharyngolaryngeal Pain		Amiodarone	C		
				Hctz	C		
				Aspirin	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585108-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539076A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Sleep Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6	WK					

Date:02/17/05ISR Number: 4585109-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0539081A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Menstrual Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585110-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0539084A  
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6	MON					

Date:02/17/05ISR Number: 4585111-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0539097A  
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Feeling Jittery		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	5	WK		Lexapro	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585112-4Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539112A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 DAY						
				Imitrex	C	Glaxosmithkline	
				Synthroid	C	Glaxosmithkline	
				Nexium	C		
				Allegra	C		
				Zocor	C		
				Lexapro	C		
				Senokot	C		
				Multivitamin	C		
				Fibercon	C		
				Glucosamine	C		
				Ibuprofen	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585113-6Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539113A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day							
				Soma	C		
				Motrin	C	Glaxosmithkline	
				Vicodin	C		

Date:02/17/05ISR Number: 4585114-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539269A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
MON							

Date:02/17/05ISR Number: 4585115-XReport Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0539270A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1	WK	Depressed Mood		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Fatigue Somnolence					

Date:02/17/05ISR Number: 4585117-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0539271A  
 Age: Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2	DAY	Hypersensitivity		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Pruritus Generalised					

Date:02/17/05ISR Number: 4585118-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0539278A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3 WK	Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Insomnia		Ativan	C		
				Desyrel	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585119-7Report Type:Periodic  
 Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539283A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	12 DAY	Agitation Crying Panic Attack Stomatitis Vision Blurred		Wellbutrin Xl Synthroid Vitamins	PS C C	Glaxosmithkline Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585120-3Report Type:Periodic  
 Age:77 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539290A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
0 DAY		Stool Analysis Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2 DAY				Wellbutrin Sr	SS	Glaxosmithkline	ORAL
				Lexapro	C		

Date:02/17/05ISR Number: 4585121-5Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539312A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1TAB Per day	0 DAY	Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
				Lipitor	C		
				Nexium	C		
				Flonase	C	Glaxosmithkline	
				Allegra	C		

Date:02/17/05ISR Number: 4585122-7Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539464A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585123-9Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539465A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	7	MON		Prozac	C		

Date:02/17/05ISR Number: 4585124-0Report Type:Periodic  
Age:41 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0539468A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bruxism		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3	DAY		Imitrex	C	Glaxosmithkline	
		Tic					

Date:02/17/05ISR Number: 4585125-2Report Type:Periodic  
Age:75 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0539469A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
272	DAY						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585126-4Report Type:Periodic  
Age:38 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0539471A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585127-6Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539479A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4 WK	Dry Mouth		Ambien	C		
		Insomnia		Zelnorm	C		
		Restlessness					
		Weight Decreased					

Date:02/17/05ISR Number: 4585128-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539483A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysphemia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
4 DAY				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585129-XReport Type:Periodic  
Age:45 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539499A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 MON			Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Twice							
per day	1 WK			No Concurrent			

Date:02/17/05ISR Number: 4585130-6Report Type:Periodic  
 Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539520A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Somnolence		Wellbutrin Xl Thyroid Medication	PS C	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585131-8Report Type:Periodic  
 Age:58 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0539669A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK			Fosamax	C		

Date:02/17/05ISR Number: 4585132-XReport Type:Periodic  
 Age:25 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539672A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Heart Rate Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	6 DAY	Vision Blurred		Centrum	C		
				Vitamin C	C	Glaxosmithkline	
				Tylenol Cold + Flu	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585133-1Report Type:Periodic  
Age:52 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0539678A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2 MON		Rash		Black Cohosh	C	Glaxosmithkline	
UNKNOWN		Swelling Face Swollen Tongue Urticaria					

Date:02/17/05ISR Number: 4585134-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539684A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Heart Rate Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6 WK	Hypertension Insomnia		Wellbutrin Sr	SS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585135-5Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0539828A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Psychomotor Hyperactivity		Cynomel	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585136-7Report Type:Periodic  
Age:79 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0539906A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown	31 DAY	Rash Generalised		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585137-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0539907A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Initial Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Unknown		Therapeutic Response Unexpected					

Date:02/17/05ISR Number: 4585138-0Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0539913A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 6 MON		Headache					

Date:02/17/05ISR Number: 4585139-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540062A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Feeling Jittery		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hyperhidrosis		Effexor	C		
		Nervousness		Wellbutrin Sr	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585140-9Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540069A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 DAY			Ritalin	C		
				Lipitor	C		

Date:02/17/05ISR Number: 4585141-0Report Type:Periodic  
Age:34 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540081A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK	Pruritus					
		Rash					
		Swelling					

Date:02/17/05ISR Number: 4585142-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0540128A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:02/17/05ISR Number: 4585143-4Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0540132A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Therapeutic Response					
		Unexpected					

Date:02/17/05ISR Number: 4585144-6Report Type:Periodic  
Age:55 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540237A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Fatigue Middle Insomnia		Tylenol	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585145-8Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540240A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4 MON	Anxiety Headache Hot Flush Insomnia Tinnitus		Ortho Novum Allegra Prozac	C C C		

Date:02/17/05ISR Number: 4585146-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540249A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	
Other				No Concurrent			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Medication C

Date:02/17/05ISR Number: 4585147-1Report Type:Periodic  
Age:31 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0540258A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR		Breast Tenderness				
			Therapeutic Aspiration	Lexapro	C		

Date:02/17/05ISR Number: 4585148-3Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0540269A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Dizziness				

Date:02/17/05ISR Number: 4585149-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540280A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day			Adverse Event				

Date:02/17/05ISR Number: 4585150-1Report Type:Periodic  
Age:45 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0540324A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other			Disorientation				
			Memory Impairment				
			Petit Mal Epilepsy				

Date:02/17/05ISR Number: 4585151-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540431A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	5 WK	Cough		Wellbutrin	PS	Glaxosmithkline	
		Pharyngolaryngeal Pain Sinusitis		Oral Contraceptives	C		

Date:02/17/05ISR Number: 4585152-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0540452A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Oedema		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585153-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0540453A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585156-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540461A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Withdrawal Syndrome		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585157-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540484A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585158-6Report Type:Periodic  
Age:48 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540512A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY	Insomnia		Hytrin	C		
				Maxide	C		
				Procardia	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585159-8Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540536A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK	Dry Mouth		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585160-4Report Type:Periodic  
Age:20 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540538A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2	MON					

Date:02/17/05ISR Number: 4585161-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0540539A  
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hyperhidrosis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6	MON					
				Trazodone	C		
				Lamictal	C	Glaxosmithkline	
				Zoloft	C		

Date:02/17/05ISR Number: 4585162-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0540540A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3	MON					
				Synthroid	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585163-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540544A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day	MON	Tinnitus					

Date:02/17/05ISR Number: 4585164-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540549A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585165-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540550A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Eye Movement Disorder Muscle Twitching		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585166-5Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540551A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585167-7Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0540552A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin Xl Zyprexa	PS C	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585168-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540556A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Eructation		No Concurrent			
		Tinnitus		Medication	C		

Date:02/17/05ISR Number: 4585169-0Report Type:Periodic  
Age:41 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0540757A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	300MG Per day 2 DAY	Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged				No Concurrent			
Other				Medication	C		

Date:02/17/05ISR Number: 4585170-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540758A

Outcome	PT
	Agitation
	Dizziness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Feeling Abnormal Pollakiuria Pruritus	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	14 DAY			Nexium	C		
				Trazodone	C		

Date:02/17/05ISR Number: 4585171-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0540765A  
 Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hot Flush		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK			Xanax	C		ORAL

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Date:02/17/05ISR Number: 4585172-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0540770A  
 Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ear Discomfort Ear Pain Insomnia Tinnitus Vomiting		Wellbutrin Xl Sonata	PS C	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585173-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0540774A  
 Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 DAY			Effexor Xr	C		

Date:02/17/05ISR Number: 4585174-4Report Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540784A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Dry Mouth		Prozac	C		
		Insomnia		Librium	C		
		Tremor		Ambien	C		
				Lopressor	C		

Date:02/17/05ISR Number: 4585175-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540791A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:02/17/05ISR Number: 4585176-8Report Type:Periodic  
Age:25 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540800A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Stool Analysis Abnormal					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585177-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540801A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Medication Error		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Pharmaceutical Product Complaint		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585178-1Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0540951A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Pharmaceutical Product Complaint		Wellbutrin	PS	Glaxosmithkline	ORAL
		Stool Analysis Abnormal		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585179-3Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540954A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	4 MON	Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Heart Rate Increased Memory Impairment		Synthroid	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585180-XReport Type:Periodic  
Age:72 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541095A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	4 DAY	Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Nausea		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585181-1Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541100A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	4 DAY			Glucophage	C		
				Dynacirc	C	Glaxosmithkline	
				Diovan	C		

Date:02/17/05ISR Number: 4585182-3Report Type:Periodic  
Age:45 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0541106A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	0 DAY						

Date:02/17/05ISR Number: 4585183-5Report Type:Periodic  
Age:48 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0541107A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	4 DAY	Drug Ineffective					
		Rash					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585184-7Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541133A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6 WK	Dizziness					
		Fatigue		Trazodone	C		

Date:02/17/05ISR Number: 4585185-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541146A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion		Xanax	C		

Date:02/17/05ISR Number: 4585186-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541147A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585187-2Report Type:Periodic  
Age:24 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541155A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Life-Threatening		Convulsion					
300MG Per day	239 DAY			Prozac	SS		
Other							
UNKNOWN	40MG Per day						

Date:02/17/05ISR Number: 4585188-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541314A

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:02/17/05ISR Number: 4585189-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0541316A  
Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
		Formication		Wellbutrin Xl No Concurrent Medication	PS  C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585190-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0541317A  
Age:41 YR Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
150MG Per day	9 DAY	Rash		Wellbutrin Xl  No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585191-4Report Type:Periodic  
Age:27 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541318A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585192-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541374A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585193-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541382A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased		Wellbutrin Xl Strattera	PS SS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585194-XReport Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0541395A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585195-1Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541533A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

300MG Per day 3 MON Panic Attack  
Therapeutic Response  
Decreased

Wellbutrin Xl PS Glaxosmithkline ORAL

Date:02/17/05ISR Number: 4585196-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0541539A  
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
1	MON			Effexor	C		

Date:02/17/05ISR Number: 4585197-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0541541A  
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	10 DAY			No Concurrent Medication	C		



Sleep Disorder

Date:02/17/05ISR Number: 4585202-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541603A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Liver Function Test Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585203-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541614A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
15 WK		Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585204-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541636A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Ill-Defined Disorder Pain In Extremity		Wellbutrin Xl	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585206-3Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541713A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	6 DAY	Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Insomnia		Ambien	C		
				Zelnorm	C		
				Lexapro	C		

Date:02/17/05ISR Number: 4585207-5Report Type:Periodic  
Age:64 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541719A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Stool Analysis Abnormal		Beta-Blocker	C		
				Pravachol	C		
				Folic Acid	C		
				Aspirin	C	Glaxosmithkline	

Date:02/17/05ISR Number: 4585208-7Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541731A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	35 DAY	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Mood Altered					

Date:02/17/05ISR Number: 4585209-9Report Type:Periodic  
Age:42 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541750A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dry Mouth		Lithobid	C	Glaxosmithkline	
		Hangover		Topomax	C		

Hypotension  
Nausea

Seroquel  
Klonopin

C  
C

Date:02/17/05ISR Number: 4585210-5Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541751A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585211-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541759A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bruxism		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300U Per day							

Date:02/17/05ISR Number: 4585212-9Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541764A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585213-0Report Type:Periodic  
Age:29 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541940A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 WK		Influenza Like Illness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Pyrexia		Prozac	C		
		Rash		Propanol	C		

Date:02/17/05ISR Number: 4585214-2Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541953A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Hormone Replacement	C		

Date:02/17/05ISR Number: 4585215-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541962A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown	2 MON	Erythema Nodosum		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585216-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541963A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anaphylactic Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585217-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541975A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hypertension

Wellbutrin Xl

PS

Glaxosmithkline

Date:02/17/05ISR Number: 4585218-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542159A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	7 DAY			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Heart Rate Increased					
		Insomnia		Blood Pressure Med	C		
		Paranoia		Claritin	C		

Date:02/17/05ISR Number: 4585219-1Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542160A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3 MON			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Drug Ineffective					
		Muscle Twitching		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585220-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542173A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
500MG Per day	1 WK			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Photosensitivity Reaction					
				Urine Incontinence			



Mood Altered

Wellbutrin Xl

PS

Glaxosmithkline

ORAL

Date:02/17/05ISR Number: 4585225-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542203A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day		Cardiac Flutter					
		Tachycardia					

Date:02/17/05ISR Number: 4585226-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542206A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Insomnia		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585227-0Report Type:Periodic  
Age:22 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542425A

Outcome	PT
	Agitation
	Energy Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Insomnia Somnolence Tinnitus	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6 WK			No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585228-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0542430A  
 Age: Gender: I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							
300MG Per day							

Date:02/17/05ISR Number: 4585229-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0542432A  
 Age:68 YR Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Dry Mouth		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK			No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585230-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0542439A  
 Age: Gender:Male I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Suicidal Ideation		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585231-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0542440A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Suicidal Ideation		Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585232-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0542445A  
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1 WK	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
UNKNOWN		Anxiety		Unknown Medication	C		
		Chest Discomfort					
		Sleep Disorder					
		Tinnitus					

Date:02/17/05ISR Number: 4585233-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0542451A  
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	4 WK	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dry Mouth		No Concurrent Medication	C		
		Nausea					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585234-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542456A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	4 WK		Wellbutrin	PS	Glaxosmithkline	ORAL
		Alopecia		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585235-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542464A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Myalgia					

Date:02/17/05ISR Number: 4585236-1Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0542478A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other		Hallucination, Auditory					

Date:02/17/05ISR Number: 4585237-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542483A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Alopecia Hypotrichosis					

Date:02/17/05ISR Number: 4585238-5Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0542632A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization -	Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
3 MON					
Initial or Prolonged		No Concurrent			
Other		Medication	C		

Date:02/17/05ISR Number: 4585239-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0542636A  
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300G Per day	8 DAY			No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585240-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0542639A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	7 YR	Rash		Metformin	C		
				Diclofenac Sodium	C	Glaxosmithkline	
				Crestor	C		
				Premarin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585241-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542643A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	1 WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Tachycardia		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585242-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542644A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
Other		Hallucination		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585243-9Report Type:Periodic  
 Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542650A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	2 DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Insomnia		Birth Control Pills	C		

Date:02/17/05ISR Number: 4585244-0Report Type:Periodic  
 Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542660A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	2 MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Nervousness		Wellbutrin Sr	SS	Glaxosmithkline	
		Pain In Extremity		Klonopin	C		

Date:02/17/05ISR Number: 4585245-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0542711A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585246-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0542828A  
Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585247-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0542836A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Stool Analysis Abnormal		Aciphex	C		
				Atenolol	C		
				Dilaudid	C		
				Lasix	C	Glaxosmithkline	
				Ativan	C		
				Paregoric	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585249-XReport Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542842A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	4	MON	Memory Impairment	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Myalgia	Effexor	C		
			Psychomotor Hyperactivity				

Date:02/17/05ISR Number: 4585250-6Report Type:Periodic  
Age:64 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542848A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	6	MON	Abdominal Distension	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Abdominal Pain Upper	Diovan	C		
				Levoxyl	C	Glaxosmithkline	
				Allegra	C		
				Singulair	C		

Date:02/17/05ISR Number: 4585251-8Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0542851A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Oedema	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585252-XReport Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542853A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	14	MON	Anorexia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Pharmaceutical Product Complaint				
			Weight Decreased				

Date:02/17/05ISR Number: 4585253-1Report Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542860A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypertension		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585254-3Report Type:Periodic  
Age:63 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0542862A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4585255-5Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0542863A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585256-7Report Type:Periodic  
Age:57 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543051A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	2 DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dyspnoea		No Concurrent Medication	C		

Date:02/17/05ISR Number: 4585257-9Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543053A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Tinnitus					

Date:02/17/05ISR Number: 4585258-0Report Type:Periodic  
Age:57 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0543066A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Unknown	81 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion		Neurontin	C		

Date:02/17/05ISR Number: 4585259-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543140A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	2 WK			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hypersensitivity					
		Urticaria					

Date:02/17/05ISR Number: 4585260-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543160A

Outcome Dose Duration PT Report Source Product Role Manufacturer Route  
Drug Ineffective Wellbutrin Xl PS Glaxosmithkline ORAL  
300MG Per day

Date:02/17/05ISR Number: 4585261-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0543378A  
Age: Gender: I/FU:I

Outcome Dose Duration PT Report Source Product Role Manufacturer Route  
Feeling Jittery Wellbutrin Xl PS Glaxosmithkline ORAL  
300MG Per day

Date:02/17/05ISR Number: 4585262-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0543779A  
Age:51 YR Gender:Female I/FU:F

Outcome Dose Duration PT Report Source Product Role Manufacturer Route  
Other Convulsion Wellbutrin Xl PS Glaxosmithkline ORAL  
300MG Per day 15 DAY  
75MG Per day Effexor Xr C ORAL  
40MG As Lasix C Glaxosmithkline ORAL  
required

Date:02/17/05ISR Number: 4585855-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544770A  
Age: Gender: I/FU:I

Outcome Dose Duration PT Report Source Product Role Manufacturer Route  
Hospitalization - Overdose Wellbutrin PS Glaxosmithkline ORAL  
Initial or Prolonged



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4585863-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0276896A

Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day 3 DAY	Acute Myocardial		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization -	1TAB Per day	Infarction		Cilest	C		ORAL
Initial or Prolonged		Cardiac Arrest					
Other		Cardioversion					
		Chest Pain					
		Coronary Artery					
		Dissection					
		Headache					

Date:02/17/05ISR Number: 4585872-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0365278A

Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day 5 DAY	Muscle Spasms		Zyban	PS	Glaxosmithkline	ORAL

Date:02/17/05ISR Number: 4586824-9Report Type:Direct Company Report #CTU 240701

Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150 MG 4	Asthenia		Zyban 150 Mg	PS		ORAL
		Dyskinesia					

DAYS ORAL;

300 MG

DURATION

ORAL

Triamterene	C
Inderol	C
Tarka T-240	C
Volmax	C

Date:02/17/05ISR Number: 4589250-1Report Type:Direct  
 Age:43 YR Gender:Female I/FU:I

Company Report #CTU 240714

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG /DAILY		Abasia		Welbutrin Xl 300 Mg	PS	Gsk	ORAL
Initial or Prolonged PO [ STARTED Required 150 ON 1/29		Confusional State					
Intervention to INCR 300 MG		Hypertension					
Prevent Permanent ON 2/2/05] Impairment/Damage		Vision Blurred					

Date:02/17/05ISR Number: 4590060-XReport Type:Direct  
 Age:74 YR Gender:Male I/FU:I

Company Report #CTU 240752

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 1 TAB QD		Agitation		Wellbutrin Xl 300 Mg	PS		
		Completed Suicide		Glucovance	C		
		Confusional State		Lotensin	C		
		Gun Shot Wound					
		Hallucination					
		Visual Disturbance					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4604975-7Report Type:Periodic  
Age:70 YR Gender:Female I/FU:F

Company Report #S04-USA-06882-01

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Health	Memantine	PS		ORAL
10 MG QD PO			Professional	Memantine	SS		ORAL
5 MG QD PO				Memantine	SS		ORAL
10 MG QD PO				Memantine	SS		ORAL
15 MG QD PO				Wellbutrin (Bupropion Hydrochloride)	SS		
150 MG QD				Wellbutrin (Bupropion Hydrochloride)	SS		
300 MG QD				Aricept (Donepezil Hydrochloride)	C		
				Claritin (Loratadine)	C		
				Estradiol	C		
				Tylenol (Paracetamol)	C		
				Aspirin	C		
				Mirapex	C		
				Zinc	C		
				Multivitamin	C		

Date:02/18/05ISR Number: 4587212-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0545451A  
Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
		Amnesia		Zyprexa	C		
		Anxiety		Depakote Er	C		
		Constipation					
		Headache					
		Irritability					
		Mania					

Muscle Twitching  
Nervousness  
Somnolence

Date:02/18/05ISR Number: 4587221-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0545970A  
Age:26 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abnormal Behaviour Completed Suicide		Wellbutrin Xl Adderall	PS C	Glaxosmithkline	ORAL

Date:02/18/05ISR Number: 4587247-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0371030A  
Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG		Arthralgia Musculoskeletal Stiffness		Zyban	PS	Glaxosmithkline	ORAL
Variable dose 1	MON			Humalog	C		
UNKNOWN				Humulin	C		
UNKNOWN							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/21/05ISR Number: 4588192-5Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0370764A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Pruritus		Zyban	PS	Glaxosmithkline	
UNKNOWN		Sleep Disorder					
		Tooth Disorder					
		Toothache					

Date:02/22/05ISR Number: 4588841-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541125A

Age:83 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Medication Error					
per day	2 WK	Paranoia		Lithium	C	Glaxosmithkline	
		Pharmaceutical Product		Ativan	C		
		Complaint					

Date:02/22/05ISR Number: 4588845-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0545918A

Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
8 DAY		Dizziness		Delsym	SS	Glaxosmithkline	
		Dysgeusia		Allegra	C		
		Hot Flush		Promethazine Dm	C		
		Loss Of Consciousness		Robitussin Dm	C		
		Muscle Twitching		Levothyroxine	C	Glaxosmithkline	
		Nausea		Z Pack	C		
		Tremor		Biaxin	C		

Date:02/22/05ISR Number: 4588846-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0545927A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Adverse Drug Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day		Constipation		Wellbutrin	SS	Glaxosmithkline	
150MG Per day		Diplopia		Duragesic	C		
		Dry Mouth		Methadone	C	Glaxosmithkline	
		Hallucination, Auditory		Effexor	C		
		Hallucination, Visual		Valium	C		
		Pruritus		Vicodin	C		
		Somnolence		Neurontin	C		
		Vision Blurred		Avandia	C	Glaxosmithkline	

Date:02/22/05ISR Number: 4588847-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0545955A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Hallucination		No Concurrent			
		Hypertension		Medication	C		
		Overdose					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/22/05ISR Number: 4588849-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0546103A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Bupropion	PS	Glaxosmithkline	
		Drug Exposure During Pregnancy					
		Pregnancy					

Date:02/22/05ISR Number: 4588857-5Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0362396A

Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Accidental Overdose		Zyban	PS	Glaxosmithkline	
UNKNOWN	150MG	Single					
dose		Dizziness					
		Hyperventilation					
		Nausea					
		Retching					

Date:02/22/05ISR Number: 4588868-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0371935A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Pressure Decreased		Zyban	PS	Glaxosmithkline	
Initial or Prolonged		Difficulty In Walking		Ventolin	C	Glaxosmithkline	
		Hypertension		Seretide	C	Glaxosmithkline	
		Lung Infection		Atrovent	C	Glaxosmithkline	
		Tremor		Acimax	C	Glaxosmithkline	
				Zanidip	C		
				Losec	C	Glaxosmithkline	

Date:02/22/05ISR Number: 4591875-4Report Type:Expedited (15-DaCompany Report #HQWYE199809FEB05

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Convulsion	Health	Effexor (Venlafaxine		
	Drug Interaction	Professional	Hydrochloride, )	PS	
	Photopsia		Wellbutrin Xl		
			(Bupropion		
			Hydrochloride,)	SS	ORAL

SEE IMAGE

Date:02/22/05ISR Number: 4592479-XReport Type:Expedited (15-DaCompany Report #2005-01-1690

Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Haemorrhagic Stroke Hypertension	Health Professional Company	Peg-Intron (Peginterferon Alfa-2b) Injectable	PS		
SUBCUTANEOUS	150 MCG QWK		Representative				
SUBCUTANEOUS				Copegus (Ribavirin) Capsules	SS		ORAL
1200 MG QD							
ORAL				Pegasys (Pegylated Interferon Alfa-2a) Injectable	SS		
SUBCUTANEOUS	SUBCUTANEOUS			Wellbutrin (Bupropion) Percocet	SS C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Benzodiazepine(Nos) C

Date:02/23/05ISR Number: 4589781-4Report Type:Expedited (15-DaCompany Report #BPC-ZN-05-048  
Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day Initial or Prolonged Disability			Blood Cholesterol Increased Confusional State Diabetes Mellitus Non-Insulin-Dependent Foaming At Mouth Grand Mal Convulsion Hyperglycaemia Hypertension Impaired Driving Ability Urinary Incontinence	Zyban	PS	Glaxosmithkline	ORAL

Date:02/23/05ISR Number: 4589786-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0546075A  
Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Blood Caffeine Increased Convulsion Fatigue Sleep Disorder Weight Decreased	Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:02/23/05ISR Number: 4589787-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0546080A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Life-Threatening Other			Death Medication Error	Wellbutrin Wellbutrin No Concurrent Medication	PS SS C	Glaxosmithkline	ORAL ORAL

Date:02/23/05ISR Number: 4589788-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0546307A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Apnoea		Effexor	C		
150MG Per day				Lipitor	C		

Date:02/23/05ISR Number: 4589806-6Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0371611A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	
Other		Haemorrhage					
10 DAY		Night Sweats					
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/23/05ISR Number: 4591382-9Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #CTU 241243

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression		Generic Wellbutrin	PS		
400 MG A DAY		Pharmaceutical Product Complaint Suicidal Ideation Therapy Non-Responder					

Date:02/23/05ISR Number: 4595153-9Report Type:Expedited (15-DaCompany Report #2004103296  
 Age:73 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Health	Zoloft (Sertraline)	PS		ORAL
50 MG (50 MG, 1 IN 1 D), ORAL		Cataract Condition Aggravated Depression Drug Ineffective Drug Intolerance Formication Vision Blurred	Professional	Bupropion Hydrochloride (Bupropion Hydrochloride) Paroxetine Hydrochloride (Paroxetine Hydrochloride) Psycholeptics (Psycholeptics) Provella-14 (Estrogens Conjugated, Medroxyprogesterone Acetate0 Ciclosporin (Ciclosporin) All Other Therapeutic Products (All Other Therapeutic Products)	SS  SS C  C C		

Date:02/24/05ISR Number: 4590559-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0540971A  
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Grand Mal Convulsion	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG In the							
morning	224 DAY		Memory Impairment				
			Speech Disorder				
			Tongue Biting				

Date:02/24/05ISR Number: 4590561-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0543779A  
Age:51 YR Gender:Female I/FU:F

Outcome	PT
Other	Convulsion
	Eye Injury
	Face Injury
	Fall
	Incontinence
	Limb Injury

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tongue Biting

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
19 DAY			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
75MG Per day			Effexor Xr	C		ORAL
40MG As			Lasix	C	Glaxosmithkline	ORAL
required						

Date:02/24/05ISR Number: 4590567-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0546813A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Cerebrovascular Accident	Wellbutrin	PS	Glaxosmithkline	ORAL
75MG Per day							

Date:02/24/05ISR Number: 4590586-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0372275A  
 Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening			Psychiatric Symptom	Zyban	PS	Glaxosmithkline	ORAL
Disability			Self-Injurious Ideation	Tylex	C		ORAL
Other							

Date:02/24/05ISR Number: 4590587-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0372277A  
 Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Alcohol Use	Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	10 DAY		Balance Disorder	Cannabis	C		
UNKNOWN			Fall	Alcohol	C		
UNKNOWN							

Gait Disturbance

Date:02/24/05ISR Number: 4591195-8Report Type:Direct Company Report #CTU 241354  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Bupropion Sr 15	PS		ORAL
150 MG PO BID		Pharmaceutical Product Complaint					

Date:02/25/05ISR Number: 4592042-0Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0547191A  
 Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice per day	WK	Anxiety					

Date:02/28/05ISR Number: 4592921-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0547245A  
 Age: Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Haemorrhage		Wellbutrin XL	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/05ISR Number: 4592948-2Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0372872A  
 Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Bupropion			
150MG See		Cognitive Disorder		Hydrochloride	PS	Glaxosmithkline	
		Hallucination, Visual					
dosage text	12 DAY	Panic Attack		No Concurrent			
		Suicidal Ideation		Medication	C		
		Tremor					

Date:02/28/05ISR Number: 4596202-4Report Type:Direct Company Report #CTU 241505  
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Asthenia		Zoloft 50mg 1 Po Qd	PS		ORAL
50MG QDAY							
Initial or Prolonged		Fatigue		Bupropion 150 Mg 1			
150 MG QDAILY		Tremor		Po Qd	SS		ORAL

Date:03/01/05ISR Number: 4594134-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0399790A  
 Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Anger		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
		Intentional Misuse					
per day	2 YR						

Date:03/01/05ISR Number: 4594139-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0546331A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Abortion Spontaneous Wellbutrin PS Glaxosmithkline ORAL  
300MG Per day 16 MON

No Concurrent Medication C

Date:03/01/05ISR Number: 4594168-4Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0357968A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other				Zyban	PS	Glaxosmithkline	ORAL
150MG See							
dosage text	48	DAY					
			Emotional Disorder				
			Haematoma	Alcohol	SS		
			Retrograde Amnesia	No Concurrent Medication	C		

Date:03/01/05ISR Number: 4594169-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0361828A  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability				Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	10	DAY					
			Depression				
			Feeling Of Despair	Haldol Decanoate	C		
INTRAMUSCULAR	100MG	Every					
two weeks		YR	Restlessness				
			Tearfulness	Artane	C		
5MG Three							
times per day		YR					
UNKNOWN	400MG	Twice		Tegretol	C		
per day		YR					
				Lexapro	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/05ISR Number: 4609105-3Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #20040200191

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Other	Reglan (Metoclopramide) (Baxter)	PS	Baxter	
10 MG DAILY							
				Wellbutrin Xl (Bupropion Hcl) Tablets Glaxo Smith Kline	SS	Glaxo Smith Kline	ORAL
300 MG DAILY							
PO	3	WK					

Date:03/02/05ISR Number: 4596699-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544022A  
 Age:13 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Convulsion Dehydration Electrolyte Imbalance Fall Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:03/02/05ISR Number: 4596701-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0545918A  
 Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
8 DAY							
		Burning Sensation		Delsym	SS	Glaxosmithkline	
		Dizziness		Allegra	C		
		Dysgeusia		Promethazine Dm	C		
		Hot Flush		Robitussin Dm	C		
		Loss Of Consciousness		Levothyroxine	C	Glaxosmithkline	
		Muscle Twitching		Z Pack	C		
		Nausea		Biaxin	C		

Tremor

Date:03/02/05ISR Number: 4596703-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0546744A  
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Rash		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	2 WK	Respiratory Disorder		Lotrel	C		

Date:03/02/05ISR Number: 4596704-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0547176A  
Age:13 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Breast Mass		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/05ISR Number: 4598310-0Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #CTU 241942

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
/5-6 WEEKS	5 WK			Zyban	PS		
		Complaint					
		Suicidal Ideation					

Date:03/02/05ISR Number: 4598857-7Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #CTU 241920

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150 BID ORAL				Bupropion 150	PS		ORAL
		Burning Sensation					
		Rash Macular					

Date:03/02/05ISR Number: 4598955-8Report Type:Expedited (15-DaCompany Report #A0546787A  
 Age:50 YR Gender:Male I/FU:I

Company Report #A0546787A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required							
Intervention to							
Prevent Permanent							
Impairment/Damage							
PER DAY /							
ORAL							
		Blood Creatinine	Study	Wellbutrin Xl			
		Increased	Health	Tablet-Extended			
		Grand Mal Convulsion	Professional	Release (Bupropion			
		Postictal State		Hydrochloride)	PS		ORAL
		Refusal Of Treatment By					
		Patient					

Date:03/03/05ISR Number: 4599299-0Report Type:Direct  
 Age: Gender:Female I/FU:I

Company Report #CTU 242000

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening							
100MG QD				Wellbutrin	PS	Gsk	ORAL
		Anaemia					

Required	Duodenal Ulcer			
ORAL				
Intervention to	Haemorrhage	Excedrin	SS	ORAL
TAB Q 4 HRS				
Prevent Permanent	Gastric Ulcer Haemorrhage			
ORAL				
Impairment/Damage	Hypotension			

Date:03/03/05ISR Number: 4599589-1Report Type:Expedited (15-DaCompany Report #SUSI-2005-00116  
Age:11 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Coma Fatigue	Other	Adderall Xr (Dextroamphetamine Sulfate, Dextroamphetamine Saccharate,	PS		
15 MG,							
1X/DAY:QD				Wellbutrin-Slow Release (Bupropion Hydrochloride)	SS		
300 MG/DAY;							
SEE IMAGE				Strattera (Atomoxetine Hydrochloride)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/04/05ISR Number: 4598249-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541138A  
 Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300MG Per day	3	WK	Joint Stiffness	Wellbutrin	PS	Glaxosmithkline	ORAL
			Musculoskeletal Stiffness	Lipitor	C		
			Oedema Peripheral	Aspirin	C	Glaxosmithkline	
			Rash				
			Serum Sickness				

Date:03/04/05ISR Number: 4598250-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541141A  
 Age:24 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300MG Per day	3	WK	Erythema	Wellbutrin	PS	Glaxosmithkline	ORAL
			Musculoskeletal Stiffness	Micronor	C		
			Oedema Peripheral				
			Orbital Oedema				
			Rash				
			Serum Sickness				
			Urticaria				

Date:03/04/05ISR Number: 4598251-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544127A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
UNKNOWN			Amnesia	Bupropion	PS	Glaxosmithkline	ORAL
			Convulsion	Ultram	C		
			Road Traffic Accident				

Date:03/04/05ISR Number: 4598252-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0546744A  
 Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 150MG Twice	Blood Pressure Increased	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day           2    WK	Heart Rate Increased				
	Hypersensitivity	Lotrel	C		
	Rash	Levaquin	C		
	Respiratory Disorder				
	Respiratory Rate Increased				
	Urticaria				

Date:03/04/05ISR Number: 4598732-8Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0357955A  
Age:44 YR   Gender:Male           I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dry Mouth		Zyban	PS	Glaxosmithkline	ORAL
150MG See							
dosage text	22	DAY					
		Ecchymosis					
		Haematoma		Micardis	C	Glaxosmithkline	ORAL
		Psoriasis		Emconcor	C		ORAL
				Lipitor	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/04/05ISR Number: 4598749-3Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLIN-B0373608A  
 Age:50 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - UNKNOWN	Duration Dyspnoea		Zyban	PS	Glaxosmithkline	
Initial or Prolonged UNKNOWN	Respiratory Disorder		Nitrofurantoin	C		
UNKNOWN			Sodium Rabeprazole	C		
UNKNOWN			Oestradiol	C		
UNKNOWN			Doxycycline	C		
UNKNOWN			Prednisolone	C	Glaxosmithkline	
UNKNOWN			Ipratropium Bromide	C	Glaxosmithkline	
UNKNOWN			Salbutamol Sulphate	C	Glaxosmithkline	

Date:03/04/05ISR Number: 4599655-0Report Type:Direct Company Report #CTU 242055  
 Age:58 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other DAILY ORAL	Duration Dyspnoea		Wellbutrin Sr 100 Mg	PS		ORAL
	Medication Error					

Date:03/04/05ISR Number: 4599818-4Report Type:Direct Company Report #CTU 242032  
 Age:38 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1 DOSE ONLY	Duration Blood Pressure Increased		Bupropion 75 Mg	PS		
	Headache		Metoprolol	C		
	Waxy Flexibility		Pantoprazole	C		
			Vitamins	C		

Date:03/04/05ISR Number: 4599974-8Report Type:Direct Company Report #CTU 242104  
Age:17 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin 150 Mg Gsk	PS	Gsk	ORAL
150 MG 2X							
DAILY ORAL							

Date:03/04/05ISR Number: 4600473-5Report Type:Expedited (15-DaCompany Report #2002133101US  
Age:34 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health	Diphenhydramine (Diphenhydramine)	PS		ORAL
ORAL							
Professional							
Isotretinoin (Isotretinoin)							
Bupropion (Bupropion)							
SS							
SS							

Date:03/04/05ISR Number: 4601322-1Report Type:Expedited (15-DaCompany Report #2002133124US  
Age:33 YR Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Multiple Drug Overdose	Literature Health	Diphenhydramine (Diphenhydramine)	PS		ORAL
ORAL							
Professional							
Risperidone (Risperidone)							
SS							
SS							
UNSPECIFIED							

AMT, ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

UNSPECIFIED  
 Bupropion (Bupropion) SS ORAL

AMT, ORAL

Date:03/07/05ISR Number: 4600276-1Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0046030A  
 Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	150MG Per day	Cardiac Failure		Zyban	PS	Glaxosmithkline	ORAL
	10MG Per day	Supraventricular		Simvastatin	C		ORAL
		Tachycardia					

Date:03/08/05ISR Number: 4601602-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0540261A  
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	450MG Per day	Dysgeusia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	200MG Per day	Mania		Lamictal	C	Glaxosmithkline	ORAL

Date:03/08/05ISR Number: 4601605-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0543066A  
 Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Unknown 81 DAY	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
		Incontinence		Neurontin	C		
		Postictal State					

Date:03/08/05ISR Number: 4601606-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0543494A  
 Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3	Contusion		Macrobid	C		
	WK	Hypersensitivity		Zyrtec	C	Glaxosmithkline	
		Hypotension					
		Rash Generalised					
		Rash Macular					
		Skin Discolouration					
		Swelling					
		Urinary Tract Infection					
		Urticaria					

Date:03/08/05ISR Number: 4604956-3Report Type:Direct  
Age:36 YR Gender:Male I/FU:I

Company Report #CTU 242383

Outcome	PT
Hospitalization -	Accidental Overdose
Initial or Prolonged	Anorexia
	Atonic Seizures
	Clonic Convulsion
	Confusional State
	Depression
	Dizziness
	Dysarthria
	Hallucination
	Headache

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Insomnia				
		Tremor				
		Vision Blurred	Report Source	Product	Role	Manufacturer
Dose	Duration			Wellbutrin	PS	Route
900 MG PO BID	2 WK					ORAL

Date:03/09/05ISR Number: 4602858-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0545928A  
 Age:42 YR Gender:Male I/FU:F

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose		Eye Swelling		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
Other		Pharyngolaryngeal Pain					
150MG Twice							
per day	1 MON						
		Rash		Zocor	C		ORAL
20MG Per day							

Date:03/09/05ISR Number: 4602864-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0548151A  
 Age: Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose		Asthma		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
Other		Dyspnoea		Advair	C	Glaxosmithkline	
100MG Unknown		Pulmonary Oedema					
RESPIRATORY							
(INHALATION)							

Date:03/09/05ISR Number: 4602879-7Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0369547A  
 Age:32 YR Gender:Female I/FU:F

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration						
Dose		Medication Error		Zyban	PS	Glaxosmithkline	
Other		Urticaria					
UNKNOWN	150MG Twice						
per day	2 WK						

Date:03/09/05ISR Number: 4602886-4Report Type:Expedited (15-DaCompany Report #FI-GLAXOSMITHKLINE-B0372907A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Consciousness Muscle Spasms		Zyban	PS	Glaxosmithkline	

Date:03/09/05ISR Number: 4602903-1Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0373985A  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Zyban	PS	Glaxosmithkline	
6 WK		Dyspnoea Hallucination Incoherent Insomnia Pyrexia Sensory Disturbance Tonsillitis					

Date:03/09/05ISR Number: 4604076-8Report Type:Direct Company Report #CTU 242399  
Age:32 YR Gender:Female I/FU:I

Outcome	PT
Disability	Depression
Other	Drug Ineffective Pharmaceutical Product

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Complaint  
Sedation

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
BID			Bupropion	PS		

Date:03/10/05ISR Number: 4603926-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0548205A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day Initial or Prolonged			Grand Mal Convulsion Medication Error Overdose	Wellbutrin Lexapro	PS C	Glaxosmithkline	ORAL

Date:03/10/05ISR Number: 4603927-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0548219A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Death Overdose Suicide Attempt	Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:03/10/05ISR Number: 4603929-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0548226A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening			Convulsion Overdose Rhabdomyolysis Suicide Attempt	Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:03/10/05ISR Number: 4603931-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0548370A  
Age:16 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300MG Per day	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Life-Threatening Hospitalization - Initial or Prolonged Other		Death Overdose Respiratory Arrest Vomiting		No Concurrent Medication	C		

Date:03/10/05ISR Number: 4603953-1Report Type:Expedited (15-DaCompany Report #FI-GLAXOSMITHKLINE-B0373882A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN	150MG	Blood Pressure Increased See		Zyban	PS	Glaxosmithkline	
Initial or Prolonged dosage text		Heart Rate Increased Respiratory Disorder		Unknown Medication	C		
UNKNOWN							

Date:03/10/05ISR Number: 4607618-1Report Type:Expedited (15-DaCompany Report #2005038205  
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Toxic Epidermal Necrolysis	Foreign Health Professional	Nifedidor Capsule (Nifedipine) Acetylsalicylic Acid	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Acetylsalicylic Acid)	SS
Furosemid Heumann (Furosemide)	SS
Enalapril Heumann (Enalapril Maleate)	SS
Carvedilol (Carvedilol)	SS
Digoxin (Digoxin)	SS
Canrenoic Acid (Canrenoic Acid)	SS
Allopurinol (Allopurinol)	SS
Bupropion (Bupropion)	SS
Technetium Tc 99m Pentetate (Technetium Tc 99m Pentetate)	SS

Date:03/11/05ISR Number: 4605917-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0516229A  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Exposure During Pregnancy Placental Insufficiency Pre-Eclampsia Pregnancy		Wellbutrin Effexor	PS C	Glaxosmithkline	ORAL

Date:03/11/05ISR Number: 4605952-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0548178A  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 2 WK		Convulsion  Fall Loss Of Consciousness Tongue Biting Tremor		Wellbutrin  Oral Contraceptive	PS C	Glaxosmithkline	ORAL

Date:03/11/05ISR Number: 4605953-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0548375A  
Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK	Drug Exposure During Pregnancy					

Date:03/11/05ISR Number: 4605954-6Report Type:Expedited (15-DaCompany Report #US-0501110039  
Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction		Wellbutrin	PS	Glaxosmithkline	
		Grand Mal Convulsion		Prozac	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/11/05ISR Number: 4605956-XReport Type:Expedited (15-DaCompany Report #2005-03-0065  
Age:46 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Collapse Of Lung		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged	Convulsion		Peg-Intron	SS		
SUBCUTANEOUS						
	Pneumonia		Ribavirin	SS		ORAL
25MG Single			Paxil Cr	C	Glaxosmithkline	ORAL
dose						
.05MG Unknown			Levoxyl	C	Glaxosmithkline	ORAL
30MG Unknown			Prevacid	C		ORAL

Date:03/11/05ISR Number: 4605957-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0548830A  
Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice						
per day	Medication Error					
3 YR			Zyprexa	C		

Date:03/11/05ISR Number: 4605958-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549055A  
Age:2 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Life-Threatening	Overdose		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -			Unknown Medication	SS		
Initial or Prolonged						
Disability						

Date:03/11/05ISR Number: 4605959-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549068A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cerebrovascular Accident		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:03/11/05ISR Number: 4605988-1Report Type:Expedited (15-DaCompany Report #139634  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hypomania		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	8 DAY	Mania		Quilonum Retard	C	Glaxosmithkline	ORAL
42MMOL per day	YR	Sleep Disorder					
45MG per day	YR			Remergil	C		ORAL
5MG per day				Zolpidem	C		ORAL
6MG per day				Risperdal	C		ORAL

Date:03/11/05ISR Number: 4607180-3Report Type:Direct Company Report #CTU 242898  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 2 TID		Grand Mal Convulsion		Tramadol	PS		
Intervention to ONE Q D				Wellbutrin Xr	SS		
Prevent Permanent Impairment/Damage				Methylphenidate	C		
				Relafen	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/11/05ISR Number: 4609084-9Report Type:Expedited (15-DaCompany Report #CEL-2005-00282-ROC  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other Required Intervention to Prevent Permanent Impairment/Damage ORAL		Anxiety Blood Pressure Increased Dehydration	Consumer	Delsym Er Suspension (Dextromethorphan Hydrobromide)	PS		ORAL
DEE IMAGE PER ORAL	8 DAY	Dizziness Drug Interaction Dysgeusia Fatigue Feeling Abnormal Feeling Hot		Weelbutrin (Bupropion Hydrochloride) Robitussin	SS SS		ORAL ORAL
PER ORAL		Heart Rate Irregular Hyperreflexia		Promethazine Dm (Promethazine Dm)	SS		ORAL
		Loss Of Consciousness Muscle Twitching Myoclonus Nausea Serotonin Syndrome Tremor		Allegra (Fexofenadine Hydrochloride) Thyroxine (Levothyroxine Sodium) Azithromycin (Azithromycin) Biaxin (Clarithromycin)	C C C C		

Date:03/11/05ISR Number: 4609533-6Report Type:Expedited (15-DaCompany Report #GXKR2005US00689  
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Bundle Branch Block Left Cardiovascular Disorder	Literature Health	Divalproex (Ngx) (Divalproex)	PS		ORAL
5920 MG, ORAL		Disorientation Electrocardiogram Qt Corrected Interval	Professional	Ziprasidone (Ziprasidone) Bupropion (Ngx)	SS		ORAL

15700 MG	Prolonged	(Bupropion)	SS	ORAL
(PROBABLY	Grand Mal Convulsion			
FEWER), ORAL	Intentional Misuse			
72 G, ORAL	Lethargy	Fluoxetine (Ngx)		
	Medication Error	(Fluoxetine)	SS	ORAL
	Supraventricular			
	Tachycardia			

Date:03/14/05ISR Number: 4607745-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0537281A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Arthralgia		Lamictal	PS	Glaxosmithkline	ORAL
2 YR							
Other		Chest Discomfort		Wellbutrin	SS	Glaxosmithkline	ORAL
200MG Twice							
per day	2 YR	Facial Pain					
		Gait Disturbance					
		Oedema Peripheral					

Date:03/14/05ISR Number: 4607746-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0543520A

Age:37 YR Gender:Female I/FU:F

Outcome	PT
Other	Amnesia
	Grand Mal Convulsion

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Insomnia Tongue Biting				
Dose	Duration		Report Source	Product	Role	Manufacturer
242 DAY				Wellbutrin	PS	Glaxosmithkline
60MG Per day				Prozac	C	
						Route
						ORAL

Date:03/14/05ISR Number: 4607747-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0546535A  
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Unknown	WK						
Hospitalization -		Grand Mal Convulsion		Trazodone	C		
UNKNOWN	100MG At						
Initial or Prolonged							
night							
Disability				Ultracet	C		
Other							

Date:03/14/05ISR Number: 4607748-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549034A  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rectal Haemorrhage		Wellbutrin	PS	Glaxosmithkline	ORAL
3 MON							
		Stool Analysis Abnormal		Zoloft	C		
				Synthroid	C	Glaxosmithkline	
				Lovastatin	C		
				Accupril	C		
				Estrace	C		
				Multivitamin	C		

Date:03/14/05ISR Number: 4607750-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549064A  
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150MG Twice Other per day			Hepatic Enzyme Increased Hepatic Failure	Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
				Paxil	C	Glaxosmithkline	
				Coumadin	C	Glaxosmithkline	
				Lipitor	C		
				Toprol	C		
				Klonopin	C		

Date:03/14/05ISR Number: 4607751-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549208A  
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - INTRAVENOUS Initial or Prolonged Other	300MG	Unknown	Intentional Misuse Local Swelling	Wellbutrin Xl	PS	Glaxosmithkline	

Date:03/14/05ISR Number: 4607753-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549209A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - INTRAVENOUS Initial or Prolonged			Intentional Misuse	Wellbutrin Xl	PS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/14/05ISR Number: 4607754-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549210A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Infection		Wellbutrin Xl	PS	Glaxosmithkline	
INTRAVENOUS		Intentional Misuse					

Date:03/14/05ISR Number: 4607756-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549256A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anger		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Fatigue					
per day	3 DAY	Homicidal Ideation		No Concurrent Medication	C		
		Irritability					
		Mood Swings					

Date:03/14/05ISR Number: 4607758-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549286A

Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin	PS	Glaxosmithkline	
300MG per day							

Date:03/14/05ISR Number: 4609724-4Report Type:Direct Company Report #CTU 243085

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abdominal Pain Upper		Bupropion Hcl	PS		
		Gastrooesophageal Reflux					
		Disease					
		Pharmaceutical Product					
		Complaint					

Outcome PT  
Other Acquired Porphyria  
Aggression  
Agitation  
Akathisia  
Amnesia  
Arthralgia  
Asthenia  
Bipolar Disorder  
Body Temperature  
Fluctuation  
Convulsion  
Coordination Abnormal  
Crying  
Depression  
Disability  
Dizziness  
Dysarthria  
Dysphemia  
Dyspnoea  
Electric Shock  
Emotional Disorder



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
42	DAY	Encephalopathy Facial Palsy Fatigue Feeling Abnormal	Wellbutrin	PS	Glaxosmithkline	ORAL
		Hair Metal Test Abnormal	Tranquilizer	SS		
		Head Banging	Depakote	SS		
		Headache	Zyprexa	SS		
		Hyperaesthesia	Alcohol	C		
		Hypersensitivity				
		Hypoxia				
		Immune System Disorder				
		Injury				
		Insomnia				
		Intentional Self-Injury				
		Mania				
		Mastocytosis				
		Metabolic Encephalopathy				
		Multiple Chemical Sensitivity				
		Muscle Spasms				
		Myalgia				
		Myoclonus				
		Nervous System Disorder				
		Nightmare				
		Overdose				
		Pain Of Skin				
		Petit Mal Epilepsy				
		Schizoaffective Disorder				
		Suicidal Ideation				
		Suicide Attempt				
		Throat Tightness				
		Tremor				
		Vestibular Disorder				
		Vomiting				

Date:03/15/05ISR Number: 4608469-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0506955A

Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice		Balance Disorder		Wellbutrin Sr	PS	Glaxosmithkline	ORAL

Initial or Prolonged  
per day

Dizziness

25MG Per day 7 MON

Drug Exposure During

Paxil Cr

SS

Glaxosmithkline

ORAL

Pregnancy

Prenatal Vitamin

C

Drug Withdrawal Syndrome

Electric Shock

Formication

Lethargy

Nausea

Paraesthesia

Peripheral Coldness

Pregnancy

Premature Baby

Premature Labour

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/15/05ISR Number: 4608476-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0545918A

Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
8	DAY						
		Burning Sensation		Delsym	SS	Glaxosmithkline	
		Dizziness		Allegra	C		
		Dysgeusia		Promethazine Dm	C		
		Hot Flush		Robitussin Dm	C		
		Hypertension		Levothyroxine	C	Glaxosmithkline	
		Loss Of Consciousness		Z Pack	C		
		Muscle Twitching		Biaxin	C		
		Nausea					
		Skin Burning Sensation					
		Tremor					

Date:03/15/05ISR Number: 4608478-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0547176A

Age:13 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Breast Mass		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Gynaecomastia		No Concurrent Medication	C		

Date:03/15/05ISR Number: 4608485-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549462A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Drug Interaction		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Ill-Defined Disorder		Lithium	SS	Glaxosmithkline	
				Cogentin	SS		
				Klonopin	SS		
				Trazodone	SS		
				Prolixin	SS		
				Seroquel	SS		

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Drug Interaction		Wellbutrin Sr Cogentin Trazodone Seroquel Klonopin Trilafon	PS SS SS SS SS SS	Glaxosmithkline     Glaxosmithkline	ORAL

Age:24 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Metrorrhagia		Zyban Loestrin	PS SS	Glaxosmithkline	ORAL ORAL

20MCG per day

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/15/05ISR Number: 4608492-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0374117A

Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Visual Field Defect		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	17	DAY					
UNKNOWN				Fusidic Acid	C		

Date:03/15/05ISR Number: 4608493-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0374119A

Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Constipation		Zyban	PS	Glaxosmithkline	ORAL
300MG Per day	10	DAY					
Other		Dizziness		Nicorette Gum	C	Glaxosmithkline	ORAL
2MG per day	11	DAY					
		Headache					
		Nausea					
		Speech Disorder					

Date:03/15/05ISR Number: 4608494-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0374120A

Age:22 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Crying		Zyban	PS	Glaxosmithkline	ORAL
300MG per day	9	DAY					
		Depression					

Date:03/15/05ISR Number: 4608495-5Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0374586A

Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Chest Pain		Bupropion			

Initial or Prolonged      Electrocardiogram      Hydrochloride      PS      Glaxosmithkline  
150MG Twice

Abnormal

per day      14      DAY

No Concurrent  
Medication      C

Date:03/15/05ISR Number: 4608496-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0374613A

Age:38 YR      Gender:Female      I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose      Duration						
Hospitalization - 150MG Per day 5      DAY	Chest Pain		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged	Electrocardiogram St Segment Depression Urticaria Generalised					

Date:03/15/05ISR Number: 4608521-3Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0548662A

Age:25 YR      Gender:Female      I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose      Duration						
Other	Breast Discharge Fatigue Oligomenorrhoea Vomiting In Pregnancy		Zyban	PS	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/15/05ISR Number: 4608824-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544253A  
 Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Unknown		Drug Exposure During Pregnancy					

Date:03/15/05ISR Number: 4608825-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544253B  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Congenital Anomaly		Wellbutrin	PS	Glaxosmithkline	
		Drug Exposure During Pregnancy					

Date:03/15/05ISR Number: 4610483-XReport Type:Expedited (15-DaCompany Report #HQWYE793203MAR05  
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Withdrawal Syndrome	Health Professional	Effexor (Venlafaxine Hydrochloride, Unspec)	PS		
150 MG -		Hallucination					
		Hyperhidrosis					

FREQUENCY

UNSPECIFIED

150 MG 1X PER

1 DAY, ORAL

Wellbutrin Xl (Bupropion Hydrochloride, )	SS		ORAL
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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Foreign	Zyban Tablet-Zyban			
Other		Decreased Appetite	Literature	(Bupropion			
		Depersonalisation	Health	Hydrochloride)	PS		
SEE DOSAGE		Drug Effect Decreased	Professional				
TEXT		Grandiosity					
		Hallucination					
		Ideas Of Reference					
		Illusion					
		Insomnia					
		Irritability					
		Mania					
		Paranoia					
		Psychotic Disorder					
		Suspiciousness					
		Tension					
		Tobacco Abuse					

Outcome	PT
Hospitalization -	Convulsion
Initial or Prolonged	Incontinence





Hospitalization -	Amnesia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Initial or Prolonged	Convulsion	Trazodone	C		
Other	Dizziness	Neurontin	C		
300MG As					
	Fall				
required					
	Head Injury				
	Loss Of Consciousness				
	Tremor				

Date:03/16/05ISR Number: 4610042-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549640A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dehydration		Wellbutrin	PS	Glaxosmithkline	ORAL
		Grand Mal Convulsion		Klonopin	SS		
UNKNOWN		Vomiting					

Date:03/16/05ISR Number: 4618087-XReport Type:Periodic Company Report #S05-USA-00304-01  
 Age:22 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -							
Initial or Prolonged		Pancreatitis Acute	Health Professional	Lexapro (Escitalopram)	PS		
				Wellbutrin			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Bupropion  
Hydrochloride) SS

Date:03/17/05ISR Number: 4610902-9Report Type:Expedited (15-DaCompany Report #200511975US  
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complication Of Delivery		Allegra	PS	Aventis	
dose: NOT		Drug Exposure During				Pharmaceuticals Inc.	
PROVIDED		Pregnancy					
		Haemorrhage		Bupropion	SS		
dose: NOT		Umbilical Cord		Celexa	SS		
PROVIDED		Abnormality					
dose: NOT				Zantac	SS		
PROVIDED							

Date:03/17/05ISR Number: 4611423-XReport Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-12888830  
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Bradycardia		Metoprolol Tartrate	PS	Apothecon	
1 YR							
Initial or Prolonged		Drug Interaction		Diltiazem Hcl	C		
1 YR							
12 DAY				Bupropion	I		

Date:03/17/05ISR Number: 4611579-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544527A  
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State		Wellbutrin	PS	Glaxosmithkline	ORAL
27 DAY							

Dysphemia  
 Grand Mal Convulsion  
 Overdose  
 Postictal State  
 Sleep Disorder  
 Tonic Clonic Movements

Adderall SS  
 Cold Medication SS

Date:03/17/05ISR Number: 4611580-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0546075A  
 Age:21 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Blood Caffeine Increased	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Drug Level Increased	Concerta	C		ORAL
108MG In the			Drug Toxicity				
morning	YR						
			Fatigue	Dextrostat	C	Glaxosmithkline	ORAL
2TAB Per day	YR						
			Grand Mal Convulsion				
			Hypertonia				
			Postictal State				
			Sleep Disorder				
			Syncope				
			Weight Decreased				

Date:03/17/05ISR Number: 4611585-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0550044A  
 Age: Gender:Male I/FU:I

Outcome PT  
 Other Arthralgia  
 Fatigue

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Systemic Lupus  
Erythematosus

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
			Wellbutrin	PS	Glaxosmithkline	ORAL

Date:03/17/05ISR Number: 4611589-1Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0357968A  
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction		Zyban	PS	Glaxosmithkline	ORAL
150MG See		Ecchymosis					
dosage text	48 DAY	Emotional Disorder		Alcohol	SS		
		Retrograde Amnesia		No Concurrent Medication	C		

Date:03/17/05ISR Number: 4611603-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0374756A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anorexia		Zyban	PS	Glaxosmithkline	ORAL
150MG See		Depersonalisation					
dosage text	3 WK	Depression					
		Emotional Distress					
		Suicidal Ideation					
		Weight Decreased					

Date:03/17/05ISR Number: 4611606-9Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0374928A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspnoea		Zyban	PS	Glaxosmithkline	
UNKNOWN	1TAB Twice						

per day

Eye Rolling

Hallucination  
Insomnia  
Nausea  
Palpitations

No Concurrent  
Medication

C

Date:03/17/05ISR Number: 4613437-2Report Type:Direct  
Age:53 YR Gender:Female I/FU:I

Company Report #CTU 243509

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150 G 1 PO		Abnormal Dreams Insomnia		Zyban Xl	PS		ORAL
DAILY X 3							
DAYS THEN 1							
PO BID	1	MON					

Date:03/17/05ISR Number: 4615820-8Report Type:Expedited (15-DaCompany Report #GXKR2005US00689  
Age:29 YR Gender:Male I/FU:I

Outcome  
Hospitalization -  
Initial or Prolonged

PT  
Bundle Branch Block Left  
Cardiovascular Disorder  
Disorientation  
Electrocardiogram Qt  
Corrected Interval  
Prolonged



Death Death Zyban PS Glaxosmithkline ORAL  
 50TAB Single

dose

Date:03/18/05ISR Number: 4614864-XReport Type:Direct Company Report #CTU 243681  
 Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Convulsion		Bupropion 75mg	PS		ORAL
Hospitalization -		Injury					
ORAL							
Initial or Prolonged		Road Traffic Accident		Formoterol	C		
Disability		Traumatic Brain Injury		Nifedipine	C		
				Isosorbide			
				Mononitrate	C		
				Metoprolol	C		
				Albuterol	C		
				Simvastatin	C		

Date:03/18/05ISR Number: 4616547-9Report Type:Expedited (15-DaCompany Report #L04-USA-07403-24  
 Age:30 YR Gender:Female I/FU:F

Outcome	PT
Death	Blood Pressure Decreased
Hospitalization -	Cardiac Arrest
Initial or Prolonged	Completed Suicide
	Convulsion
	Ejection Fraction



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
340 MG ONCE		Decreased Intentional Misuse Mental Status Changes Nausea	Literature Health	Lexapro (Escitalopram)	PS		ORAL
PO		Nervous System Disorder Ventricular Dysfunction	Professional				
2550 MG ONCE				Bupropion (Long-Acting) (Bupropion)	SS		ORAL
PO							
340 MG ONCE				Paroxetine	SS		ORAL
PO							
8.5 MG ONCE				Clonazepam	SS		ORAL
PO							

Date:03/21/05ISR Number: 4613914-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0374737A  
Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Unknown	21 DAY		Zyban	PS	Glaxosmithkline	ORAL
UNKNOWN				Femoston	C		
UNKNOWN				Co-Codamol	C		
UNKNOWN				Noctura	C		
		Mood Swings 12 WK Palpitations					

Date:03/21/05ISR Number: 4613919-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0375117A  
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	1TABS Per day 3 DAY	Abnormal Behaviour		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged	UNKNOWN	Aggression		Vasten	C		
	40MG Per day	Insomnia		Sectral	C		ORAL
50MG per day		Overdose		Triatec 2.5	C		
UNKNOWN		Suicidal Ideation		Lexomil	C		ORAL
.25UD See		Suicide Attempt					
dosage text				Kardegic 160	C		
UNKNOWN	160MG Per day						

Date:03/21/05ISR Number: 4616353-5Report Type:Expedited (15-DaCompany Report #IMP\_0840\_2005  
Age:43 YR Gender:Female I/FU:I

Outcome	PT
Other	Abdominal Pain Upper
	Bone Pain
	Cardiac Disorder
	Chest Pain
	Eating Disorder
	Feeling Abnormal
	Gastroesophageal Reflux
	Disease
	Hepatic Steatosis
	Hyperhidrosis
	Hypoaesthesia
	Nausea
	Pain
	Pain In Jaw
	Palpitations
	Pharmaceutical Product
	Complaint

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Weight Decreased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG QDAY		Consumer	Bupropion	PS		ORAL
PO			Atenolol	C		
			Ezetimibe And			
			Simvastatin	C		
			Nexium /Usa/	C		

Date:03/22/05ISR Number: 4615168-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0374668A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Zyban	PS	Glaxosmithkline	ORAL
50TAB Single		Overdose					
dose							

Date:03/22/05ISR Number: 4617548-7Report Type:Expedited (15-DaCompany Report #HQWYE150815MAR05  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Withdrawal Syndrome Suicidal Ideation Suicide Attempt	Consumer	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		ORAL
225 MG 1X PER				Ativan (Lorazepam, Unspec)	SS		
1 DAY, ORAL				Wellbutrin (Bupropion Hydrochoride, )	SS		

Date:03/22/05ISR Number: 4617549-9Report Type:Expedited (15-DaCompany Report #HQWYE007210MAR05  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Crying Suicidal Ideation Treatment Noncompliance Vomiting	Consumer	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		ORAL
SEE IMAGE				Ativan (Lorazepam, Unspec) Wellbutrin (Bupropion Hydrochloride)	SS  SS		

Date:03/22/05ISR Number: 4620242-XReport Type:Direct Company Report #CTU 243983  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash Erythematous Rash Pruritic		Roxicet 5mg/325mg 5/325 Roxane Lab	PS	Roxane Lab	ORAL
ONE TAB Q8H							
ORAL				Bupropion Hcl Extended Rele 150 Mg			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Eon Labs SS Eon Labs ORAL

ONE TAB BID

ORAL

Date:03/23/05ISR Number: 4617153-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544500A  
Age:11 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day 11 MON		Cardio-Respiratory Arrest	Wellbutrin	PS	Glaxosmithkline	ORAL
Life-Threatening	40MG per day			Strattera	SS		

Date:03/23/05ISR Number: 4617166-0Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0365531A  
Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Twice		Anorexia	Zyban	PS	Glaxosmithkline	
UNKNOWN	per day 8 DAY		Anxiety				
			Confusional State				
			Coordination Abnormal				
			Disorientation				
			Feeling Abnormal				
			Feeling Of Despair				
			Hallucination				
			Mental Disorder				

Date:03/23/05ISR Number: 4617176-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0374974A  
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Unknown 7 DAY		Anger	Zyban	PS	Glaxosmithkline	ORAL
	40MG Per day		Injury	Atorvastatin	C		ORAL

300MG Per day	Aspirin	C	Glaxosmithkline	ORAL
5MG Per day	Ramipril	C		ORAL
75MG Per day	Clopidogrel	C		ORAL

Date:03/23/05ISR Number: 4617181-7Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0375449A  
 Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence		Zyban	PS	Glaxosmithkline	
UNKNOWN							

Date:03/23/05ISR Number: 4617183-0Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0375519A  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain		Zyban	PS	Glaxosmithkline	ORAL
150MG See		Colitis Ischaemic					
dosage text	1	MON					
		Insomnia					
		Rectal Haemorrhage					

Date:03/23/05ISR Number: 4618905-5Report Type:Expedited (15-DaCompany Report #S05-USA-00523-01  
 Age:60 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Catheter Sepsis
Initial or Prolonged	Coma
	Confusional State

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Interaction Hypotension Respiratory Failure	Report Source	Product	Role	Manufacturer	Route
OROPHARINGEAL	20 MG QD PO		Health Professional	Lexapro (Escitalopram)	PS		
300 MG QD PO				Lexapro (Escitalopram)	SS		
				Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
150 MG QD PO				Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
150 MCG BID				Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
PO				Atenolol	C		
				Isosorbide	C		
				Potassium	C		
				Zocor	C		
				Aspirin	C		
				Avandia (Rosiglitazone Maleate)	C		
				Nexium (Esomeprazole)	C		
				Aricept (Donepezil Hydrochloride)	C		
				Neurontin (Gabapentin)	C		
				Duragesic (Fentanyl)	C		
				Hydrocodone	C		
				Centrum Silver	C		
				Magnesium	C		
				Vitamin B12	C		
				Diazepam	C		

Date:03/23/05ISR Number: 4621313-4Report Type:Direct  
Age:18 YR Gender:Female I/FU:I

Company Report #CTU 244005

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150 PO Initial or Prolonged	Grand Mal Convulsion		Wellbutrin Sr	PS		ORAL

Date:03/23/05ISR Number: 4621356-0Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 244100

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 75MG 2 Initial or Prolonged TABS BID  ORAL	Confusional State  Fall  Gait Disturbance  Head Injury Subdural Haemorrhage		Bupropion 75mg	PS		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/24/05ISR Number: 4617555-4Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20050105542  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
OROPHARINGEAL			Abdominal Distension	Topamax	PS		
UNKNOWN			Fatigue	Wellbutrin Xl	SS		
			Nausea	Lamictal	C		
			Sjogren'S Syndrome				
			Somnolence				
			Tremor				

Date:03/24/05ISR Number: 4617606-7Report Type:Expedited (15-DaCompany Report #US-MERCK-0503USA03482  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -			Adverse Event	Cogentin	PS	Merck & Co., Inc	ORAL
Initial or Prolonged			Drug Interaction	Wellbutrin Sr	SS		
UNKNOWN				Lithium Carbonate	SS		ORAL
UNKNOWN				Klonopin	SS		
UNKNOWN				Trazodone			
UNKNOWN				Hydrochloride	SS		
UNKNOWN				Seroquel	SS		
UNKNOWN				Prolixin	SS		

Date:03/24/05ISR Number: 4617822-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0534857A  
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -			Abdominal Pain Upper	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3 DAY							
Initial or Prolonged			Anger	Tagamet	C	Glaxosmithkline	
			Anorexia	Vicodin	C		
			Bipolar Disorder				

Coordination Abnormal  
Dementia  
Diarrhoea  
Disturbance In Attention  
Fatigue  
Nausea

Date:03/24/05ISR Number: 4617834-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0550280A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Death		Gastrointestinal Haemorrhage		Lamictal	SS	Glaxosmithkline	ORAL
2	DAY			Klonopin	C		
				Symbyax	C		

Date:03/24/05ISR Number: 4617837-6Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0550469A

Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain		Zyban	PS	Glaxosmithkline	ORAL
150MG See		Dizziness					
dosage text	7	DAY					
		Insomnia					
		Nausea					
		Oedematous Pancreatitis					

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Date:03/24/05ISR Number: 4617862-5Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0375746A  
Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
UNKNOWN							
1MG At night	YR						

13 DAY  
Erythema  
Rash  
Rash Papular  
Skin Disorder

Zyban  
Flunitrazepam  
PS  
C  
Glaxosmithkline  
ORAL

Date:03/24/05ISR Number: 4617864-9Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0375754A  
Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Anger
Initial or Prolonged	Crying
	Fall
	Feeling Abnormal
	Intentional Misuse
	Mood Swings
	Multiple Drug Overdose
	Night Sweats

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Suicide Attempt Tearfulness	Report Source	Product	Role	Manufacturer	Route
UNKNOWN				Zyban	PS	Glaxosmithkline	
				Contraceptive Pill	C		ORAL
				Valium	C		
				Sleeping Tablets	C		

Date:03/24/05ISR Number: 4619554-5Report Type:Expedited (15-DaCompany Report #2005-00954  
Age:56 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Drug Interaction Potentiation Sinus Bradycardia	Literature Health Professional	Metoprolol (Watson Laboratories)(Emtopr olol Tartrate) Tablet	PS	Watson Laboratories	ORAL
	75 MG, DAILY, ORAL	365 DAY						
	150 MG, BID, ORAL	12 DAY			Bupropiain (Bupropion)	SS		ORAL
					Diltiazem	C		

Date:03/25/05ISR Number: 4619133-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0546075A  
Age:21 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Blood Caffeine Increased Drug Toxicity		Wellbutrin Xl Concerta	PS C	Glaxosmithkline	ORAL ORAL
	108MG In the morning	YR	Fatigue					
	2TAB Per day	YR	Grand Mal Convulsion		Dextrostat	C	Glaxosmithkline	ORAL
			Hypertonia Loss Of Consciousness					

Postictal State  
Sleep Disorder  
Stress  
Syncope  
Weight Decreased

Date:03/25/05ISR Number: 4619138-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549286A  
Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300MG per day		Abortion Spontaneous		Wellbutrin	PS	Glaxosmithkline	
		Drug Exposure During Pregnancy Pregnancy					

Date:03/25/05ISR Number: 4619147-XReport Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0364019A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
UNKNOWN		Hyperthyroidism		Zyban	PS	Glaxosmithkline	
		Personality Change		Thyroxine	SS	Glaxosmithkline	
UNKNOWN		Visual Acuity Reduced Visual Field Defect					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/25/05ISR Number: 4619154-7Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0374919A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Zyban	PS	Glaxosmithkline	
UNKNOWN		4 DAY					
		Feeling Abnormal					
		Hypersensitivity					
		Loss Of Consciousness					

Date:03/25/05ISR Number: 4619162-6Report Type:Expedited (15-DaCompany Report #BE-GLAXOSMITHKLINE-B0375805A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Abortion Induced		Zyban	PS	Glaxosmithkline	
Congenital Anomaly		Asphyxiating Thoracic Dystrophy					
		Drug Exposure During Pregnancy					
		Limb Deformity					

Date:03/25/05ISR Number: 4619753-2Report Type:Expedited (15-DaCompany Report #US-MERCK-0304USA00834

Age:63 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Acute Myocardial
Initial or Prolonged	Infarction
Disability	Angina Unstable
Other	Anxiety
	Arthralgia
	Arthropathy
	Back Pain
	Benign Prostatic
	Hyperplasia
	Cardiac Failure
	Congestive
	Chest Pain
	Coronary Artery Disease
	Coronary Artery
	Restenosis
	Depression

Diabetes Mellitus  
Non-Insulin-Dependent  
Diabetic Nephropathy  
Diabetic Neuropathy  
Diabetic Retinopathy  
Drug Hypersensitivity  
Drug Ineffective  
Fall  
Hepatitis  
Herpes Zoster  
Hypercholesterolaemia  
Infection  
Injury  
Intervertebral Disc  
Degeneration  
Intervertebral Disc  
Disorder  
Ischaemic Cardiomyopathy  
Oedema Peripheral  
Pneumonia



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Renal Cyst					
		Sepsis					
		Sexual Dysfunction					
		Weight Increased		Vioxx	PS	Merck & Co., Inc	ORAL
10	DAY			Vioxx	SS	Merck & Co., Inc	ORAL
146	DAY			Vioxx	SS	Merck & Co., Inc	ORAL
UNKNOWN				Vioxx	SS	Merck & Co., Inc	ORAL
10	DAY			Zyban	SS		
146	DAY			Vioxx	SS	Merck & Co., Inc	ORAL
UNKNOWN		2170 DAY		Zestril	C		
UNKNOWN		2170 DAY		Zestril	C		
UNKNOWN				Tenormin	C		
UNKNOWN				Tenormin	C		
UNKNOWN				Glucophage	C		
UNKNOWN				Micronase	C		
UNKNOWN		1051 DAY		Lanoxin	C		
UNKNOWN				[Therapy Unspecified]	C		
UNKNOWN				Aciphex	C		

Date:03/25/05ISR Number: 4620882-8Report Type:Expedited (15-DaCompany Report #S05-USA-01449-01  
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Exposure During Pregnancy	Health Professional	Celexa (Citalopram Hydrobromide)	PS		
		Haemorrhage		Wellbutrin (Bupropion			
		Pregnancy					

300 MG QD; PO	Umbilical Cord	Hydrochloride)	SS	ORAL
	Abnormality	Wellbutrin (Bupropion Hydrochloride)	SS	ORAL
150 MG QD; PO		Wellbutrin (Bupropion Hydrochloride)	SS	ORAL
300 MG QD; PO		Allegra (Fexofenadine Hydrochloride)	SS	
		Zantac (Ranitidine Hydrochloride)	SS	

Date:03/25/05ISR Number: 4622592-XReport Type:Direct Company Report #CTU 244365  
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other Required Intervention to Prevent Permanent 200MG Q 12 Impairment/Damage ORAL		Pharmaceutical Product Complaint Urticaria		Bupropion Hcl Sr 200mg Watson/ Express Script	PS	Watson/Express Script	ORAL

Date:03/28/05ISR Number: 4620510-1Report Type:Expedited (15-DaCompany Report #A03200400600  
 Age:48 YR Gender:Male I/FU:F

Outcome  
Life-Threatening  
Hospitalization -

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Initial or Prolonged  
Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Impaired Work Ability		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
5MG See		Suicide Attempt		Ambien	SS		ORAL
dosage text	9 DAY						
150MG Per day				Wellbutrin	SS	Glaxosmithkline	
				Paroxetine Hydrochloride	SS	Glaxosmithkline	ORAL
20MG Per day	3 DAY						

Date:03/28/05ISR Number: 4620524-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549256A  
Age: Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Twice		Anger		Zyban	PS	Glaxosmithkline	ORAL
per day	3 DAY		Fatigue					
			Headache		No Concurrent Medication	C		
			Homicidal Ideation					
			Insomnia					
			Irritability					
			Mood Swings					

Date:03/28/05ISR Number: 4620529-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0550255A  
Age:48 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	3 MON		Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL
			Suicide Attempt		Prozac	C		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2 WK		Agitation Chills Drug Exposure During Pregnancy Insomnia Paraesthesia		No Concurrent Medication	C		

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Eczema		Wellbutrin	PS	Glaxosmithkline	ORAL
4 WK		Excoriation Pruritus		Cardiac Anti-Arrhythmic Medication Anti-Hypercholesterolaemic	C C		
Initial or Prolonged Other							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/28/05ISR Number: 4620532-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0550641A

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Condition Aggravated		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 11 MON						
Initial or Prolonged	Convulsion		Unknown	C		

Date:03/28/05ISR Number: 4622638-9Report Type:Direct

Company Report #CTU 244472

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
30 MG @ HS PO	Communication Disorder		Remeron	PS		ORAL
	Dizziness		Wellbutrin 150 Daily			
300 MG Q DAY	Feeling Abnormal		X 4 Then	SS		
	Headache					
	Nausea					
	Vision Blurred					

Date:03/29/05ISR Number: 4621894-0Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0550686A

Age:54 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Angina Pectoris		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day 2 DAY						
1TAB Weekly	Asthenia		Diazepam	C		ORAL
	Diarrhoea					
	Headache					

Date:03/29/05ISR Number: 4621905-2Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0364019A

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					

Other UNKNOWN	Hyperthyroidism	Zyban	PS	Glaxosmithkline
UNKNOWN	Personality Change	Thyroxine	SS	Glaxosmithkline
	Visual Acuity Reduced Visual Field Defect			

Date:03/29/05ISR Number: 4621909-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0375382A  
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Unknown		Abortion Induced		Zyban	PS	Glaxosmithkline	ORAL
		Drug Exposure During Pregnancy		Oral Contraception	C		ORAL

Date:03/29/05ISR Number: 4621915-5Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0375976A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other UNKNOWN		Aggression		Zyban	PS	Glaxosmithkline	
		Drug Interaction		Alcohol	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/05ISR Number: 4621934-9Report Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-12908679  
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Drug Interaction		Trazodone Hcl	PS	Apothecon	
Initial or Prolonged			Prolixin	I	Apothecon	
			Wellbutrin Sr	I		
			Lithium	I		ORAL
			Benztropine Mesylate	I		
			Clonazepam	I		
			Quetiapine Fumarate	I		

Date:03/30/05ISR Number: 4622837-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541137A  
Age:22 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Alcoholic Pancreatitis		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day						
Initial or Prolonged	Disease Recurrence		Alcohol	SS		
Other			Lexapro	C		
40MG Per day						

Date:03/30/05ISR Number: 4622838-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541603A  
Age:68 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Abdominal Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day						
Initial or Prolonged	Alanine Aminotransferase		Percocet	C		
Disability	Increased		Fosamax	C		
70MG Weekly						
Other	Anorexia		Allegra	C		
180MG Per day						
	Aspartate		Bextra	C		
20MG Per day						
	Aminotransferase		Nu-Iron	C		
150MG Per day						
	Increased		Magnesium Oxide	C		
400MG Three						

times per day	Blood Alkaline			
	Phosphatase Increased	Duoneb	C	
	Diarrhoea	Combivent	C	
	Electrolyte Imbalance	Advair	C	Glaxosmithkline
1PUFF Twice	Ileus			
per day	Intestinal Perforation	Tiazac	C	Glaxosmithkline
300MG Per day	Liver Function Test	Lotensin	C	Glaxosmithkline
10MG Per day	Abnormal	Bethanecol	C	
50MG Four	Nausea			
times per day	Peritonitis	Effexor Xr	C	
150MG Per day	Volvulus Of Bowel	Sonata	C	
10MG At night	Vomiting	Aciphex	C	
20MG Per day	Weight Decreased			

Date:03/30/05ISR Number: 4622841-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549064A  
Age:55 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150MG Twice Other per day	Hepatic Enzyme Increased Hepatic Failure Jaundice		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
	Prothrombin Time Prolonged		Paxil Coumadin Lipitor Toprol Klonopin	C C C C C	Glaxosmithkline Glaxosmithkline	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/31/05ISR Number: 4624153-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0551834A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion Dizziness Drug Toxicity Feeling Abnormal Headache Loss Of Consciousness		Wellbutrin	PS	Glaxosmithkline	

Date:03/31/05ISR Number: 4624178-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0375831A

Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 15MG Twice per day	17 DAY	Blood Glucose Decreased Circulatory Collapse		Zyban	PS	Glaxosmithkline	ORAL
4MG per day		Hypoglycaemia		Simvastatin	C		ORAL
15MG per day		Loss Of Consciousness		Lansoprazole	C		ORAL
20MG per day				Enalapril	C		ORAL
SUBCUTANEOUS				Mixtard	C		

Date:03/31/05ISR Number: 4627220-5Report Type:Expedited (15-DaCompany Report #2005UW04517

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Drug Interaction	Health Professional	Seroquel Wellbutrin - Slow Release Lithium Cogentin Klonopin Trazodone Prolixin	PS SS SS SS SS SS SS		

Date:03/31/05ISR Number: 4632038-3Report Type:Periodic  
Age:22 YR Gender:Male I/FU:I

Company Report #200411259BCC

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1320 MG, ONCE, ORAL		No Adverse Drug Effect	Health Professional  Other	Aleve Tablets (Naproxen Sodium)	PS		ORAL
1500 MG, ONCE, ORAL				Zyban (Bupropion Hydrochloride)	SS		ORAL

Date:04/01/05ISR Number: 4625324-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549225A  
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Amnesia Aura Grand Mal Convulsion Mania		Wellbutrin No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/01/05ISR Number: 4625333-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0551303A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness		Wellbutrin	PS	Glaxosmithkline	ORAL
225MG Twice							
		Medication Error					
per day							
		Meniere'S Disease		Premarin	C		
		Otosclerosis		Maxalt	C		
				Flexeril	C		
				Temazepam	C		

Date:04/01/05ISR Number: 4625334-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0551305A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
10 WK							
		Anger		Clonopin	C		
		Crying		Sudafed	C	Glaxosmithkline	
		Distractibility		Nasonex	C		
		Disturbance In Attention		Antibiotic	C		
		Dizziness					
		Emotional Disorder					
		Memory Impairment					
		Nausea					
		Panic Attack					
		Suicidal Ideation					

Date:04/01/05ISR Number: 4625335-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0551463A

Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Testosterone		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Increased		Testosterone Gel	SS	Glaxosmithkline	
		Gynaecomastia		Ativan	C		
		Haematocrit Increased		Viramune	C		
		Weight Decreased		Videx Ec	C		
				Famvir	C	Glaxosmithkline	

Date:04/01/05ISR Number: 4625336-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0551501A  
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice			Disturbance In Attention	Bupropion	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	2 WK		Feeling Abnormal				
			Loss Of Consciousness	Percocet	C		
			Vomiting				

Date:04/01/05ISR Number: 4625340-2Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0302490A  
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Headache	Zyban	PS	Glaxosmithkline	
Initial or Prolonged			Leukopenia	Bricanyl	C		
			Meningitis	Pulmicort	C		
			Neutropenia				
			Pain				
			Pyrexia				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/01/05ISR Number: 4625364-5Report Type:Expedited (15-DaCompany Report #CH-GLAXOSMITHKLINE-B0376363A  
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day	18 DAY	Abnormal Behaviour		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Amnesia Blood Prolactin Increased Cerebral Atrophy Clonus Confusional State Convulsion Crying Cyanosis Disorientation Dyskinesia Grand Mal Convulsion Posture Abnormal Respiration Abnormal					

Date:04/01/05ISR Number: 4628372-3Report Type:Expedited (15-DaCompany Report #IMP\_0840\_2005  
 Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150 MG QDAY		Abdominal Pain Upper	Consumer	Bupropion	PS		ORAL
PO		Alopecia Back Pain Chest Pain Constipation Decreased Activity Eating Disorder Feeling Abnormal Gastrooesophageal Reflux Disease Hepatic Steatosis Hyperhidrosis Hypoaesthesia Micturition Disorder Nausea Palpitations Pharmaceutical Product		Atenolol Ezetimibe Nexium	C C C		

Complaint  
Skin Exfoliation  
Weight Decreased

Date:04/04/05ISR Number: 4626324-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549621A  
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	20 DAY	Decreased Appetite					
		Decreased Interest					
		Irritability					
		Libido Decreased					
		Mood Swings					
		Sleep Disorder					
		Suicidal Ideation					

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Freedom Of Information (FOI) Report

Date:04/04/05ISR Number: 4626335-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0551784A  
Age:16 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS	Glaxosmithkline	

Date:04/04/05ISR Number: 4626340-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0551947A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 5 MON Initial or Prolonged		Convulsion Loss Of Consciousness Myalgia Tremor		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:04/05/05ISR Number: 4627135-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0520475A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG per day		Drug Exposure During Pregnancy Growth Retardation Intra-Uterine Death Pregnancy		Wellbutrin	PS	Glaxosmithkline	

Date:04/05/05ISR Number: 4627136-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0548375A  
Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 12 DAY		Abortion Spontaneous Drug Exposure During Pregnancy		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:04/05/05ISR Number: 4627137-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549655A  
Age:48 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Amnesia		Wellbutrin XL	PS	Glaxosmithkline	ORAL
Initial or Prolonged	Convulsion		Trazodone	C		
	Dizziness		Neurontin	C		
300MG As						
required	Fall					
	Head Injury					
	Loss Of Consciousness					
	Tremor					

Date:04/05/05ISR Number: 4627145-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0363857A  
Age:29 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Arthralgia
Initial or Prolonged	Asthenia
	Back Pain
	Cerebrovascular Accident
	Chest Pain
	Dizziness
	Fatigue
	Flank Pain



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
8 DAY		Headache Limb Discomfort Muscle Spasms	Zyban	PS	Glaxosmithkline	ORAL
INTRAVENOUS day	1391MG 15 DAY	Nausea Renal Failure Acute Per Rhabdomyolysis	Sodium Stibogluconate	SS	Glaxosmithkline	
500MG Three times per day	16 DAY	Vomiting	Keflex	C	Glaxosmithkline	ORAL
200MG Per day	34 DAY		Fluconazole	C		ORAL
			Rifampin	C		
			Ketoconazole	C		

Date:04/05/05ISR Number: 4627148-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0369079A  
Age: Gender:Male I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose Other		PT Suicidal Ideation	Zyban	PS	Glaxosmithkline	ORAL

Date:04/05/05ISR Number: 4627167-4Report Type:Expedited (15-DaCompany Report #DK-GLAXOSMITHKLINE-B0376672A  
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose Other UNKNOWN	300MG Per day	PT Delusion Flushing Pruritus Psoriasis Swelling Urticaria	Zyban	PS	Glaxosmithkline	

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Crying	Consumer	Ativan	PS		
		Suicidal Ideation		Effexor	SS		ORAL
225 MG Q DAY							
		Treatment Noncompliance					
PO							
		Vomiting		Effexor	SS		ORAL
225 MG Q DAY							
PO							
				Effexor	SS		ORAL
150 MG Q DAY							
PO							
				Effexor	SS		ORAL
75 MG Q DAY							
PO							
				Effexor	SS		ORAL
75 MG QOD PO							
				Wellbutrin	SS		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Exposure During	Foreign	Wellbutrin			
		Pregnancy	Study	Unspecified Tablet			
		Stillbirth	Literature	(Bupropion			
			Health	Hydrochloride)	PS		
			Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/05/05ISR Number: 4629327-5Report Type:Expedited (15-DaCompany Report #B0375846A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Exposure During Pregnancy Premature Separation Of Placenta	Foreign Study Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		

Date:04/05/05ISR Number: 4629328-7Report Type:Expedited (15-DaCompany Report #B0375846B

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death Neonatal Drug Exposure During Pregnancy	Foreign Study Literature Health Professional	Wellbutrin Unspecified Tablet (Bupropion Hydrochloride)	PS		
TRANSPLACENTA							
RY							

Date:04/05/05ISR Number: 4629509-2Report Type:Expedited (15-DaCompany Report #HQWYE364525MAR05

Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Withdrawal Syndrome Hypotension Therapeutic Response Decreased Tremor	Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		

SEE IMAGE

100 MG 1X PER

1 DAY

Bupropion (Amfebutamone, ) SS

Singulair (Montelukast Sodium) C Advair (Fluticasone

Propionate/Salmetero  
 l Xinafoate) C  
 Lorazepam  
 (Lorazepam) C  
 Trazodone  
 (Trazodone) C

Date:04/06/05ISR Number: 4628098-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549235A  
 Age:47 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 450MG Per day 367 DAY	Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Initial or Prolonged 200MG per day	Convulsion		Lamictal	C	Glaxosmithkline	ORAL
Other 15MG per day	Humerus Fracture		Ambien	C		ORAL
.5MG per day	Loss Of Consciousness		Xanax	C		ORAL
5MG per day	Tongue Biting		Zyprexa	C		ORAL
60MG per day			Paxil	C	Glaxosmithkline	ORAL
40MG per day 198 DAY			Strattera	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/06/05ISR Number: 4628099-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549643A  
Age:17 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	WK	Eye Rolling		Adderall Xr	C		
DAY		Fall					

Date:04/06/05ISR Number: 4628101-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0551938A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blister		Wellbutrin	PS	Glaxosmithkline	ORAL
5 MON		Dermatitis Exfoliative					
Initial or Prolonged		Skin Reaction					
		Urticaria					

Date:04/06/05ISR Number: 4628103-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0551946A  
Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bite		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2 MON	Mouth Injury		No Concurrent			
		Oral Pain		Medication	C		
		Tourette'S Disorder					

Date:04/06/05ISR Number: 4628104-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0551948A  
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Mood Altered		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice							

Initial or Prolonged Psychotic Disorder  
per day  
75MG Per day  
Suicide Attempt

Effexor	SS	ORAL
Remeron	C	
Effexor	C	
Metformin	C	
70/30 Humulin		
Insulin	C	
Lescol	C	
Lisinopril	C	
Seroquel	C	
Depakote Er	C	

Date:04/07/05ISR Number: 4629067-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549208A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Intentional Misuse		Wellbutrin Xl	PS	Glaxosmithkline	
INTRAVENOUS	300MG	Unknown					
Initial or Prolonged		Local Swelling					
Other		Medication Error					

Date:04/07/05ISR Number: 4629068-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549209A  
Age:45 YR Gender:Male I/FU:F

Outcome  
Hospitalization -  
Initial or Prolonged

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Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
INTRAVENOUS		Intentional Misuse		Wellbutrin Xl	PS	Glaxosmithkline	
		Medication Error					

Date:04/07/05ISR Number: 4629069-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549210A  
 Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	
INTRAVENOUS		Intentional Misuse					
Hospitalization -		Medication Error					
Initial or Prolonged							
Other							

Date:04/07/05ISR Number: 4629083-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0376590A  
 Age:68 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Toxic Epidermal		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day 11 DAY		Necrolysis					
Initial or Prolonged							

Date:04/07/05ISR Number: 4631156-3Report Type:Direct Company Report #CTU 245564  
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG ONCE		Depression		Wellbutrin Xl 300 Mg			
		Feeling Abnormal		Glaxosmithklkline	PS	Glaxosmithkline	ORAL
DAILY ORAL		Pain					
		Panic Attack					
		Pharmaceutical Product					
		Complaint					

Tremor

Date:04/08/05ISR Number: 4630076-8Report Type:Expedited (15-DaCompany Report #US-ABBOTT-03P-163-0225847-00  
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Toxicity		Hydrocodone	PS		
Not reported				Bupropion	SS		
Not reported				Venlafaxine	SS		
Not reported							

Date:04/08/05ISR Number: 4630527-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0545489A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Amnesia		Benzodiazepine	SS		
		Drug Withdrawal Syndrome					
		Grand Mal Convulsion					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/08/05ISR Number: 4630533-4Report Type:Expedited (15-DaCompany Report #2005/03/0457  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Per day	3	MON	Anger	Wellbutrin	PS	Glaxosmithkline	ORAL
			Homicidal Ideation	Peg-Intron	SS		
INTRAVENOUS							
			Mood Altered	Ribavirin	SS		ORAL
			Paranoia	Vicodin	C		ORAL
				Ziac	C		ORAL

Date:04/08/05ISR Number: 4630534-6Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0552669A  
 Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Per day		DAY	Haematemesis	Zyban	PS	Glaxosmithkline	ORAL
			Insomnia				
			Nausea				

Date:04/08/05ISR Number: 4630547-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0376592A  
 Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
150MG Per day	8	DAY	Blood Pressure Increased	Zyban	PS	Glaxosmithkline	ORAL
			Cardiovascular Disorder	Esomeprazole	C		ORAL
40MG per day			Drug Withdrawal Syndrome				
			Headache				

Date:04/08/05ISR Number: 4630548-6Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0376594A  
 Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Erythema Zyban PS Glaxosmithkline ORAL  
 150MG Twice  
 per day 6 DAY Hypersensitivity  
 Oropharyngeal Swelling  
 Rash Papular

Date:04/08/05ISR Number: 4630553-XReport Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0376835A  
 Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Laryngeal Oedema		Zyntabac	PS	Glaxosmithkline	ORAL
300MG per day	19 DAY	Rash					

Date:04/08/05ISR Number: 4630754-0Report Type:Expedited (15-DaCompany Report #US-ROCHE-399988  
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Asthenia		Klonopin	PS	Roche	ORAL
TAKEN EVERY		Confusional State					
NIGHT.		Decreased Appetite					
STARTED PRIOR		Dehydration					
TO 15		Drug Interaction					
NOVEMBER		Lethargy		Allopurinol	I	Roche	ORAL
TAKEN AT		Nausea					

STARTED  
 PRIOR TO 15  
 NOVEMBER 2004

FDA - Adverse Event Reporting System (AERS)

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INTRAVENOUS	37 DAY	Intron A	I	Roche	ORAL
TAKEN EVERY		Wellbutrin	I		ORAL
NIGHT.		Trazodone	I		ORAL
STARTED PRIOR					
TO 15					
NOVEMBER					
STARTED PRIOR		Zocor	I		ORAL
TO 15					
NOVEMBER 2004					
AND REPORTED					
TO BE 'LONG					
TAKEN AT		Remeron	I		ORAL
NIGHT.					
STARTED PRIOR					
TO 15					
NOVEMBER 2004					
STARTED PRIOR		Multivitamin Nos	I		ORAL
TO 15					
NOVEMBER 2004					
AND REPORTED					
AS 'LONG-TERM					
STARTED PRIOR		Glucosamine	I		ORAL
TO 15					

NOVEMBER 2004

AND REPORTED

AS 'LONG-TERM

Compazine I ORAL

Date:04/08/05ISR Number: 4630755-2Report Type:Expedited (15-DaCompany Report #US-ROCHE-400153

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Drug Interaction		Klonopin	PS	Roche	
UNKNOWN						
Initial or Prolonged			Wellbutrin Sr	I		ORAL
UNKNOWN			Lithium Salt	I		
UNKNOWN			Cogentin	I		
UNKNOWN			Trazodone	I	Roche	
UNKNOWN			Prolixin	I		
UNKNOWN			Seroquel	I		

Date:04/08/05ISR Number: 4630775-8Report Type:Expedited (15-DaCompany Report #US-ROCHE-400318

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Drug Interaction		Klonopin	PS	Roche	
UNKNOWN						
Initial or Prolonged			Wellbutrin Sr	I		ORAL
UNKNOWN			Benztropine	I		
UNKNOWN			Trazodone	I	Roche	
UNKNOWN			Seroquel	I		
UNKNOWN			Trilafon	I		

Date:04/08/05ISR Number: 4632745-2Report Type:Direct

Age:47 YR Gender:Female I/FU:I

Company Report #CTU 245717

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia		Wellbutrinxl 150mg	PS		ORAL
2 A DAY							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL

Date:04/11/05ISR Number: 4631870-XReport Type:Expedited (15-DaCompany Report #PHEH2005US03916  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Lamisil	PS	Novartis Sector: Pharma	
UNKNOWN				Wellbutrin	SS		ORAL

Date:04/11/05ISR Number: 4633096-2Report Type:Direct Company Report #CTU 245806  
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
20 MG 3 TIMES		Chest Pain		Baclofen	PS		ORAL
A DAY ORAL		Pain In Extremity					
300 MG 1 TIME				Wellbutrin 300 Mg	SS		ORAL
A DAY ORAL				Vioxx	C		
				Celebrex	C		
				Bextra	C		

Date:04/11/05ISR Number: 4635391-XReport Type:Expedited (15-DaCompany Report #2005051512  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 400 MG (200		Anxiety	Consumer	Celebrex (Celecoxib)	PS		ORAL
Initial or Prolonged MG, 2 IN 1		Back Pain					
Other		Depression					
D), ORAL		Nervousness		Bupropion			

Suicidal Ideation

Hydrochloride  
(Bupropion  
Hydrochloride)

SS

ORAL

300 MG (150

MG, 2 IN 1

D), ORAL

Escitalopram (Escitalopram)	C
Trazodone (Trazodone)	C
Seretide Mite (Fluticasone Propionate, Salmeterol Xinafoate)	C
All Other Therapeutic Products (All Other Therapeutic Products)	C

Date:04/12/05ISR Number: 4633101-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0537624A

Age: Gender:Female I/FU:F

Outcome	PT
Other	Alopecia Colitis Ulcerative Lethargy Rectal Haemorrhage

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Somnolence Stool Analysis Abnormal				
Dose	Duration		Report Source	Product	Role	Manufacturer
100MG Twice				Wellbutrin	PS	Glaxosmithkline
per day	3 MON			Wellbutrin Interferon	SS C	Glaxosmithkline
						Route
						ORAL

Date:04/12/05ISR Number: 4633102-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0542699A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose				Wellbutrin	PS	Glaxosmithkline
Other		Anger Depression Fear Irritability Suicidal Ideation				Route
						ORAL

Date:04/12/05ISR Number: 4633103-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0543321A  
Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose				Wellbutrin	PS	Glaxosmithkline
Other		Blood Albumin Increased				Route
100MG Twice		Liver Function Test				ORAL
per day		Abnormal		Nsaid	SS	
		Protein Total Increased		Lamictal	C	Glaxosmithkline
100MG Twice						
per day						

Date:04/12/05ISR Number: 4633109-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0552814A  
Age: Gender: I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Energy Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Loss Of Consciousness					
per day				Alcohol	C		

Date:04/12/05ISR Number: 4633138-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0376516A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris		Zyban	PS	Glaxosmithkline	

Date:04/12/05ISR Number: 4633982-3Report Type:Direct Company Report #CTU 245903  
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anxiety		Welbutrin 300 Mg	PS		ORAL
ONCE DAILY		Dizziness					
Initial or Prolonged		Feeling Abnormal					
ORAL		Insomnia					
		Palpitations					
		Panic Attack					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/12/05ISR Number: 4634800-XReport Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #CTU 245977

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Dizziness		Paxil Cr/ Paroxetine			
Other		Headache		50mg A Day	PS		
Required		Tinnitus		Wellbutrin 450mg			
Intervention to Prevent Permanent Impairment/Damage		Treatment Noncompliance		A Day	SS		

Date:04/13/05ISR Number: 4634081-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544500A  
 Age:11 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	11 MON	Cardio-Respiratory Arrest		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Life-Threatening		Drug Ineffective		Strattera	SS		
UNKNOWN		Fatigue		Amphetamine	SS		
		Feeling Drunk		Adderall	C		
		Lethargy					

Date:04/13/05ISR Number: 4634082-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0549256A  
 Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anger		Zyban	PS	Glaxosmithkline	ORAL
per day	3 DAY	Fatigue					
		Headache		No Concurrent Medication	C		
		Homicidal Ideation					
		Insomnia					
		Irritability					
		Mood Swings					

Date:04/13/05ISR Number: 4634090-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553014A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypokalaemia		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:04/13/05ISR Number: 4634092-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553163A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Medication Error		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:04/13/05ISR Number: 4634093-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553177A

Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
49 WK		Medication Error		No Concurrent Medication	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/13/05ISR Number: 4634094-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553283A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Adverse Event Convulsion		Wellbutrin Unspecified Medications	PS  C	Glaxosmithkline	ORAL
UNKNOWN							

Date:04/13/05ISR Number: 4634730-3Report Type:Direct Company Report #CTU 246047

Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death 450 MG 1X		Convulsion		Wellbutrin Xl	PS		
		Drowning					
DAILY							

Date:04/13/05ISR Number: 4636473-9Report Type:Direct Company Report #CTU 246010

Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Pharmaceutical Product Complaint		Bupropriion 100 Mg Eon Bupropriion 150 Mg Eon	PS  SS	Eon  Eon	

Date:04/14/05ISR Number: 4635159-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553424A

Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability 150MG In the morning	11 MON	Agitation  Anorexia Disturbance In Attention Drug Withdrawal Syndrome		Wellbutrin Xl  Protonix	PS  C	Glaxosmithkline	ORAL

Feeling Hot  
Insomnia  
Nausea

Date:04/14/05ISR Number: 4635163-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0347372A  
Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anorexia		Zyban	PS	Glaxosmithkline	
150MG	Unknown	21 DAY					
		Diplopia					
		Iiird Nerve Paralysis					
		Nystagmus					
		Vertigo					

Date:04/14/05ISR Number: 4636495-8Report Type:Direct Company Report #CTU 246227  
Age:13 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Agitation		Bupropion Sr			
		Depression		150mg	PS		ORAL
150MG	DAILY						
ORAL		Drug Effect Decreased					
		Pharmaceutical Product					
		Complaint					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/14/05ISR Number: 4636704-5Report Type:Expedited (15-DaCompany Report #2005052957  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Self-Medication	Health Professional Company Representative	Geodon (Ziprasidone) Bupropion Hydrochloride (Bupropion Hydrochloride) Olanzapine/Fluoxetine Hydrochloride (Fluoxetine Hydrochloride, Olanzapine) Mirtazapine (Mirtazepine) Obetrol (Amfetamine Aspartate, Amfetamine Sulfate, Dexamfetamine Saccharate,	PS		
					SS		
					SS		
					SS		

Date:04/14/05ISR Number: 4638581-5Report Type:Direct Company Report #CTU 246206  
 Age:11 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ONE Q AM		Abnormal Behaviour		Bupropion	PS		
		Aggression					
		Crying					
		Headache					
		Mood Swings					
		Pharmaceutical Product Complaint					
		Therapeutic Response					
		Unexpected With Drug					
		Substitution					

Date:04/15/05ISR Number: 4635879-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0552604A  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	10MG Per day	10 DAY	Aggression	Paroxetine	PS	Glaxosmithkline	ORAL
Initial or Prolonged	50MG Per day		Amnesia	Paxil Cr	SS	Glaxosmithkline	ORAL
	150MG Twice		Confusional State	Wellbutrin	SS	Glaxosmithkline	ORAL
	per day		Drug Ineffective				
	150MG Twice		Drug Withdrawal Syndrome	Wellbutrin Sr	C	Glaxosmithkline	ORAL
	per day		Hypertension				
			Suicidal Ideation	Metoprolol	C		
			Suicide Attempt				
			Tremor				

Date:04/15/05ISR Number: 4635880-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0552663A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	450MG Per day		Convulsion	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Mouth Haemorrhage	Lithium	C	Glaxosmithkline	
			Tongue Disorder	Seroquel	C		
				Ambien	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Prevacid C

Date:04/15/05ISR Number: 4635881-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0552994A  
 Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day			Grand Mal Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged			Petit Mal Epilepsy Staring	Klonopin	C		

Date:04/15/05ISR Number: 4635883-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553354A  
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 4 WK			Condition Aggravated	Wellbutrin	PS	Glaxosmithkline	ORAL
			Depression Overdose Suicide Attempt	Valium No Concurrent Medication	SS C		

Date:04/15/05ISR Number: 4635884-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553362A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Atrophic Vulvovaginitis	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:04/15/05ISR Number: 4635885-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553377A  
 Age:15 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Per day			Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged			Electrocardiogram Qt Prolonged	Topamax Xanax	C C		



Disability Overdose Clonidine C  
Other

Date:04/15/05ISR Number: 4636525-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0552854A  
Age:17 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Cystitis		Unspecified			
		Overdose		Medication	C		

Date:04/15/05ISR Number: 4640133-8Report Type:Direct Company Report #CTU 246361  
Age:11 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 150 MG QAM		Depression		Bupropion Sr 150 Mg	PS		ORAL
Initial or Prolonged ORAL		Drug Effect Decreased					
		Educational Problem					
		Pharmaceutical Product Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/18/05ISR Number: 4637628-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0547369A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Accidental Overdose		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Convulsion		Effexor	SS		
225MG In the morning		Muscle Twitching					

Date:04/18/05ISR Number: 4637631-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553387A

Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day		Feeling Of Despair					
		Hallucination		Aspirin	C	Glaxosmithkline	
		Medication Error					
		Suicidal Ideation					

Date:04/18/05ISR Number: 4638605-5Report Type:Direct Company Report #USP 57143

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
TABLET		Medication Error		Wellbutrin	PS	Mylan	
TABLET				Serzone	SS	Ivax	

Date:04/18/05ISR Number: 4638734-6Report Type:Direct Company Report #CTU 246414

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash Erythematous		Roxicet 5/325 Roxane			

ONE TAB Q6H	Rash Pruritic	Lab	PS	Roxane Lab	ORAL
ORAL					
		Bupropion Hcl 150 Mg			
ONE TAB BID		Eon Labs	SS	Eon Labs	ORAL
ORAL					

Date:04/19/05ISR Number: 4638921-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553367A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day 2 YR	Cardiomegaly		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Hypertension		Lopressor	C		
				Coumadin	C	Glaxosmithkline	
				Hct	C		
				Lisinopril	C		
				Aspirin	C	Glaxosmithkline	
				Unknown Medication	C		
				Progestin	C		

Date:04/19/05ISR Number: 4638922-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553397A

Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	400MG Unknown	Parkinson'S Disease		Wellbutrin	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/19/05ISR Number: 4638929-1Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0554063A

Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erythema		Clavulin	PS	Glaxosmithkline	ORAL
250MG Three							
		Fibromyalgia					
times per day	11	DAY					
		Musculoskeletal Stiffness		Zyban	SS	Glaxosmithkline	ORAL
150MG Twice							
		Myalgia					
per day	9	DAY					
		Pruritus		Apo-Salvent	C	Glaxosmithkline	
RESPIRATORY							
(INHALATION)		Wheezing					
				Becloforte	C	Glaxosmithkline	
RESPIRATORY							
(INHALATION)							

Date:04/19/05ISR Number: 4638937-0Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0374613A

Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Acute Coronary Syndrome		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	6	DAY					
Hospitalization -		Barrett'S Oesophagus					
Initial or Prolonged		Chest Pain					
		Electrocardiogram St					
		Segment Depression					
		Hiatus Hernia					
		Oesophagitis					
		Urticaria Generalised					

Date:04/19/05ISR Number: 4638948-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0377367A

Age:27 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other 150MG Twice per day	Alcohol Interaction Hallucination Injury	Zyban	PS	Glaxosmithkline	ORAL
20MG per day		Alcohol Omeprazole	SS C	Glaxosmithkline	ORAL

Date:04/19/05ISR Number: 4638958-8Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0378012A  
 Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening UNKNOWN		Convulsion		Zyban	PS	Glaxosmithkline	
Hospitalization - Initial or Prolonged		Overdose Respiratory Arrest					

Date:04/20/05ISR Number: 4640251-4Report Type:Expedited (15-DaCompany Report #13611  
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Deficiency Anaemia Gastrointestinal Infection Polyhydramnios		Bupropion Prenatal Vitamins Calcium	PS C C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/20/05ISR Number: 4640252-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553608A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaemia		Wellbutrin	PS	Glaxosmithkline	ORAL
200MG Twice		Epistaxis					
per day		Gastrointestinal		Plavix	SS		
		Haemorrhage		Aspirin	SS	Glaxosmithkline	

Date:04/20/05ISR Number: 4640254-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553794A

Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 2 YR		Death		Effexor	C		
Hospitalization -		Overdose					
Initial or Prolonged							

Date:04/20/05ISR Number: 4640256-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0554033A

Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Post Procedural		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day 1 YR		Haemorrhage		Klonopin	C		
				Nexium	C		
				Lasix	C	Glaxosmithkline	

Date:04/20/05ISR Number: 4640270-8Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0377668A

Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown 39 DAY							

Joint Crepitation  
 10MG per day Amitriptyline C ORAL  
 200MG per day Ibuprofen C Glaxosmithkline ORAL

Date:04/20/05ISR Number: 4640273-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0378145A  
 Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema Arthralgia Blood Immunoglobulin E Increased Pruritus		Zyban	PS	Glaxosmithkline	ORAL

Date:04/20/05ISR Number: 4640316-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0530626A  
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	6 WK			Eskalith	PS	Glaxosmithkline	ORAL
	300MG Per day 6 WK			Wellbutrin	SS	Glaxosmithkline	ORAL
	100MG Four times per day 5 YR			Wellbutrin	SS	Glaxosmithkline	ORAL
				Humibid	C		
				Robaxin	C		
				Ultram	C		
				Neurontin	C		
				Valium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/20/05ISR Number: 4642083-XReport Type:Expedited (15-DaCompany Report #2005052957

Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Self-Medication	Health Professional Company Representative	Geodon (Ziprasidone) Bupropion Hydrochloride (Bupropion Hydrochloride) Olanzapine/Fluoxetine Hydrochloride (Fluoxetine Hydrochloride, Olanzapine) Mirtazapine (Mirtazapine) Obetrol (Amphetamine Aspartate, Amphetamine Sulfate, Dexamethasone Saccharate,	PS		
					SS		
					SS		
					SS		

Date:04/21/05ISR Number: 4641477-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553582A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiovascular Disorder Death Intentional Misuse		Wellbutrin	PS	Glaxosmithkline	

Date:04/21/05ISR Number: 4641498-3Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0370764A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability UNKNOWN	150MG	Pruritus Unknown Sleep Disorder Tooth Disorder Toothache		Zyban	PS	Glaxosmithkline	



Date:04/21/05ISR Number: 4642187-1Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 246814

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams Morbid Thoughts Nightmare Pharmaceutical Product Complaint		Wellbutrin Sr	PS		

Date:04/21/05ISR Number: 4643116-7Report Type:Expedited (15-DaCompany Report #2005057334  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthma	Consumer	Bextra (Valdecoxib)	PS		ORAL
20 MG, ORAL		Blood Pressure Increased Bronchospasm		Dilantin (Phenytoin Sodium)	SS		ORAL
ORAL		Convulsion Weight Increased		Corticosteroids (Corticosteroids)	SS		

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Freedom Of Information (FOI) Report

Bupropion  
 Hydrochloride  
 (Bupropion Hydrochloride) SS  
 Neurontin  
 (Gabapentin) SS  
 Etanercept  
 (Etanercept) C  
 Prednisone  
 (Prednisone) C  
 Vicodin (Hydrocodone  
 Bitartrate,  
 Paracetamol) C  
 Cyclobenzaprine  
 Hydrochloride  
 (Cyclobenzaprine  
 Hydrochloride) C  
 Lansoprazole  
 (Lansoprazole) C

Date:04/22/05ISR Number: 4642369-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0545927A  
 Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
100MG Per day		Adverse Drug Reaction		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Constipation		Wellbutrin	SS	Glaxosmithkline	
		Diplopia		Duragesic	C		
		Dry Mouth		Effexor	C		
		Hallucination, Auditory		Valium	C		
		Hallucination, Visual		Vicodin	C		
		Pruritus		Neurontin	C		
		Somnolence		Avandia	C	Glaxosmithkline	
		Treatment Noncompliance					
		Vision Blurred					

Date:04/22/05ISR Number: 4642376-6Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0302490A  
 Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization -	Anaemia	Zyban	PS	Glaxosmithkline
Initial or Prolonged	Dizziness	Bricanyl	C	
	Eye Pain	Pulmicort	C	
	Facial Pain	Zyrlex	C	Glaxosmithkline
UNKNOWN	Headache			
	Leukopenia			
	Meningitis			
	Metrorrhagia			
	Nausea			
	Neutropenia			
	Palpitations			
	Paraesthesia			
	Photosensitivity Reaction			
	Pyrexia			
	White Blood Cell Count Decreased			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/22/05ISR Number: 4642380-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0369124A

Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	300MG per day 9 DAY	Bradycardia		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Dyspnoea Exertional		Vaccine	SS	Glaxosmithkline	
SUBCUTANEOUS	2UNIT Single	Hepatocellular Damage					
dose	1 DAY	Pitting Oedema		Lescol	C		ORAL
1UNIT per day		Transaminases Increased					
		Weight Increased					

Date:04/22/05ISR Number: 4643809-1Report Type:Direct

Company Report #CTU 246932

Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Rash		Bupropion	PS		
				Isosorbide Dinitrate	C		
				Gemfibrozil	C		
				Metoprolol Tartrate	C		
				Diltiazem (Inwood)	C		
				Irbesartan	C		
				Glyburide	C		
				Sertraline Hcl	C		
				Aspirin	C		
				Calcipotriene	C		
				Alprazolam	C		
				Meloxicam	C		
				Omeprazole	C		
				Salicylic Acid			
				2/Sulfur 2%	C		
				Atorvastatin Calcium	C		
				Hydrocortisone	C		
				Clobetasol			
				Propionate	C		
				Albuterol	C		
				Coal Tar	C		
				Ipratropium Bromide	C		
				Acetaminophen	C		
				Metformin Hcl	C		

Date:04/22/05ISR Number: 4644497-0Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 246867

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Abdominal Pain Upper		Wellbutrin Sr (Or			
Initial or Prolonged	Haematemesis		Placebo)	PS	Glaxosmithkline	
150 BID						
	Ulcer Haemorrhage		Bupropion Sr	C		
			Vicodin	C		
			Levothyroxine	C		
			Acetaminophen	C		
			Albuterol Inhaler	C		
			Diphenhydramine Hcl	C		
			Hydrochlorothiazide	C		
			Aspirin	C		
			Lisinopril	C		
			Simvastatin	C		
			Cyclobenzaprine Hcl	C		
			Trazodone	C		
			Felodipine	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/25/05 ISR Number: 4644122-9 Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0554382A  
 Age:41 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Convulsion		Lexiva	PS	Glaxosmithkline	ORAL
Initial or Prolonged	Rash Erythematous		Wellbutrin	SS	Glaxosmithkline	ORAL
	Rash Generalised		Atazanavir	C		
	Rash Maculo-Papular		Ritonavir	C		
			Epzicom	C	Glaxosmithkline	
			Lipitor	C		
			Neurontin	C		
			Celexa	C		

Date:04/25/05 ISR Number: 4644133-3 Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0374776A  
 Age:40 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Anxiety		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice						
Initial or Prolonged	Chest Discomfort					
per day						
8 WK	Dyspnoea		No Concurrent			
	Hyperventilation		Medication	C		
	Paraesthesia					

Date:04/25/05 ISR Number: 4644142-4 Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0376835A  
 Age:33 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Other	Laryngeal Oedema		Zyntabac	PS	Glaxosmithkline	ORAL
300MG per day						
19 DAY	Rash					

Date:04/25/05 ISR Number: 4644172-2 Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0378887A  
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						

Other	Aggression	Zyban	PS	Glaxosmithkline	ORAL
	Anxiety				
	Disturbance In Attention				
	Drug Withdrawal Syndrome				
	Dry Mouth				
	Insomnia				

Date:04/25/05ISR Number: 4644738-XReport Type:Direct Company Report #CTU 247098  
 Age:36 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Convulsion		Bupropion	PS		ORAL
200MG QAM						
Initial or Prolonged	Muscle Spasms					
ORAL	Paraesthesia					

Date:04/25/05ISR Number: 4645859-8Report Type:Expedited (15-DaCompany Report #HQWYE023014APR05  
 Age:53 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Mood Altered	Consumer	Effexor Xr			
Initial or Prolonged	Psychotic Disorder		(Venlafaxine			
Other	Suicide Attempt		Hydrochloride,			
			Capsule, Extended			

Freedom Of Information (FOI) Report

75 MG 1X PER 1 DAY, ORAL	Release)	PS	ORAL
200 MG 2X PER 1 DAY, ORAL	Wellbutrin - Slow Release (Amfebutamone Hydrochloride)	SS	ORAL
	Remeron (Mirtazapine)	C	
	Metformin (Metformin)	C	
	Humulin 70/30 (Insulin Human Injection, Isophane/Insulin Human Zinc	C	
	Lescol (Fluvastatin Sodium)	C	
	Lisinopril (Lisinopril)	C	
	Seroquel (Quetiapine)	C	
	Valproate Semisodium (Valproate Semisodium)	C	

Date:04/25/05ISR Number: 4645914-2Report Type:Expedited (15-DaCompany Report #HQWYE017014APR05  
Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion Overdose	Health Professional Company Representative	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		ORAL
ORAL				Wellbutrin Xl (Bupropion Hydrochloride, )	SS		ORAL
ORAL							



Date:04/25/05ISR Number: 4646752-7Report Type:Direct  
Age:51 YR Gender:Male I/FU:I

Company Report #CTU 247099

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 150MG BID	Hallucination, Visual		Bupropion 150mg	PS		ORAL
Initial or Prolonged ORAL	Nightmare					

Date:04/26/05ISR Number: 4645199-7Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0554466A  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Congenital Anomaly 150MG Per day	Hypospadias		Bupropion	PS	Glaxosmithkline	
.137MG Per day	Microtia		Synthroid	C	Glaxosmithkline	
10MG Eight times per day			Domperidone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/26/05ISR Number: 4645200-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0554507A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Anger		Bupropion			
150MG Three		Asthenia		Hydrochloride	PS	Glaxosmithkline	ORAL
times per day		Depression					
150MG Three		Drug Ineffective		Wellbutrin	SS	Glaxosmithkline	ORAL
times per day		Irritability					
		Pathological Gambling					
		Suicidal Ideation					

Date:04/26/05ISR Number: 4645236-XReport Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0379013A

Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Malignant Respiratory		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization -		Tract Neoplasm					
Initial or Prolonged		Metastases To Lymph Nodes					
		Metastases To Skin					

Date:04/27/05ISR Number: 4646701-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553401A

Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Fall		Wellbutrin	PS	Glaxosmithkline	ORAL
11 DAY		Grand Mal Convulsion		Effexor Xr	C		ORAL
300MG In the		Myalgia					
morning		Periorbital Haematoma		Remeron	C		ORAL
45MG At night				Premarin	C		
UNKNOWN	.125MG						

Unknown

Date:04/27/05ISR Number: 4646703-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0554522A  
Age:28 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 300MG Per day	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged	Dizziness		Paxil Cr	C	Glaxosmithkline	
	Migraine		Klonopin	C		
			Alcohol	C		

Date:04/27/05ISR Number: 4646704-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0554814A  
Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Drug Screen False Positive Overdose Rash		Wellbutrin No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

Date:04/27/05ISR Number: 4646705-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0554881A  
Age:54 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Adverse Drug Reaction Blister Malaise

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Musculoskeletal Stiffness Oedema Peripheral Skin Infection					
		Toxic Epidermal Necrolysis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:04/27/05ISR Number: 4646706-0Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0555062A  
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	47 DAY	Deafness		Wellbutrin	PS	Glaxosmithkline	ORAL
		Deafness Unilateral Tinnitus		Topiramate	C		

Date:04/27/05ISR Number: 4647391-4Report Type:Direct Company Report #CTU 247266  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other 450 MG DAILY		Aggression Anxiety		Wellbutrin Xl 450 Mg Glaxosmithkline	PS	Glaxosmithkline	ORAL
ORAL		Asthenia					
75 MG DAILY		Confusional State		Zoloft 75 Mg	SS		ORAL
ORAL		Depression					
		Drug Effect Decreased Dysstasia Hyperhidrosis Tension Tremor					

Date:04/28/05ISR Number: 4647823-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0550641A  
Age:16 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 446 DAY		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged Other		Aura Grand Mal Convulsion Tic		Unknown	C		

Date:04/28/05ISR Number: 4650532-6Report Type:Expedited (15-DaCompany Report #2005UW6257  
Age:9 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 600 MG PO		Dystonia	Health	Seroquel	PS		ORAL
Intervention to 400 MG PO			Professional	Seroquel	SS		ORAL
Prevent Permanent 6 MG Impairment/Damage				Risperdal	SS		
				Depakote	SS		
				Wellbutrin	SS		
				Strattera	SS		
				Trileptal			
				"Ciba-Geigy"	SS		
				Trileptal			
				"Ciba-Geigy"	SS		

DECREASING

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/28/05ISR Number: 4650654-XReport Type:Direct  
 Age:33 YR Gender:Female I/FU:I

Company Report #CTU 247385

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	XL 150 MG BID	Anxiety		Bupropion Generic	PS		
		Crying		Lexapro	C		
		Drug Effect Decreased		Zoloft	C		
		Fatigue		Celexa	C		
		Therapeutic Response		Remeron	C		
		Unexpected With Drug					
		Substitution					

Date:04/29/05ISR Number: 4649086-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0554844A  
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	15MG Per day	Bronchitis		Paxil	PS	Glaxosmithkline	ORAL
Initial or Prolonged	150MG Twice	Conversion Disorder		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
per day		Multiple Sclerosis					
		Tremor		Avonex	C		
				Ambien	C		
				Xanax	C		
				L-Thyroxine	C	Glaxosmithkline	
				Methadone	C	Glaxosmithkline	
				Baclofen	C		
				Neurotin	C		
				Lisinopril	C		
				Macrochantin	C		
				Gemfibrozil	C		
				Detrol La	C		
				Protonix	C		
				Zetia	C		
				Propoxyphene	C		
				Provigil	C		
				Keppra	C		

Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Compartment Syndrome		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Swelling		Geodon	C		

Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Bupropion	PS	Glaxosmithkline	
300MG per day				Lamotrigine	SS	Glaxosmithkline	
				Augmentin	C	Glaxosmithkline	
				Excedrin	C		
				Zyrtec	C	Glaxosmithkline	
				Multivitamin	C		
				Fibercon	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/29/05ISR Number: 4649094-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0555404A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS	Glaxosmithkline	
Life-Threatening		Overdose		Tylenol	SS	Glaxosmithkline	
Hospitalization -				Alcohol	SS		

Date:04/29/05ISR Number: 4649128-1Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0378887A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aggression Anxiety Disturbance In Attention Drug Withdrawal Syndrome Dry Mouth Headache Insomnia Tremor		Zyban	PS	Glaxosmithkline	ORAL

Date:04/29/05ISR Number: 4649130-XReport Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0378949A

Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anxiety		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Diplegia					
per day	13 DAY	Hallucination		Alprazolam	C		
UNKNOWN		Insomnia Monoplegia Panic Reaction Paranoia Rash					



Date:04/29/05ISR Number: 4650500-4Report Type:Expedited (15-DaCompany Report #2005-DE-01433GD

Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Overdose	Foreign Literature	Mirtazapine (Mirtazapine)	PS		
				Quetiapine (Quetiapine)	SS		
				Bupropion (Bupropion)	SS		
				Amitriptyline (Amitriptyline)	SS		
				Valproate (Valproic Acid)	C		

Date:05/02/05ISR Number: 4649640-5Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20050204345

Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Alcohol Poisoning		Fentanyl	PS		
UNKNOWN		Drug Toxicity		Citalopram	SS		
				Clonazepam	SS		
				Bupropion	SS		

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Freedom Of Information (FOI) Report

Alcohol SS

Date:05/02/05ISR Number: 4651622-4Report Type:Expedited (15-DaCompany Report #S05-USA-02072-01  
 Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Grand Mal Convulsion	Health Professional	Lexapro (Escitalopram)	PS		ORAL
10 MG QD PO				Lexapro (Escitalopram)	SS		ORAL
5 MG QD PO				Wellbutrin Xl (Bupropion)	SS		ORAL
450 MG QD PO				Wellbutrin Xl (Bupropion)	SS		ORAL
400 MG QD PO							

Date:05/03/05ISR Number: 4650727-1Report Type:Expedited (15-DaCompany Report #US-2004-BP-09934BP  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Oedema Peripheral Platelet Count Decreased		Spiriva Handihaler	PS	B.I. Pharmaceuticals, Inc. /Ridgefield	
RESPIRATORY (INHALATION)	1	MON		Bextra	SS		ORAL
				Wellbutrin	SS		ORAL
				Foradil	SS		
RESPIRATORY (INHALATION)				Prednisone	C		ORAL
				Fosamax	C		
				Foltz (Folic Acid)	C		ORAL
				Glucosamine	C		ORAL
				Duoneb	C		

RESPIRATORY

( INHALATION)

Date:05/03/05ISR Number: 4652895-4Report Type:Expedited (15-DaCompany Report #B0378357A

Age:24 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG/ PER DAY / ORAL	Convulsion Sinus Tachycardia	Foreign Literature Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
			Budesonide	C		

Date:05/04/05ISR Number: 4651808-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508446A

Age:39 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Twice Initial or Prolonged per day	Amnesia Convulsion Loss Of Consciousness Night Sweats		Wellbutrin	PS	Glaxosmithkline	ORAL
6 MON			Pepcid	C		
			Vicodin	C		
			Flexeril	C		
			Nasonex	C		
			Klonopin	C		
1MG Twice per day						

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Freedom Of Information (FOI) Report

Date:05/04/05ISR Number: 4651851-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0555664A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly			Congenital Eye Disorder Drug Exposure During Pregnancy	Wellbutrin Xl	PS	Glaxosmithkline	

Date:05/04/05ISR Number: 4651852-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0555841A

Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day 1 MON		Belligerence Emotional Disorder Feeling Abnormal Mental Status Changes Screaming	Wellbutrin Alcohol Remeron Klonopin	PS SS C C	Glaxosmithkline	ORAL

Date:05/04/05ISR Number: 4651853-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0555847A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Nephrolithiasis	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/04/05ISR Number: 4651881-8Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0379081A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	3 WK		Aggression Anorexia Depression Emotional Disorder Hypoaesthesia Insomnia Mania	Zyban No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Night Sweats  
Paraesthesia

Date:05/04/05ISR Number: 4653998-0Report Type:Expedited (15-DaCompany Report #10953  
Age:23 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 7.5 MG NOCTE 150 MG	Acute Psychosis Heart Rate Decreased Intentional Misuse Irritability Multiple Drug Overdose Paranoia Persecutory Delusion Psychomotor Agitation Somnolence Suicide Attempt	Literature Health Professional	Midazolam Bupropion	PS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/04/05ISR Number: 4654649-1Report Type:Direct  
Age:29 YR Gender:Female I/FU:I

Company Report #CTU 247789

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Erythema		Wellbutrin	PS		
Intervention to Prevent Permanent Impairment/Damage		Swelling Face Urticaria					

Date:05/04/05ISR Number: 4654674-0Report Type:Direct  
Age:21 YR Gender:Female I/FU:I

Company Report #CTU 247796

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300 MG	Drug Level Increased		Wellbutrin Xl	300 PS		
PARENTERAL		QAM Medication Error					
PARENTERAL		Multiple Drug Overdose					

Date:05/05/05ISR Number: 4653447-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0548370A  
Age:16 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300MG Per day	Cardio-Respiratory Arrest		Wellbutrin	PS	Glaxosmithkline	ORAL
Life-Threatening Other		Convulsion Overdose Respiratory Arrest Vomiting		No Concurrent Medication	C		

Date:05/05/05ISR Number: 4653450-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553385A  
Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	300MG Unknown	Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Cyanosis		No Concurrent			

Grand Mal Convulsion  
Muscle Twitching

Medication

C

Date:05/05/05ISR Number: 4653451-4Report Type:Expedited (15-DaCompany Report #HQWYE150815MAR05  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Crying		Wellbutrin	PS	Glaxosmithkline	ORAL
		Drug Withdrawal Syndrome		Effexor Xr	SS		ORAL
		Suicidal Ideation		Ativan	SS		
UNKNOWN		Suicide Attempt					
		Vomiting					

Date:05/05/05ISR Number: 4653453-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0556329A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pituitary Haemorrhage		Wellbutrin	PS	Glaxosmithkline	ORAL
				Topamax	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/05/05ISR Number: 4653484-8Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0045843A  
 Age:50 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 9 DAY		Coordination Abnormal	Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged 350MG per day		Drug Interaction	Saroten Retard	SS	Glaxosmithkline	ORAL
40MG per day		Muscle Twitching	Pantozol	C		ORAL
45MG per day			Remergil	C		ORAL
300MG per day			Seroquel	C		ORAL
300MG per day 2 DAY			Paracetamol	C	Glaxosmithkline	ORAL

Date:05/06/05ISR Number: 4654850-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0522799A  
 Age:76 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose 1TSP Three times per day 1 DAY		Diarrhoea	Amoxicillin	PS	Glaxosmithkline	ORAL
150MG Per day 1 DAY		Drug Ineffective				
		Nausea	Wellbutrin	SS	Glaxosmithkline	ORAL
			Paxil	SS	Glaxosmithkline	ORAL
			Nexium	C		
			Lithium	C	Glaxosmithkline	
			Synthroid	C	Glaxosmithkline	
			Toprol	C		

Date:05/06/05ISR Number: 4655170-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0536358A  
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 10MG Per day		Anxiety	Paxil	PS	Glaxosmithkline	ORAL



150MG Per day 19 DAY Chest Pain Wellbutrin SS Glaxosmithkline ORAL  
 Depression  
 Drug Withdrawal Syndrome  
 Irritability  
 Pain In Extremity  
 Suicidal Ideation

Date:05/06/05ISR Number: 4655171-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0551303A  
 Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness Neurosensory		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
		Drug Administration Error		Premarin	C		
		Meniere'S Disease		Maxalt	C		
		Otosclerosis		Flexeril	C		
		Tinnitus		Temazepam	C		
		Vertigo					

Date:05/06/05ISR Number: 4655188-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0557107A  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Flight Of Ideas		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 0 DAY		Hallucination, Auditory		Zoloft	C		
		Pharmaceutical Product		Prozac	C		
		Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/06/05ISR Number: 4655240-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0379751A  
 Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State		Zyban	PS	Glaxosmithkline	
UNKNOWN	150MG	Twice					
per day	17	DAY					
		Dysarthria					
TRANSDERMAL	15MG	Per day		Nicotine	C	Glaxosmithkline	
		Headache					
UNKNOWN		Somnolence		Zopiclone	C		
				Aspirin	C	Glaxosmithkline	
UNKNOWN	75MG	Per day		Atenolol	C		
UNKNOWN	25MG	Per day		Glyceryl Trinitrate	C	Glaxosmithkline	
				Clarithromycin	C		
UNKNOWN	500MG	Per day		Erythromycin	C	Glaxosmithkline	
UNKNOWN							

Date:05/06/05ISR Number: 4655834-5Report Type:Direct Company Report #CTU 247969  
 Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache		Zyban 150 Mg	PS		ORAL
1 PO BID							

Date:05/09/05ISR Number: 4657533-2Report Type:Direct Company Report #CTU 248037  
 Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pruritus		Bupropion Hcl 150 Mg	PS		ORAL
150 MG BID							
		Urticaria					
ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Dispensing Error		Wellbutrinxl150	PS	Glaxo Smithkline	
TABLET		Medication Error					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Anxiety Balance Disorder	Study Health	Lexiva (Fosamprenavir)	PS		ORAL
700 MG/TWICE		Depressed Mood	Professional				
PER DAY/ORAL		Dizziness Drug Interaction Headache	Other	Epzicom (Abacavir So4 + Lamivudine)	SS		ORAL
1 TABLET/PER		Insomnia					
DAY/ORAL		Irritability Photophobia		Ritonavir Capsule (Ritonavir)	SS		ORAL
100/TWICE		Tremor					
PER DAY/ORAL				Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
450 MG/PER							
DAY/ORAL				Loperamide Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/10/05ISR Number: 4657955-XReport Type:Expedited (15-DaCompany Report #2005UW06747

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Duration Drug Interaction		Seroquel	PS	Zeneca Pharmaceutical	ORAL
			Wellbutrin Sr Tablet Controlled Release	I		
			Cogentin	I		
			Trazodone	I		
			Klonopin	I		
			Trilafon	I		

Date:05/10/05ISR Number: 4658431-0Report Type:Direct Company Report #CTU 248090

Age:11 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100 MG PO BID	Duration Therapeutic Response Unexpected With Drug Substitution		Wellbutrin Sr 100 Mg Po Bid (Brand Name Only)	PS		ORAL

Date:05/10/05ISR Number: 4659314-2Report Type:Expedited (15-DaCompany Report #B0307360A

Age:38 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other 150 MG PER DAY ORAL	Duration Brain Stem Infarction Dysphagia Eyelid Ptosis Headache Hiccups Nausea Neutrophil Count Increased Photosensitivity Reaction Pupils Unequal Vomiting Wallenberg Syndrome	Foreign Literature Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

White Blood Cell Count  
Increased

Date:05/11/05ISR Number: 4658469-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0369124A  
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG per day	9 DAY	Bradycardia		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged SUBCUTANEOUS	2UNIT Single	Dyspnoea Exertional Hepatocellular Damage		Vaccine	SS	Glaxosmithkline	
dose 1UNIT per day	1 DAY	Pitting Oedema Transaminases Increased Weight Increased		Lescol Dhea	C C		ORAL

Date:05/11/05ISR Number: 4658470-XReport Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0375449A  
Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other UNKNOWN	150MG	Drug Dependence Twice No Adverse Effect		Zyban	PS	Glaxosmithkline	
per day				Waran	C	Glaxosmithkline	

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Spiriva	C
Simvastatin	C
Kalcipos-D	C
Zomig	C
Oxis	C
Pulmicort	C
Fosamax	C

Date:05/11/05ISR Number: 4658479-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0380219A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation Angina Pectoris		Zyban	PS	Glaxosmithkline	ORAL

Date:05/11/05ISR Number: 4658480-2Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0380229A  
 Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged UNKNOWN	150MG Per day 18 DAY	Hypertension Micturition Disorder Vision Blurred		Bupropion Hydrochloride	PS	Glaxosmithkline	
20MG per day				Insulin Actrapid	C		
500MG Twice per day				Metoprolol	C		
100MG per day				Omeprazole	C	Glaxosmithkline	
10MG per day				Metformin Hydrochloride	C		
				Calcium Carbaspirin	C		
				Oxazepam	C		
				Simvastatin	C		
				Aspirin	C	Glaxosmithkline	

Date:05/11/05ISR Number: 4659009-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0556917A  
Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 900MG Single Initial or Prolonged dose	2 HR	Anxiety Convulsion Fatigue Heart Rate Increased Hypoaesthesia Overdose Posture Abnormal Self-Medication		Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:05/11/05ISR Number: 4659010-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0556946A  
Age:4 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Other		Delirium Hallucination Overdose		Wellbutrin	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/11/05ISR Number: 4659013-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0557105A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Wellbutrin	PS	Glaxosmithkline	ORAL
Life-Threatening		Overdose					
Other							

Date:05/11/05ISR Number: 4659027-7Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0379971A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN	150MG	Palpitations Twice		Zyban	PS	Glaxosmithkline	
per day		Vision Blurred					
UNKNOWN				Insulin	C		
UNKNOWN				Atorvastatin	C		

Date:05/11/05ISR Number: 4659497-4Report Type:Expedited (15-DaCompany Report #2005AP000329

Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anaphylactic Reaction Asthma Drug Hypersensitivity Pyrexia	Foreign Consumer Health Professional	Apo-Citalopram  (Citalopram Hydrobromide)	PS		ORAL
60 MG; QD; PO	1 YR		Distributor Other	Buspar (Buspirone Hydrochloride)	SS		ORAL
10 MG; TID; PO	2 YR			Wellbutrin (Bupropion Hydrochloride) Avalox	SS		



Date:05/11/05ISR Number: 4660966-1Report Type:Expedited (15-DaCompany Report #KII-2005-0016466  
Age:65 YR Gender:Female I/FU:I

Outcome	PT
Life-Threatening	Alanine Aminotransferase
Hospitalization -	Increased
Initial or Prolonged	Aspartate
Other	Aminotransferase
	Increased
	Blood Alkaline
	Phosphatase Increased
	Blood Creatine
	Phosphokinase Increased
	Blood Creatinine
	Increased
	Blood Ph Decreased
	Blood Pressure
	Fluctuation
	Blood Urea Increased
	Bradycardia
	Cardio-Respiratory Arrest
	Confusional State
	Electromechanical

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dissociation Haemodynamic Instability Hypoxia Loss Of Consciousness	Report Source				
ORAL		Myoclonus	Study	Hydromorphone Hcl	PS		ORAL
ORAL		Pneumonia Sinus Tachycardia Vomiting	Health Professional Other	Bupropion (Amfebutamone)	SS		ORAL
ORAL				Citalorpan (Citalorpan)	SS		ORAL
ORAL				Trazodone (Trazodone)	SS		ORAL
ORAL				Norvasc (Amlodipine Besilate)	SS		ORAL
ORAL				Neurontin (Gabapentin)	SS		ORAL
ORAL				Propranolol	SS		ORAL
ORAL				Methocarbamol (Methocarbamol)	SS		ORAL
ORAL				Herbal Preparation	SS		ORAL
				Benzodiazepine Derivatives	SS		

Date:05/12/05ISR Number: 4660417-7Report Type:Expedited (15-DaCompany Report #2005/03/0457

Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anger		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 MON	Homicidal Ideation		Peg-Intron	SS		
INTRAVENOUS		Mood Altered Paranoia		Rebetol Vicodin Ziac	SS C C		ORAL ORAL ORAL

Date:05/12/05ISR Number: 4660419-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0554027A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Amnesia		Birth Control Pills	C		
		Aura					
		Convulsion					

Date:05/12/05ISR Number: 4660420-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0554033A

Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Post Procedural					
300MG Per day	1 YR	Haemorrhage		Klonopin	C		
				Nexium	C		
				Lasix	C	Glaxosmithkline	

Date:05/12/05ISR Number: 4660428-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0557266A

Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -		Convulsion		Effexor	C		
Initial or Prolonged		Overdose		Valium	C		

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Date:05/12/05ISR Number: 4660429-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0557281A  
 Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	100MG Per day 1 DAY	Choking		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged		Convulsion		No Concurrent Medication	C		

Date:05/12/05ISR Number: 4660430-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0557283A  
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 16 DAY		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Convulsion Head Injury		Dilantin	C		

Date:05/12/05ISR Number: 4660432-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0557413A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Ventricular Arrhythmia		Lamictal	PS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	ORAL

Date:05/12/05ISR Number: 4661722-0Report Type:Direct Company Report #CTU 248344  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Adverse Event Condition Aggravated		Generic Wellbutrin 200 Mg Bid	PS		ORAL
WELLBUTRIN		Headache					
GENERIC 200		Insomnia					
MG PO BID		Therapeutic Response					

Decreased

Date:05/12/05ISR Number: 4661779-7Report Type:Direct  
Age:46 YR Gender:Female I/FU:I

Company Report #CTU 248341

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache Migraine		Bupropriion 150 Mg Bid Po	PS		ORAL
150 MG PO BID		Pharmaceutical Product Complaint					

Date:05/13/05ISR Number: 4661403-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0548217A  
Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
474 DAY		Grand Mal Convulsion Incontinence Tongue Biting		Alcohol	C		

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Freedom Of Information (FOI) Report

Date:05/13/05ISR Number: 4661408-2Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0554466A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly	150MG Per day		Hypospadias	Wellbutrin	PS	Glaxosmithkline	
	.137MG Per day		Microtia	Synthroid	C	Glaxosmithkline	
	10MG Eight times per day			Domperidone	C		

Date:05/13/05ISR Number: 4661415-XReport Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0557615A

Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day		Angina Pectoris	Zyban	PS	Glaxosmithkline	ORAL
			Back Pain				
			Hypoaesthesia				

Date:05/13/05ISR Number: 4661417-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0557786A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Anger	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Convulsion	Alcohol	C		
			Intentional Self-Injury				

Date:05/13/05ISR Number: 4661454-9Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0379324A

Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Depression	Zyban	PS	Glaxosmithkline	ORAL

Initial or Prolonged Mood Altered  
Rash

Date:05/13/05ISR Number: 4661457-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0380052A  
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anhedonia		Zyban	PS	Glaxosmithkline	ORAL
150MG per day	27 DAY	Depressed Mood		Niquitin Cq	C	Glaxosmithkline	
TRANSDERMAL		Insomnia					
		Suicidal Ideation					

Date:05/13/05ISR Number: 4661458-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0380140A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Zyban	PS	Glaxosmithkline	
UNKNOWN		Depression					
		Psychiatric Symptom					
		Suicidal Ideation					

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Freedom Of Information (FOI) Report

Date:05/13/05ISR Number: 4661459-8Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0380489A  
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cold Sweat		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day		Dizziness		No Concurrent			
		Dry Mouth		Medication	C		
		Gait Disturbance					
		Hallucination, Auditory					
		Hallucination, Visual					
		Insomnia					
		Malaise					
		Tremor					

Date:05/13/05ISR Number: 4662497-1Report Type:Direct Company Report #CTU 248490  
 Age:16 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Diabetes Mellitus		Risperdol 2 Mg.	PS		ORAL
2 MG.		Non-Insulin-Dependent					
Intervention to				Wellbutrin	SS		
B.I.D. ORAL				Depakote	C		
Prevent Permanent				Adderal	C		
Impairment/Damage				Paxil	C		

Date:05/13/05ISR Number: 4662499-5Report Type:Direct Company Report #CTU 248475  
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Bupropion			
		Arthralgia		(Budeprion)	150		
		Drug Ineffective		Mg Sr Unk	PS		ORAL
1 PER DAY		Fatigue					
ORAL		Feeling Cold					
		Fluid Retention					



Headache  
 Pain In Extremity  
 Speech Disorder  
 Vision Blurred  
 Weight Increased

Date:05/13/05ISR Number: 4663159-7Report Type:Expedited (15-DaCompany Report #S05-USA-02393-01  
 Age:18 YR Gender:Female I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
20 MG QD PO		Convulsion Discomfort	Health Professional	Lexapro (Escitalopram)	PS		ORAL
10 MG QD PO		Fatigue Hypoaesthesia		Lexapro (Escitalopram)	SS		ORAL
		Joint Contracture Paraesthesia Suicide Attempt Tremor		Wellbutrin (Bupropion Hydrochloride)	SS		
900 MG ONCE PO				Wellbutrin (Bupropion Hydrochloride)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/16/05ISR Number: 4662672-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0508446A  
 Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Twice Initial or Prolonged per day	6	MON		Wellbutrin	PS	Glaxosmithkline	ORAL
Other			Loss Of Consciousness Night Sweats	Pepcid Vicodin Flexeril Nasonex Klonopin	C C C C C		
1MG Twice per day							

Date:05/16/05ISR Number: 4662680-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0554507A  
 Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Three times per day			Agitation Anger	Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
150MG Three times per day			Asthenia Depression	Wellbutrin Sr	SS	Glaxosmithkline	ORAL
UNKNOWN 600MG Five times per day			Disturbance In Attention Drug Ineffective	Clonazepam	C		
UNKNOWN 50MG At night			Irritability Mood Swings	Neurontin	C		
UNKNOWN			Pathological Gambling	Atenolol	C		
UNKNOWN			Sleep Disorder Suicidal Ideation	Trazodone	C		

Date:05/16/05ISR Number: 4662685-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0557263A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG Three Initial or Prolonged times per day		Colectomy Partial Coronary Artery Surgery		Wellbutrin	PS	Glaxosmithkline	ORAL
3 MON		Diarrhoea		Paxil	SS	Glaxosmithkline	ORAL
		Diverticulitis		Atenolol	C		
		Drug Ineffective		Prevacid	C		
		Myocardial Infarction		Lotrel	C		
		Pneumonia		Lipitor	C		

Date:05/16/05ISR Number: 4662690-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0557567A

Age:17 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG Per day 3 WK		Anger		Wellbutrin	PS	Glaxosmithkline	ORAL
		Depression		Seroquel	C		
		Drug Ineffective		Paxil Cr	C	Glaxosmithkline	
		Paranoia		Yasmin	C		
		Suicidal Ideation		Multivitamin	C		

Date:05/16/05ISR Number: 4662694-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0557761A

Age: Gender:Female I/FU:I

Outcome	PT
Disability	Drug Ineffective Pruritus Rash

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tremor Urticaria Urticaria Generalised					
4 YR				Lamictal	PS	Glaxosmithkline	ORAL
2 WK				Wellbutrin	SS	Glaxosmithkline	ORAL
				Lexapro	C		
				Naproxen	C		

Date:05/16/05ISR Number: 4663871-XReport Type:Expedited (15-DaCompany Report #2005069676  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 80 MG (80 MG, Initial or Prolonged 1 IN 1 D) Other		Coma Fatigue Hemiplegia Loss Of Consciousness Metabolic Encephalopathy Nasopharyngitis Pneumonia Aspiration Unresponsive To Pain Stimuli Vomiting	Health Professional	Geodon (Ziprasidone)	PS		
				Sudafed (Pseudoephedrine Hydrochloride)(Pseud oephedrine)	SS		
				Metformin (Metformin)	SS		
				Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		
				Olanzapine (Olanzapine)	C		
				Morphine (Morphine)	C		

Date:05/16/05ISR Number: 4664358-0Report Type:Direct Company Report #CTU 248728  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 200 MG QD PO		Malaise Pharmaceutical Product Complaint		Bupropion Sr 200 Mg Global	PS	Global	
				Humalog U100	C		

Vomiting Projectile

Cozaar	C
Digoxin	C
Plavix	C
Ms Contin	C
Relafen	C
Bumex	C
Verapamil	C
Clonazepam	C

Date:05/16/05ISR Number: 4664544-XReport Type:Direct  
Age:12 YR Gender:Male I/FU:I

Company Report #CTU 248692

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression Pharmaceutical Product Complaint		Generic Wellbutrin	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/17/05ISR Number: 4663498-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0552663A

Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	450MG Per day	Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	450MG Twice	Fatigue		Eskalith	C	Glaxosmithkline	
	per day	Tongue Biting					
	25MG At night			Seroquel	C		
	10MG At night			Ambien	C		
	30MG Unknown			Prevacid	C		
	180MG Unknown			Allegra	C		
	.025MG			Synthroid	C	Glaxosmithkline	
	Unknown						
	145MG At			Tricor	C		
	night						

Date:05/17/05ISR Number: 4663539-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0380426A

Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Unknown 16 DAY	Dermatitis Allergic		Zyban	PS	Glaxosmithkline	ORAL
		Oedema Peripheral					
		Periorbital Oedema					

Date:05/17/05ISR Number: 4666432-1Report Type:Expedited (15-DaCompany Report #S05-USA-00523-01

Age:60 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Acute Sinusitis

Initial or Prolonged

Anaemia Of Chronic  
Disease  
Asthenia  
Back Pain  
Bacterial Sepsis  
Blood Glucose Increased  
Catheter Sepsis  
Chest Pain  
Chest Wall Pain  
Clostridium Colitis  
Coma  
Confusional State  
Depression  
Disease Recurrence  
Drug Interaction  
Electrolyte Imbalance  
Fall  
Fluid Overload  
Hypercapnia  
Hyperkalaemia  
Hypotension  
Hypoxia  
Metabolic Acidosis  
Musculoskeletal  
Discomfort  
Neuropathy  
Pneumonia Bacterial  
Pulmonary Hypertension  
Renal Failure Acute  
Respiratory Failure  
Staphylococcal Infection

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Freedom Of Information (FOI) Report

Dose	Duration	Stenotrophomonas Infection Syncope	Report Source	Product	Role	Manufacturer	Route
20 MG QD PO		Tremor Urinary Tract Infection	Health Professional	Lexapro (Escitalopram)	PS		ORAL
30 MG QD PO				Lexapro (Escitalopram)	SS		ORAL
10 MG QD PO				Lexapro (Escitalopram)	SS		ORAL
20 MG QD PO				Lexapro (Escitalopram)	SS		ORAL
30 MG QD PO				Lexapro (Escitalopram)	SS		ORAL
300 MG QD PO				Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
150 MG BID				Wellbutrin (Bupropion Hydrochloride)	SS		
300 MG QD PO				Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
150 MG QD PO				Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
				Atenolol	C		
				Potassium	C		
				Zocor	C		
				Enteric Coated Aspirin	C		
				Nexium(Esomeprazole)	C		
				Aricept(Donepezil Hydrochloride)	C		
				Duragesic(Fentanyl)	C		
				Hydrocodone	C		
				Centrum Silver	C		
				Magnesium	C		



Vitamin B12	C
Diazepam	C
Altace(Ramipril)	C
Demadex(Torasemide)	C
Skelaxin(Metaxalone)	C
Isordil(Isosorside Dinitrate)	C
Glucophage (Metformin)	C
Acetazolamide	C
Xalatan (Latanoprost)	C
Oxycontin(Oxycodone Hydrochloride)	C
Miconazole	C
Humulin	C
Astelin(Azelastine Hydrochloride)	C
Meclizine	C
Norvasc(Amlodipine Besilate)	C
Rocephin(Ceftriaxone Sodium)	C
Tobramycin	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydralazine	C
Protonix	C
Valium (Diazepam)	C
Lorcet Plus	C
Diamox	
(Acetazolamide)	C
Avandia	
(Rosiglitazone	
Maleate)	C
Nitrol (Glyceryl	
Trinitrate)	C

Date:05/18/05ISR Number: 4664550-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0548149A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	300MG Per day	Contusion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Grand Mal Convulsion					
		Laceration					

Date:05/18/05ISR Number: 4664570-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0558980A  
Age:18 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2 DAY	Death		Wellbutrin	PS	Glaxosmithkline	ORAL
Life-Threatening		Status Epilepticus					
Hospitalization -							
Initial or Prolonged							
Disability							
Other							

Date:05/18/05ISR Number: 4664588-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0380740A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other 300MG per day		Analgesic Effect		Zyban	PS	Glaxosmithkline	ORAL
		Drug Abuser		Imiject	SS	Glaxosmithkline	
SUBCUTANEOUS	4INJ per day	Therapeutic Response Unexpected					

Date:05/18/05ISR Number: 4664606-7Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0381313A  
 Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1TAB Twice per day	4 DAY	Coordination Abnormal Depressed Level Of Consciousness Impaired Driving Ability		Zyban  No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

Date:05/18/05ISR Number: 4664610-9Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0381415A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN		Overdose Suicidal Ideation Suicide Attempt		Zyban Mogadon	PS C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/18/05ISR Number: 4665271-5Report Type:Expedited (15-DaCompany Report #US-MERCK-0505USA01221

Age:82 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Accident		Vioxx	PS	Merck & Co., Inc	ORAL
		Hypertension		Wellbutrin Sr	SS		ORAL
4 YR							
		Hypotension		Wellbutrin Sr	SS		ORAL
UNKNOWN				Norvasc	C		
				Dyrenium	C		
UNKNOWN							
				Clonazepam	C		
UNKNOWN							
				Levoxyl	C		
UNKNOWN							
				Macrobid	C		
UNKNOWN							
				Prevacid	C		
UNKNOWN							

Date:05/18/05ISR Number: 4665403-9Report Type:Direct

Company Report #CTU 248958

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Blood Pressure Increased		Cephalexin 250 Mg			
Other		Diarrhoea		Capsules (Teva)	PS	Teva	ORAL
250 MG Q ID							
PO [ONLY 2		Headache					
DOSES]		Heart Rate Increased					
		Muscle Spasms		Bupropion Sr 150 Mg			
150 MG DAILY		Oral Intake Reduced		Tablet (Watson)	SS	Watson	ORAL
PO							
[DISCHARGED		Respiratory Rate					
		Increased					
ON IT]		Tremor					
		Vision Blurred					

Date:05/18/05ISR Number: 4666447-3Report Type:Expedited (15-DaCompany Report #S05-USA-01759-01  
Age:36 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Diplopia	Health	Lexapro			
Initial or Prolonged	Drug Interaction	Professional	(Escitalopram)	PS		ORAL
40 MG QD PO						
	Nausea	Company	Wellbutrin			
	Vertigo	Representative	(Bupropion			
	Viith Nerve Paralysis		Hydrochloride)	SS		
150 MG QD						
	Vomiting					

Date:05/18/05ISR Number: 4668506-8Report Type:Expedited (15-DaCompany Report #B0379971A  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Blindness	Foreign	Zyban Tablet-Zyban			
	Palpitations	Consumer	(Bupropion			
150 MG/TWICE			Hydrochloride)	PS		
PER DAY						
			Insulin	C		
			Atorvastatin Calcium	C		

Date:05/18/05ISR Number: 4668645-1Report Type:Expedited (15-DaCompany Report #A0556337A  
Age:32 YR Gender:Female I/FU:I

Outcome	PT
Other	Condition Aggravated
	Dizziness
	Drug Withdrawal Syndrome

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
25 MG/ PER		Heart Rate Increased Hyperhidrosis Night Sweats	Consumer	Paxil Cr Tablet-Controlled Release (Paroxetine Hydrochloride)	PS		ORAL
ORAL/ ORAL		Paraesthesia Psoriasis					
300 MG/ ORAL				Bupropion Hydrochloride Tablet (Bupropion Hydrochloride)	SS		ORAL

Date:05/18/05ISR Number: 4668665-7Report Type:Expedited (15-DaCompany Report #A0553988A  
Age:17 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage	300 MG/ ORAL		Overdose Vomiting	Health Professional Company Representative	Wellbutrin Xl Tablet-Extended Release (Bupropion Hydrochloride)	PS		ORAL

Date:05/18/05ISR Number: 4668666-9Report Type:Expedited (15-DaCompany Report #A0549034A  
Age:64 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	PER DAY/ ORAL		Rectal Haemorrhage Rectal Polyp Stool Analysis Abnormal	Health Professional	Wellbutrin Xl Tablet-Extended Release (Bupropion Hydrochloride)	PS		ORAL
					Sertraline Hydrochloride	C		
					Thyroxine Sodium	C		

Lovastatin	C
Quinapril	
Hydrochloride	C
Oestradiol	C
Multivitamin	C

Date:05/19/05ISR Number: 4666079-7Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20050307425  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction		Valtrex	PS	Glaxosmithkline	ORAL
		Throat Tightness		Wellbutrin	SS	Glaxosmithkline	ORAL
				Tylenol Cold + Flu	SS		ORAL

Date:05/19/05ISR Number: 4666103-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0381009A  
 Age:34 YR Gender:Female I/FU:F

Outcome	PT
Disability	Anxiety
	Dry Mouth
	Ill-Defined Disorder
	Logorrhoea

Freedom Of Information (FOI) Report

Psychomotor Hyperactivity

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day			Zyban	PS	Glaxosmithkline	ORAL
UNKNOWN			Marvelon	C		
UNKNOWN required	200MCG As		Salbutamol	C	Glaxosmithkline	
UNKNOWN per day	200MCG Twice		Beclometasone	C	Glaxosmithkline	

Date:05/19/05ISR Number: 4666108-0Report Type:Expedited (15-DaCompany Report #CH-GLAXOSMITHKLINE-B0381417A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Other		Erythema Systemic Lupus Erythematosus		Zyban Spiriva	PS SS	Glaxosmithkline	

Date:05/20/05ISR Number: 4667745-XReport Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0557615A  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Per day		Angina Pectoris Back Pain Hypoaesthesia		Zyban	PS	Glaxosmithkline	ORAL

Date:05/20/05ISR Number: 4667750-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0558376A  
Age:26 YR Gender:Female I/FU:I



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day		Overdose		Crack Cocaine	C		
		Stress		Valium	C		

Date:05/20/05ISR Number: 4667752-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0558525A

Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	5 WK	Feeling Abnormal					
		Suicidal Ideation					

Date:05/20/05ISR Number: 4667753-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0558531A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
		Head Injury		No Concurrent			
		Skull Fracture		Medication	C		

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Date:05/20/05ISR Number: 4667754-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0558535A  
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
1TAB Per day	30 MON	Dyspnoea		Trazodone	C		
		Hallucination					
		Nausea					
		Overdose					
		Tremor					
		Vomiting					

Date:05/20/05ISR Number: 4667756-4Report Type:Expedited (15-DaCompany Report #BPC-SR-033  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Congenital Anomaly		Drug Exposure During		Bupropion	PS	Glaxosmithkline	
150MG Per day		Pregnancy					
		Failure To Thrive					
		Laryngomalacia					
		Stridor					

Date:05/20/05ISR Number: 4667757-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0558836A  
 Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Overdose		Wellbutrin	PS	Glaxosmithkline	ORAL
1 DAY		Suicide Attempt		Provigil	SS		ORAL
Initial or Prolonged		Tachycardia		Ambien	SS		ORAL
60MG Single		Vomiting					
dose	1 DAY						

Date:05/20/05ISR Number: 4667790-4Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0381597A  
 Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	8 DAY	Dizziness		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged	40MG Per day	Migraine		Inderal	C		
UNKNOWN		Mydriasis					
		Sensory Loss					

Date:05/23/05ISR Number: 4668948-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0555404A

Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG Per day	Completed Suicide		Wellbutrin	PS	Glaxosmithkline	
Life-Threatening		Cough		Tylenol	SS	Glaxosmithkline	
UNKNOWN							
Hospitalization -		Diarrhoea		Alcohol	SS		
UNKNOWN							
Initial or Prolonged		Dyspnoea		Nexium	C		
Disability		Headache		Celebrex	C		
		Hepatic Failure		Lipitor	C		
		Insomnia					
		Metabolic Acidosis					
		Overdose					
		Pneumonia					
		Pneumonia Aspiration					
		Vomiting					

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Date:05/23/05ISR Number: 4670055-8Report Type:Expedited (15-DaCompany Report #IMP\_0854\_2005

Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Literature	Bupropion Sr	PS		ORAL
Other		Condition Aggravated	Health				
100 MG DAILY		Depression	Professional	Bupropion Sr	SS		ORAL
PO		Dizziness		Bupropion Sr	SS		ORAL
100 MG QAM PO		Inappropriate		Bupropion Sr	SS		ORAL
200 MG BID PO		Antidiuretic Hormone Secretion		Divalproex Sodium	SS		
100 MG QAM PO				Oxycarbazepine	SS		ORAL
900 MG QDAY		Irritable Bowel Syndrome					
PO		Orthostatic Hypotension					
		Tremor					

Date:05/23/05ISR Number: 4670060-1Report Type:Expedited (15-DaCompany Report #IMP\_0847\_2005

Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Bupropion	PS		ORAL
Other		Crying					
1 TAB QDAY PO		Feeling Abnormal					
		Paranoia					
		Pharmaceutical Product Complaint					
		Suicidal Ideation					

Date:05/23/05ISR Number: 4673228-3Report Type:Direct

Company Report #CTU 249268

Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Dyspnoea Wellbutrin PS  
150 MG ONCE A  
Initial or Prolonged Heart Rate Increased  
DAY

Date:05/23/05ISR Number: 4673284-2Report Type:Direct Company Report #CTU 249294  
Age:41 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Palpitations		Bupropion Sr	PS		ORAL
150MG PO BID						
Initial or Prolonged			Rocephin	C		
			Prednisone	C		
			Zithromax	C		
			Lovenox	C		
			Nicotine Patch	C		
			Combivent	C		
			Insulin	C		

Date:05/23/05ISR Number: 4673736-5Report Type:Expedited (15-DaCompany Report #S05-USA-01454-01  
Age:27 YR Gender:Female I/FU:F

Outcome	PT
Other	Convulsion
	Depressed Level Of Consciousness
	Disorientation
	Fall
	Head Injury
	Hypersomnia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tongue Biting Tremor	Health Professional	Lexapro (Escitalopram)	PS		ORAL
30 MG QD PO			Company Representative	Welbutrin (Bupropion Hydrochloride)	SS		
300 MG QAM				Lexapro (Escitalopram)	SS		ORAL
20 MG QD PO							

Date:05/24/05ISR Number: 4672555-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0541125A  
Age:83 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	100MG Twice		Drug Dispensing Error		Wellbutrin	PS	Glaxosmithkline	ORAL
per day		2 WK	Drug Ineffective					
			Hallucination		Lithium	C	Glaxosmithkline	
			Medication Error		Ativan	C		
			Paranoia					
			Pharmaceutical Product Complaint					

Date:05/24/05ISR Number: 4672564-4Report Type:Expedited (15-DaCompany Report #S05-USA-02393-01  
Age:18 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Initial or Prolonged			Discomfort		Lexapro	SS		ORAL
20MG Per day		15 DAY	Fatigue					
			Hypoaesthesia					
			Ill-Defined Disorder					
			Intentional Misuse					
			Muscle Contractions					
			Involuntary					

Paraesthesia  
Tremor

Date:05/24/05ISR Number: 4672566-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0558989A  
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Depression					
per day	2 YR	Dysphagia		Unknown Medication	C		
		Oesophageal Carcinoma		Respiradol	C		

Date:05/24/05ISR Number: 4672568-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0559383A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Unknown		Depression		Eskalith Cr	C	Glaxosmithkline	ORAL
450MG Twice		Dyskinesia					
per day		Feeling Abnormal		Sleeping Pill	C		
		Suicidal Ideation					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/24/05ISR Number: 4672575-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0559750A  
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Anger		Eskalith	PS	Glaxosmithkline	ORAL
450MG Twice							
Hospitalization -		Crying					
per day	3 WK						
Initial or Prolonged		Depression		Paxil	SS	Glaxosmithkline	ORAL
20MG Per day	9 YR						
Disability		Poisoning		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
150MG Per day	2 WK						
Other		Psychotic Disorder		Seroquel	SS		
UNKNOWN							
		Suicide Attempt		Risperdal	C		
				Klonopin	C		

Date:05/24/05ISR Number: 4672599-1Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0380967A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Disorientation		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Fall					
		Malaise					
		Pneumonia					
		Sleep Disorder					

Date:05/24/05ISR Number: 4672600-5Report Type:Expedited (15-DaCompany Report #CH-GLAXOSMITHKLINE-B0381417A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Cutaneous Lupus		Zyban	PS	Glaxosmithkline	
Other		Erythematosus		Spiriva	SS		
		Erythema					

Date:05/24/05ISR Number: 4674327-2Report Type:Expedited (15-DaCompany Report #US-2005-003875  
 Age:42 YR Gender:Female I/FU:F



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Delusion	Consumer	Betaseron(Interferon Beta - 1b) Injection, 250ug	PS		
SUBCUTANEOUS	SUBCUTANEOUS			Wellbutrin (Bupropion Hydrochloride)	SS		

Date:05/24/05ISR Number: 4674399-5Report Type:Expedited (15-DaCompany Report #US-2005-003875  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Delusion	Consumer	Betaseron (Interferon Beta-1b) Injection, 250 Ug	PS		
SUBCUTANEOUS	SUBCUTANEOUS			Wellbutrin (Bupropion Hydrochloride)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/25/05ISR Number: 4673418-XReport Type:Expedited (15-DaCompany Report #13611

Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deficiency Anaemia		Bupropion	PS	Glaxosmithkline	
		Gastrointestinal		Prenatal Vitamins	C		
		Infection		Calcium	C		
		Polyhydramnios					

Date:05/25/05ISR Number: 4673421-XReport Type:Expedited (15-DaCompany Report #MCN # S05-USA-01759-01

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Diplopia		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day							
Initial or Prolonged		Drug Interaction		Lexapro	SS		ORAL
40MG Per day	357 DAY						
		Facial Palsy					
		Nausea					
		Vertigo					
		Vomiting					

Date:05/25/05ISR Number: 4673423-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0558943A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Anger					
		Emotional Distress					
		Suicide Attempt					

Date:05/25/05ISR Number: 4673424-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0558956A

Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Blood Alcohol Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							

Death

Lexapro	C	
Lamotrigine	C	Glaxosmithkline
Minocycline	C	
Alcohol	C	

Date:05/25/05ISR Number: 4673425-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0559188A

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Exposure During Pregnancy Jaundice Neonatal		Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	

Date:05/25/05ISR Number: 4673459-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0381069A

Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression		Zyban	PS	Glaxosmithkline	ORAL
150MG per day	11 DAY	Headache		Microval	C		ORAL
30MCG per day	7 MON	Hyperhidrosis		Vitamin C	C	Glaxosmithkline	ORAL
500MG per day		Pruritus Salivary Hypersecretion Skin Disorder		Nurofen	C	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/25/05ISR Number: 4673466-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0381911A  
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 450MG per day	15 DAY	Amnesia		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Convulsive Threshold Lowered Dyskinesia Grand Mal Convulsion Hypertonia Loss Of Consciousness Malaise Postictal State					

Date:05/25/05ISR Number: 4674012-7Report Type:Expedited (15-DaCompany Report #IMP\_0861\_2005  
 Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 400 MG ONCE		Aggression	Foreign	Bupropion Sr	PS		ORAL
Initial or Prolonged PO		Body Temperature Increased	Literature Health	Bupropion Sr	SS		ORAL
150 MG QDAY PO	2 WK	Depression	Professional				
300 MG QDAY PO		Intentional Misuse	Other	Bupropion Sr	SS		ORAL
PO	2 DAY	Irritability Paranoia		Midazolam	SS		ORAL
105 MG ONCE PO		Persecutory Delusion					
7.5 MG NIGHTLY PO		Psychomotor Agitation Psychotic Disorder		Midazolam	SS		ORAL
		Respiratory Rate Increased Restlessness Somnolence					

Suicide Attempt  
Thinking Abnormal

Date:05/25/05ISR Number: 4674799-3Report Type:Expedited (15-DaCompany Report #231577K05USA  
Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anxiety Depression	Consumer Health	Rebif (Interferon Beta)	PS		
SUBCUTANEOUS	44 MCG, 3 IN	Faecal Incontinence	Professional				
1 WEEKS, SUBCUTANEOUS		Psychotic Disorder					
		Self-Injurious Ideation Suicidal Ideation		Wellbutrin (Bupropion Hydrochloride)	SS		
75 MG, 1 IN 1 DAYS, ORAL				Effexor (Venlafaxine Hydrochloride)	SS		ORAL

Date:05/26/05ISR Number: 4674645-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0559557A  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hypothyroidism		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	8 MON	Myopathy		Lexapro	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/26/05ISR Number: 4674654-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0560003A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Wellbutrin	PS	Glaxosmithkline	

Date:05/26/05ISR Number: 4674656-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0560118A

Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Dizziness Multiple Sclerosis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/26/05ISR Number: 4674672-0Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0381410A

Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Unknown 8 DAY	Agitation		Zyban	PS	Glaxosmithkline	ORAL
TRANSDERMAL		Headache		Nicotine	C	Glaxosmithkline	
6MG Single		30 DAY		Nicotel	C		ORAL
dose		Personality Change					
		Psychotic Disorder					

Date:05/26/05ISR Number: 4674680-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0382285A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	9 DAY	Anorexia		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged	UNKNOWN	Dry Mouth		Digoxin	C	Glaxosmithkline	
UNKNOWN	250MG Per day	Dry Throat		Zestril	C		
UNKNOWN	20MG per day						

UNKNOWN	20MG per day	Eating Disorder	Lipex	C	
		Headache	Aspirin	C	Glaxosmithkline
UNKNOWN	300MG See	Hyperhidrosis			
dosage text		Influenza Like Illness	Diaformin	C	
UNKNOWN	500MG Twice	Malaise			
per day		Nausea			
		Palpitations			
		Pyrexia			
		Somnolence			
		Vomiting			

Date:05/26/05ISR Number: 4677370-2Report Type:Direct Company Report #CTU 249605  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agoraphobia		Bupropin (Generic)			
Other		Anxiety		150 Mg One Bid	PS		
		Condition Aggravated					
		Depression					
		Therapeutic Response					
		Unexpected With Drug					
		Substitution					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4675920-3Report Type:Periodic  
Age:54 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0548813A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	2 WK	Palmar Erythema	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Pruritus	Caltrate	C	Glaxosmithkline	
			Rash Generalised	Asa	C	Glaxosmithkline	
				Vitamin C	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4675921-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548834A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Tinnitus	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Zoloft	C		

Date:05/27/05ISR Number: 4675922-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548839A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	6 MON	Insomnia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Restlessness	Lexapro	C		
				Vitamin	C		
				Green Tea	C		
				Saw Palmetto	C		
				Vitamin E	C		
				Garlic	C		

Date:05/27/05ISR Number: 4675923-9Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549008A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Twice		Agitation	Wellbutrin Xl	PS	Glaxosmithkline	ORAL



per day 16 MON  
Anxiety  
Circadian Rhythm Sleep  
Disorder  
Delusion  
Impulsive Behaviour  
Paranoia  
Social Phobia  
Weight Decreased

No Concurrent  
Medication C

Date:05/27/05ISR Number: 4675925-2Report Type:Periodic  
Age:17 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0549009A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Fentanyl	SS		
				Anesthesia	SS		
				Concerta	SS		
				Ortho Cyclen	C		
				Tapazole	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4675927-6Report Type:Periodic  
Age:53 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549011A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Excedrin	SS		
				Celebrex	SS		
UNKNOWN							

Date:05/27/05ISR Number: 4675928-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554240A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	6	WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:05/27/05ISR Number: 4675929-XReport Type:Periodic  
Age:77 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0549027A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:05/27/05ISR Number: 4675930-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554249A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
1 DAY				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Neurontin	C		

Date:05/27/05ISR Number: 4675931-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549028A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2	WK		Birth Control	C		

Date:05/27/05ISR Number: 4675932-XReport Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554250A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	13	DAY		Prevacid	C		
		Tinnitus					

Date:05/27/05ISR Number: 4675933-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549040A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lethargy		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3	MON		No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4675934-3Report Type:Periodic  
Age:52 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0554283A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day 5 DAY Initial or Prolonged		Amnesia Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4675935-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549108A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Alanine Aminotransferase Increased Aspartate Aminotransferase Increased Hepatic Enzyme Abnormal		Wellbutrin Xl Alcohol	PS C	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4675936-7Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0554368A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day 6 MON		Movement Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4675937-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549110A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4675938-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554397A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK	Rash		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4675939-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0549112A  
 Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysuria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
6 MON		Urine Output Decreased					

Date:05/27/05ISR Number: 4675940-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0554414A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK	Tinnitus					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4675941-0Report Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549227A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3	WK		Ortho Evra	C		

Date:05/27/05ISR Number: 4675942-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0554455A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4675943-4Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549265A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1	WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Ear Pain		Clonazepam	C		
		Headache		Estradiol	C		
				Tylenol	C	Glaxosmithkline	
				Fosamax	C		

Date:05/27/05ISR Number: 4675944-6Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0554474A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Libido Decreased					

Date:05/27/05ISR Number: 4675945-8Report Type:Periodic  
Age:14 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549269A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Abnormal Behaviour		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
5		DAY						

Date:05/27/05ISR Number: 4675946-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0554511A  
 Age:41 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Complex Partial Seizures		Wellbutrin	PS	Glaxosmithkline	ORAL
	300MG Per day	1 YR	Disorientation		Paxil	C	Glaxosmithkline	
4		YR			Synthroid	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4675947-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0549272A  
 Age:55 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Fatigue		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	100MG Per day	1 MON	Therapeutic Response Decreased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4675948-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554561A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Halitosis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Three							
times per day							

Date:05/27/05ISR Number: 4675949-5Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549412A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 1	WK	Confusional State		Multivitamins	C		
		Somnolence					
		Thinking Abnormal					

Date:05/27/05ISR Number: 4675950-1Report Type:Periodic  
Age:35 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554578A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Derealisation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2 DAY		Drug Screen Positive		Trazodone	C		
UNKNOWN				Citalopram	C		
UNKNOWN							

Date:05/27/05ISR Number: 4675951-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549416A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alanine Aminotransferase		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other		Increased					
150MG Per day							



Aspartate  
Aminotransferase  
Increased

Date:05/27/05ISR Number: 4675952-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0554599A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Platelet Count Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4675953-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0549419A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				No Concurrent Medication	C		

Date:05/27/05ISR Number: 4675954-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0554813A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bruxism		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4675955-0Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549424A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	4 DAY	Influenza		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Myalgia Painful Respiration Rash		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4675956-2Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554816A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day 2 WK	Vaginal Haemorrhage		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Effexor	C		
				Lexapro	C		

Date:05/27/05ISR Number: 4675957-4Report Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549426A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	3 WK	Swelling Face		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Vicodin	C		

Date:05/27/05ISR Number: 4675958-6Report Type:Periodic  
Age:83 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554835A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day 1 YR	Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Drug Interaction		Lipitor	SS		
				Vitamins	C		

Date:05/27/05ISR Number: 4675959-8Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549440A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hot Flush		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR	Hyperhidrosis		Birth Control	C		

Date:05/27/05ISR Number: 4675960-4Report Type:Periodic  
Age:49 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554839A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Arthralgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	6 MON						

Date:05/27/05ISR Number: 4675961-6Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549445A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK	Heart Rate Increased		Darvon	C		
		Palpitations					
		Panic Attack					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4675962-8Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554863A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	Vaginal Burning Sensation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4675963-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549460A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Unknown	Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4675964-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554864A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Unknown	Burning Sensation		Wellbutrin Xl	PS	Glaxosmithkline	OTHER

Date:05/27/05ISR Number: 4675965-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549502A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Unknown	Anorexia Initial Insomnia Middle Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4675966-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554886A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Pharmaceutical Product  
Complaint

Date:05/27/05ISR Number: 4675967-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549613A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	

Date:05/27/05ISR Number: 4675968-9Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555015A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Bone Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	24 DAY	Mobility Decreased		Trileptal	C		
		Myalgia		Prevacid	C		
		Swelling					

Date:05/27/05ISR Number: 4675969-0Report Type:Periodic  
Age:66 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549629A

Outcome	PT
	Abnormal Dreams
	Dry Mouth

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Thinking Abnormal

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day	10 WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Naproxen	C		
			Cardura	C		
			Prilosec	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4675970-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0555029A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	5 WK			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Photosensitivity Reaction					
		Rash		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4675971-9Report Type:Periodic Company Report #CN050048  
 Age:16 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 YR			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Medication Error					
		Pharmaceutical Product Complaint		Concerta	C		

Date:05/27/05ISR Number: 4675972-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0555030A  
 Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY						
		Hallucination					
		Tremor		Depakote	C		
				Celexa	C		
				Celebrex	C		
				Plavix	C		

Vitamin E	C	
Vitamin C	C	Glaxosmithkline
Avandia	C	Glaxosmithkline
Albuterol	C	Glaxosmithkline

Date:05/27/05ISR Number: 4675973-2Report Type:Periodic  
 Age:65 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549637A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hyperhidrosis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Vitamins	C		

Date:05/27/05ISR Number: 4675974-4Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0555034A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4675975-6Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549644A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	11 DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dysphagia					
		Gingival Pain		Mobic	C		
		Rash		Ultram	C		
		Urticaria					

Date:05/27/05ISR Number: 4675976-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555038A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion					

Date:05/27/05ISR Number: 4675977-XReport Type:Periodic  
Age:37 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549652A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	15 DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Disorientation					
		Dizziness		No Concurrent Medication	C		
		Insomnia					
		Nausea					

Date:05/27/05ISR Number: 4675978-1Report Type:Periodic  
Age:26 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555050A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Headache					
		Malaise		Alesse	C		
		Sinus Headache					



Date:05/27/05ISR Number: 4675979-3Report Type:Periodic  
Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549654A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY	Paraesthesia Paraesthesia Oral		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4675980-XReport Type:Periodic  
Age:59 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555224A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Paraesthesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day	1 MON			Zoloft Ativan	C C		

Date:05/27/05ISR Number: 4675981-1Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0549669A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal Insomnia		Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4675982-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555229A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	1 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
		Lethargy		Atenolol	C		
				Temaze	C		
				Mevacor	C		

Date:05/27/05ISR Number: 4675983-5Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0549671A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropion Hydrochloride	PS	Glaxosmithkline	ORAL
		Feeling Abnormal					
		Insomnia					
		Pruritus					

Date:05/27/05ISR Number: 4675984-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555234A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	450MG Per day	8 WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Therapeutic Response		Lexapro	C		
		Decreased					

Date:05/27/05ISR Number: 4675985-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549717A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Adverse Event					

Date:05/27/05ISR Number: 4675986-0Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555259A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthritis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
MON							

Date:05/27/05ISR Number: 4675987-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0549722A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	

Date:05/27/05ISR Number: 4675988-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0555370A  
 Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Wellbutrin	PS	Glaxosmithkline	ORAL
4 WK		Pruritus		Cymbalta	C		
		Rash		Allegra D	C		
		Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4675989-6Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549832A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	4 WK			Amaryl	C		
				Prednisone	C		
				Hctz	C		
				Zyrtec	C	Glaxosmithkline	
				Furosemide	C	Glaxosmithkline	
				Ambien	C		
				Spiroonolactone	C		
				Hydrocodone	C		
				Cortisone	C		

Date:05/27/05ISR Number: 4675990-2Report Type:Periodic  
 Age:20 YR Gender:Male I/FU:F

Company Report #S05-USA-02072-01

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Lexapro	SS		
450MG Per day	3 WK						

Date:05/27/05ISR Number: 4675991-4Report Type:Periodic  
 Age:63 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549836A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Jittery		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Hyperhidrosis		Amiodarone	C		
		Insomnia		Aspirin	C	Glaxosmithkline	
		Nervousness		Tenormin	C		
		Restlessness		Lipitor	C		
				Coumadin	C	Glaxosmithkline	
				Norvasc	C		
				Diovan Hct	C		
				Tylenol	C	Glaxosmithkline	
				Citrucel	C	Glaxosmithkline	
				Dulcolax	C		

Date:05/27/05ISR Number: 4675992-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555394A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3	WK	Dizziness	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
UNKNOWN			Heart Rate Increased	Claritin D	C		
			Hot Flush				

Date:05/27/05ISR Number: 4675993-8Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549838A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3	DAY	Anxiety	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Insomnia	Lexapro	C		
			Panic Reaction	Trazodone	C		
			Psychomotor Hyperactivity	Alprazolam	C		
			Restlessness	Vitamins	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4675994-XReport Type:Periodic  
Age:61 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0555398A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK			Lisinopril	C		ORAL
40MG Per day				Atenolol	C		ORAL
				Lovastatin	C		ORAL
				Tetracycline	C		

YR

Date:05/27/05ISR Number: 4675995-1Report Type:Periodic  
Age:38 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0549844A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Heart Rate Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Prozac	C		
20MG Per day							

Date:05/27/05ISR Number: 4675996-3Report Type:Periodic  
Age:83 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0555561A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Drug Eruption		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	0 DAY						
Initial or Prolonged		Rash Generalised					
Other							

Date:05/27/05ISR Number: 4675997-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549850A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Emotional Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK						

Date:05/27/05ISR Number: 4675998-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0555565A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Menstruation Irregular		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4675999-9Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549856A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Valium	C		

Date:05/27/05ISR Number: 4676000-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555579A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Nasopharyngitis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676001-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549880A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperhidrosis		Wellbutrin Xl No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676002-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555587A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Deafness Unilateral		Wellbutrin Effexor Ritalin	PS C C	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676003-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549893A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 300MG Unknown		Agitation		Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676004-0Report Type:Periodic  
Age:55 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0555594A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 300MG Per day UNKNOWN		Amnesia Anger Confusional State Dizziness Hypersomnia Tremor		Wellbutrin Xl Cymbalta	PS SS	Glaxosmithkline	ORAL



Date:05/27/05ISR Number: 4676005-2Report Type:Periodic Company Report #0011380  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mouth Ulceration		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676006-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0555596A  
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Ambien	C		
				Depakote	C		

Date:05/27/05ISR Number: 4676007-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0549925A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Unknown	MON	Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676008-8Report Type:Periodic  
 Age:54 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555598A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Effexor	SS		
				Clarinet	C		
				Nasonex	C		
				Zantac	C	Glaxosmithkline	
				Vitamin B	C		
				Vitamin C	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676009-XReport Type:Periodic  
 Age:73 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550036A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Energy Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	18 MON			Darvocet	C		
		Mouth Ulceration		Pain Medication	C		
				Blood Pressure Medication	C		

Date:05/27/05ISR Number: 4676010-6Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555630A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Amnesia Memory Impairment		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676011-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550038A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		Mouth Ulceration		Wellbutrin	PS	Glaxosmithkline	

No Concurrent  
Medication C

Date:05/27/05ISR Number: 4676012-XReport Type:Periodic  
Age:32 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0555727A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Grand Mal Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 DAY						

Date:05/27/05ISR Number: 4676013-1Report Type:Periodic  
Age:26 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550052A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	27 DAY			No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676014-3Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555834A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG							
Variable dose							

Date:05/27/05ISR Number: 4676015-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550069A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl No Concurrent Medication	PS C	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676016-7Report Type:Periodic  
Age:13 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0555836A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tic		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 1	MON			Elavil	C	Glaxosmithkline	ORAL
10MG At night							

Date:05/27/05ISR Number: 4676017-9Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0550072A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Burning Sensation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 3	WK	Pruritus Urticaria					

Date:05/27/05ISR Number: 4676018-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0555840A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin	PS	Glaxosmithkline	
3	YR						

Date:05/27/05ISR Number: 4676019-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0550073A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Syncope		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other				Blood Pressure Medication	C		

Date:05/27/05ISR Number: 4676020-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0555844A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	6 WK						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676021-0Report Type:Periodic  
Age:25 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550081A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	7 DAY	Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Rash		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676022-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555848A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	2 WK	Pollakiuria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676023-4Report Type:Periodic  
Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550088A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	6 WK	Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dizziness		Oral Contraceptives	C		
		Nausea					
		Retching					
		Tinnitus					

Date:05/27/05ISR Number: 4676024-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555849A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	2 WK	Pollakiuria		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676025-8Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0550092A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Lethargy		Wellbutrin Sr	C	Glaxosmithkline	ORAL
150MG Twice							
per day				Celexa	C		
				Adderall	C		

Date:05/27/05ISR Number: 4676026-XReport Type:Periodic  
Age:62 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555864A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG							
Variable dose	8	MON		Atenolol	C		
				Levoxyl	C	Glaxosmithkline	
				Maxide	C		
				Arthrotec	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676027-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550148A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menorrhagia		Wellbutrin Xl Birth Control	PS C	Glaxosmithkline	ORAL
YR							

Date:05/27/05ISR Number: 4676028-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555897A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676029-5Report Type:Periodic  
 Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550246A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 DAY			Nasonex	C		
250MG Per day	10 YR			Imipramine	C		

Date:05/27/05ISR Number: 4676030-1Report Type:Periodic  
 Age:16 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555905A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Somnolence		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:05/27/05ISR Number: 4676031-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550253A



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Disorientation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY	Visual Disturbance		Lexapro	SS		

Date:05/27/05ISR Number: 4676032-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0556002A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676033-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0550256A  
 Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Flushing		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Pyrexia		Buspar	C		
		Tinnitus		Prozac	C		
300MG Per day	2 YR						
10 YR				Klonopin	C		
				Ventolin	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676034-9Report Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0556011A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Acne		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 YR			No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676035-0Report Type:Periodic  
Age:50 YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0550257A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:05/27/05ISR Number: 4676036-2Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0556022A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Dry Mouth		Zoloft	C		
		Hyperhidrosis		Crestor	C		
		Muscle Twitching		Nexium	C		
		Nausea		Actonel	C		
		Pollakiuria					
		Rash					
		Tremor					
		Weight Decreased					

Date:05/27/05ISR Number: 4676037-4Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550262A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day							

Ativan

C

Date:05/27/05ISR Number: 4676038-6Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0556058A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1	YR		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Depression					
		Dry Eye		Depakote	C		
		Somnolence		Premarin	C		
		Tinnitus		Calcium	C		
		Vision Blurred		Multivitamins	C		

Date:05/27/05ISR Number: 4676039-8Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0550277A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Muscle Spasms					
		Muscle Twitching		Ativan	C		
				Zoloft	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676040-4Report Type:Periodic  
Age:15 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0556105A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin Xl Adderall	PS C	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676041-6Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0550308A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Feeling Jittery		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676042-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0556107A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Arthralgia Insomnia Joint Stiffness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676043-XReport Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0550312A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
2 YR		Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676044-1Report Type:Periodic  
Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0556192A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	9 DAY	Flushing		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Flonase C Glaxosmithkline  
 Flax Seed C  
 Multivitamin C

ORAL

Date:05/27/05ISR Number: 4676045-3Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550433A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	5 YR	Renal Impairment		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Tremor		Espazine	C	Glaxosmithkline	
				Mysoline	C		
				Unknown Medication	C		
				Cogentin	C		
				Mevacor	C		
				Glucotrol	C		
				Zithromax	C		

Date:05/27/05ISR Number: 4676046-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0556203A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Unknown	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676047-7Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550437A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	4 DAY	Panic Attack		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Vision Blurred		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676048-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0556210A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	2 YR	Oedema Peripheral		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Rash		Cymbalta	C		
				Cymbalta	C		

Date:05/27/05ISR Number: 4676049-0Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550443A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 YR	Fatigue		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Prozac	C		
				Lasix	C	Glaxosmithkline	
				Tenormin	C		
				Diovan	C		

Date:05/27/05ISR Number: 4676050-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0556213A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Confusional State					
		Crying					

Hypoaesthesia

Date:05/27/05ISR Number: 4676051-9Report Type:Periodic  
 Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550465A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3	WK		Levothyroxine	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676052-0Report Type:Periodic  
 Age:25 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0556217A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1	YR		Oral Contraceptive	C		

Date:05/27/05ISR Number: 4676053-2Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0550475A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2	MON					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676054-4Report Type:Periodic  
Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0556218A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR			Estrogen	C		
				Diuretics	C		
				Allegra	C		

Date:05/27/05ISR Number: 4676055-6Report Type:Periodic  
Age:50 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0550488A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other		Hypomania					

Date:05/27/05ISR Number: 4676056-8Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0556310A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Apathy		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Crying		Prozac	C		
		Depression		Fioricet	C		
		Drug Ineffective		Klonopin	C		

Date:05/27/05ISR Number: 4676057-XReport Type:Periodic  
Age:24 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0550621A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amenorrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day							



Date:05/27/05ISR Number: 4676058-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0556320A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3 WK				Serzone	C		

Date:05/27/05ISR Number: 4676059-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0550627A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Spasms		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 2 WK				No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676060-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0556340A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Lamictal	C	Glaxosmithkline	
450MG Per day				Symbyax	C		
				Zyprexa	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676061-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550634A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK	Urticaria		Lescol	C		
				Imitrex	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676062-3Report Type:Periodic  
Age:53 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0556342A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 WK	Rash Maculo-Papular		Lipitor	C		

Date:05/27/05ISR Number: 4676063-5Report Type:Periodic  
Age:27 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550674A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Feeling Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK	Headache		Zonegran	C		
		Nausea		Lexapro	C		

Date:05/27/05ISR Number: 4676064-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0556346A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Vision Blurred		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150G Unknown				Effexor	C		

Date:05/27/05ISR Number: 4676065-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0550675A  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion					
300MG Per day		Dizziness Headache		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676066-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0556349A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
		Rash Generalised		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676067-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0550826A  
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	10 WK	Abnormal Dreams		Zoloft	C		
		Headache Nausea Tension					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676068-4Report Type:Periodic  
 Age:63 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0556441A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Feeling Jittery Insomnia Palpitations					

Date:05/27/05ISR Number: 4676069-6Report Type:Periodic  
 Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550855A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoaesthesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Single		Irritability					
dose	1 DAY			Cozaar	C		
				Hctz	C		
				Theophylline	C		
				Synthroid	C	Glaxosmithkline	
				Corgard	C		
				Combivent	C		
				Lopid	C		
				Flonase	C	Glaxosmithkline	
				Motrin	C	Glaxosmithkline	
				Methotrexate	C		
				Folic Acid	C		
				Glucophage	C		
				Pepcid	C		
				Tums	C	Glaxosmithkline	
				Soy Protein	C		
				Garlic Capsule	C		

Date:05/27/05ISR Number: 4676070-2Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0556673A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Unknown 10 DAY Headache  
Nausea

Wellbutrin Xl PS Glaxosmithkline ORAL

Date:05/27/05ISR Number: 4676071-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0550866A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Overdose		Subutex	C		

Date:05/27/05ISR Number: 4676072-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0556674A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Withdrawal Syndrome		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Sleep Disorder					
		Stomach Discomfort					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676073-8Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551011A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Ultram	C		

Date:05/27/05ISR Number: 4676074-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0556675A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	1 YR	Epistaxis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Fatigue					
		Insomnia					

Date:05/27/05ISR Number: 4676075-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551019A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK			Xanax	C		
Initial or Prolonged				Ativan	C		

Date:05/27/05ISR Number: 4676076-3Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0557127A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
6 MON		Lymphadenopathy		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676077-5Report Type:Periodic  
 Age:66 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551021A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2	MON		Lipitor	C		
				Blood Pressure Med	C		

Date:05/27/05ISR Number: 4676078-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0551026A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Acne		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day	13	MON		Strattera	C		
				Lexapro	C		
				Synthroid	C	Glaxosmithkline	
				Allergy Injections	C		
				Iron Supplement	C		
				Wellbutrin Sr	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676080-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0551043A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	
Other							
UNKNOWN	300MG Per day						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676082-9Report Type:Periodic  
Age:64 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551046A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Heart Rate Increased		Wellbutrin Xl Beta Blocker	PS C	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676084-2Report Type:Periodic  
Age:27 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0551047A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	3 WK	Chest Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Pruritus		Ortho Tri-Cyclen	C		
		Urticaria		Percocet	C		

Date:05/27/05ISR Number: 4676086-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551072A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Benign Prostatic Hyperplasia		Wellbutrin Xl Buspar	PS SS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676088-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551105A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 WK	Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Anxiety		Synthroid	C	Glaxosmithkline	
		Heart Rate Increased		Yasmin	C		
		Insomnia					
		Weight Decreased					



Date:05/27/05ISR Number: 4676090-8Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0551107A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676092-1Report Type:Periodic  
Age:58 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551116A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	2	MON	Abdominal Pain Upper	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Diarrhoea	Asacol	C	Glaxosmithkline	
			Dizziness	Lipitor	C		
			Dry Mouth	Toprol Xl	C		
			Fatigue	Valium	C		
			Hyperhidrosis	Codeine	C		
			Nausea	Tylenol	C	Glaxosmithkline	
			Tremor	Dicyclomine Hcl	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676093-3Report Type:Periodic  
Age:38 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551123A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion		Cymbalta	C		

12 DAY  
MON

Date:05/27/05ISR Number: 4676094-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551130A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion					

Date:05/27/05ISR Number: 4676095-7Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0551258A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Joint Swelling Paraesthesia					

Date:05/27/05ISR Number: 4676096-9Report Type:Periodic  
Age:42 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0551277A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Depression					

300MG Per day 17 DAY

Date:05/27/05ISR Number: 4676097-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551293A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

300MG Per day

Mouth Ulceration

Wellbutrin Xl PS Glaxosmithkline ORAL

Date:05/27/05ISR Number: 4676098-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0551301A  
 Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	4 DAY	Dizziness		Prenatal Vitamins	C		
		Dry Mouth					
		Myalgia					
		Nausea					
		Somnolence					

Date:05/27/05ISR Number: 4676099-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0551324A  
 Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Disability							
6 MON							
Other							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676100-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551328A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Spasms		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676101-XReport Type:Periodic  
Age:8 MON Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551340A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	
150MG Per day		Drug Exposure Via Breast Milk Irritability Somnolence		Prenatal Vitamins	C		

Date:05/27/05ISR Number: 4676102-1Report Type:Periodic  
Age:48 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0551347A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Twitching		Wellbutrin Xl Adderall	PS C	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676103-3Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551453A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - 150MG Per day 6 MON Initial or Prolonged				Lipitor Zyrtec	C C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676104-5Report Type:Periodic  
Age:36 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0551720A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Decreased Appetite Rash		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676105-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0551733A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Irritability		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676106-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0551734A  
 Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Headache		Aspirin	C	Glaxosmithkline	
		Heart Rate Increased		Premarin	C		
				Nexium	C		
				Plavix	C		
				Synthroid	C	Glaxosmithkline	
				Avalide	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Doxycycline C  
 Restoril C  
 Valium C  
 Miacalcin C  
 Vitamin C  
 Clonidine C

Date:05/27/05ISR Number: 4676107-0Report Type:Periodic  
 Age:44 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551735A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Per day Initial or Prolonged	34 WK	Suicidal Ideation		Wellbutrin	PS	Glaxosmithkline	ORAL
				Lexapro	C		
				Nexium	C		
				Lipitor	C		
				Lorazepam	C		
				Nasonex	C		
				Unknown Drug	C		

Date:05/27/05ISR Number: 4676108-2Report Type:Periodic  
 Age:48 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551907A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG In the morning	41 DAY	Dizziness Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Premarin	C		
				Actonel	C		

Date:05/27/05ISR Number: 4676109-4Report Type:Periodic  
 Age:22 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551935A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day Initial or Prolonged	2 MON	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676110-0Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551952A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	2	WK		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676111-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551954A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MGK Three							
times per day	7	DAY		No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676112-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551966A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
5 MON		Heart Rate Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
YR				Wellbutrin	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676113-6Report Type:Periodic  
 Age:25 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551971A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 MON		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676114-8Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551979A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Nausea					

Date:05/27/05ISR Number: 4676115-XReport Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0551983A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Unknown	3 DAY	Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676116-1Report Type:Periodic  
 Age:25 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551985A



Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Rash Macular		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3	DAY			Wellbutrin Sr	C	Glaxosmithkline	ORAL
12	DAY			Effexor	C		

Date:05/27/05ISR Number: 4676117-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0552009A  
 Age:41 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG In the morning	12 DAY	Pruritus					
200MG Twice per day				Bupropion Sr	SS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676118-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0552119A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Glucose Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 MON			No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676119-7Report Type:Periodic  
Age:70 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552126A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2	MON		Lexapro	C		

Date:05/27/05ISR Number: 4676120-3Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552132A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Weight Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1	YR		Prozac	C		
				Klonopin	C		

Date:05/27/05ISR Number: 4676121-5Report Type:Periodic  
Age:59 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552134A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Blood Sodium Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Three		Depression					
times per day	6	Gastrointestinal Disorder		Depakote	C		
	MON	Mental Disorder		Remeron	C		
		Thinking Abnormal		Gabitril	C		
				Ranitidine	C	Glaxosmithkline	
				Valtrex	C	Glaxosmithkline	
				Flomax	C		
				Celebrex	C		
				Aricept	C		
				Ditropan Xl	C		
				Propranolol	C		
				Restoril	C		
				Delatestryl	C		
				Prilosec	C	Glaxosmithkline	
				Benicar	C		
				Norvasc	C		

Date:05/27/05ISR Number: 4676122-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0552142A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin XL	PS	Glaxosmithkline	ORAL
300MG Per day		Libido Increased		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676123-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552159A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin XL	PS	Glaxosmithkline	ORAL
150MG Per day	4 WK	Paraesthesia		Klonopin	C		
		Tic					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676124-0Report Type:Periodic  
 Age:40 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552165A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Skin Lesion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK	Stevens-Johnson Syndrome					

Date:05/27/05ISR Number: 4676125-2Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552166A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Movement Disorder		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676126-4Report Type:Periodic  
 Age:40 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0552168A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK	Chills		Effexor Xr	C		
		Dyspnoea		Vasotec	C		
		Pyrexia		Hydrochlorothiazide	C		
		Swelling					
		Urticaria					
		White Blood Cell Count Increased					

Date:05/27/05ISR Number: 4676127-6Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0552294A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676128-8Report Type:Periodic  
Age:34 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552302A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Dizziness		Zoloft	C		
		Somnolence		Lopressor	C		
				Zanaflex	C		

Date:05/27/05ISR Number: 4676129-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552319A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged				Unknown	C		

Date:05/27/05ISR Number: 4676130-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552323A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676131-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552324A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3 DAY	Pharyngitis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Allopurinol	C	Glaxosmithkline	
				Cipro	C	Glaxosmithkline	
				Gemfibrozil	C		
				Lisinopril	C		
				Insulin	C		
				Glyburide	C		
				Metoprolol Xl	C		

Date:05/27/05ISR Number: 4676132-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552400A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
3 DAY		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676133-1Report Type:Periodic  
Age:39 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0552434A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG In the morning		Vision Blurred		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Ambien	C		
				Allegra	C		

Date:05/27/05ISR Number: 4676134-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552576A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Tinnitus  
 300MG Per day 2 YR  
 Wellbutrin Xl PS Glaxosmithkline ORAL

Date:05/27/05ISR Number: 4676135-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0552587A  
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR			Topomax	C		
				Evista	C		
				Lipitor	C		
				Nexium	C		
				Paxil	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676136-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0552589A  
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cognitive Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Disturbance In Attention Tension		Xanax	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676137-9Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552616A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
100MG Per day	6	WK		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
100MG Twice							
per day	1	MON		Synthroid	C	Glaxosmithkline	
				Prevacid	C		
				Demadex	C		
				Altace	C		
				Claritin	C		
				Glucophage	C		
				Lipitor	C		
				Risperdal	C		
				Singulair	C		
				Baby Aspirin	C	Glaxosmithkline	
				Colace	C		
				Xalatan	C		
				Miralax	C		
				Advair	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676138-0Report Type:Periodic  
Age:65 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552632A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Suicidal Ideation		Depakote	C		
150MG Per day	3	DAY					

Date:05/27/05ISR Number: 4676139-2Report Type:Periodic  
Age:28 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552786A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	20	DAY		Singulair	C		



Claritin C  
Effexor Xr C

Date:05/27/05ISR Number: 4676140-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552807A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Nausea		Oral Contraceptive	C		

Date:05/27/05ISR Number: 4676141-0Report Type:Periodic  
Age:15 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0552815A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Palpitations		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Amoxil	C	Glaxosmithkline	ORAL
500MG Twice							
per day	4	DAY		Advil	C	Glaxosmithkline	
200MG Twice							
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676142-2Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552818A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	2	MON		Lexapro	C		

Date:05/27/05ISR Number: 4676143-4Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552819A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	12	DAY		Claritin	C		

Date:05/27/05ISR Number: 4676144-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552837A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	1	MON		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676145-8Report Type:Periodic  
Age:33 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0552858A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Dyspnoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
8 DAY							
Other		Rash		Adderall Xr	C		ORAL
25MG Per day							
		Swelling					
		Urticaria					

Date:05/27/05ISR Number: 4676146-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552873A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
2 WK		Constipation					

Date:05/27/05ISR Number: 4676147-1Report Type:Periodic  
Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552875A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676148-3Report Type:Periodic  
Age:35 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0552985A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Unknown				Neurontin	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676149-5Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552993A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676150-1Report Type:Periodic  
Age:58 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0553000A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	MON	Sluggishness		Prenatal Vitamins	C		
				Potassium Supp	C		
				Nexium	C		
				Bextra	C		
				Nitroglycerin	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676151-3Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0553008A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other							

Date:05/27/05ISR Number: 4676152-5Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0553013A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	450MG Per day	Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676153-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0553038A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				Alcohol	SS		

Date:05/27/05ISR Number: 4676154-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0553053A  
Age:17 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG In the morning	13 DAY			Lexapro	C		
20MG per day							

Date:05/27/05ISR Number: 4676155-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0553164A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Discomfort		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
150MG Per day		Muscle Tightness		No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676156-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553165A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	3 WK		Wellbutrin	PS	Glaxosmithkline	ORAL
		Alopecia		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676157-4Report Type:Periodic  
Age: Gender:I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553167A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676158-6Report Type:Periodic  
Age: Gender:I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553180A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Affect Lability					

Date:05/27/05ISR Number: 4676159-8Report Type:Periodic  
Age:44 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553181A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	61 WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Gynaecomastia		Unknown Medication	C		
				Toprol Xl	C		
				Potassium	C		
				B12 Injection	C	Glaxosmithkline	
				Decadurabolin	C		
				Videx	C		
				Viramune	C		
				Paxil	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676160-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0553182A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Feeling Jittery					

Date:05/27/05ISR Number: 4676161-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0553183A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
Other		Convulsion					

Date:05/27/05ISR Number: 4676162-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0553194A  
 Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -		Convulsion					
300MG Per day	8 WK			Percodan	C		
Initial or Prolonged							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676163-XReport Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553197A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 MON	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Anorexia		Vitamins	C		
		Blepharospasm					
		Dry Skin					
		Dysgeusia					
		Nausea					
		Thinking Abnormal					

Date:05/27/05ISR Number: 4676164-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553198A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG In the morning		Abdominal Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Diarrhoea					

Date:05/27/05ISR Number: 4676165-3Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0553223A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Sensory Disturbance		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Sleep Disorder					

Date:05/27/05ISR Number: 4676166-5Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0553355A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1 WK	Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL



Date:05/27/05ISR Number: 4676167-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0553357A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypertension		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	0 DAY						

Date:05/27/05ISR Number: 4676168-9Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553358A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Anxiety		Benadryl	C	Glaxosmithkline	
		Dry Mouth					
		Insomnia					
		Micturition Frequency					
		Decreased					
		Orgasm Abnormal					
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676169-0Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0553411A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG In the morning		Insomnia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

No Concurrent Medication C

Date:05/27/05ISR Number: 4676170-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553418A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 YR UNKNOWN UNKNOWN		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Wellbutrin Sr	SS	Glaxosmithkline	
				Paxil	SS	Glaxosmithkline	

Date:05/27/05ISR Number: 4676171-9Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553425A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676172-0Report Type:Periodic  
 Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553565A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG In the morning		Diarrhoea Wheezing		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Synthroid	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676173-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553570A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	7 DAY	Arthralgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Synthroid	C	Glaxosmithkline	
				Prilosec	C	Glaxosmithkline	
				Tricor	C		
				Singulair	C		
				Celebrex	C		
				Nalfon	C		
				Glucophage	C		

Date:05/27/05ISR Number: 4676174-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553574A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
450MG In the		Faeces Discoloured		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
morning	6 WK	Nausea					
		Stomach Discomfort		Valium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676175-6Report Type:Periodic  
Age:53 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553581A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 YR			Geodon	C		

Date:05/27/05ISR Number: 4676176-8Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553607A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK			No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676177-XReport Type:Periodic  
Age:29 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553615A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Breast Cyst		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 YR	Fibroadenoma Of Breast		Zoloft	C		

Date:05/27/05ISR Number: 4676178-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553620A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3 DAY		Drug Ineffective		Lexapro	SS		
UNKNOWN	1 YR	Drug Interaction					
		Hyperhidrosis					
		Tremor					

Date:05/27/05ISR Number: 4676179-3Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0553621A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676180-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553622A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination, Auditory		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other							

Date:05/27/05ISR Number: 4676181-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553653A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
1	WK						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676182-3Report Type:Periodic  
Age:59 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553777A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	4	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Eye Pain					
		Headache		Benicar	C		
		Hyperhidrosis		Skelaxin	C		
		Hypertension		Protonix	C		
		Nausea		Potassium Supplement	C		
		Neuralgia		Celebrex	C		
		Photopsia		Vicodin	C		
		Red Blood Cell Count Decreased		Maxalt	C		
		Vomiting		Prednisolone Acetate	C		
		White Blood Cell Count Decreased		Betanol 0.3%	C		
				Alphagan	C		
				Ditropan	C		
				Benadryl	C	Glaxosmithkline	
				Zyrtec	C	Glaxosmithkline	
				Gerovit (Multivitamin)	C		
				Tylenol	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676183-5Report Type:Periodic  
Age:57 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553792A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1	DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Tinnitus					
				Vitamins	C		
				Blood Pressure Medication	C		

Date:05/27/05ISR Number: 4676184-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553799A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	2	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dizziness					

Date:05/27/05ISR Number: 4676185-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553805A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cough		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676186-0Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553812A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676187-2Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0553833A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676188-4Report Type:Periodic  
Age:41 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0553985A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Per day	Pollakiuria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Sleep Disorder		Prozac	C		
				Seroquel	C		
	UNKNOWN	25MG Per day					

Date:05/27/05ISR Number: 4676189-6Report Type:Periodic  
Age:73 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553989A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	159MG Per day	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	8 DAY	Hyperhidrosis		Cipro	C	Glaxosmithkline	
		Nervousness		Xanax	C		
				Vasotec	C		

Date:05/27/05ISR Number: 4676190-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554018A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Pruritus		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676191-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554044A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	2 MON	Oligomenorrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676192-6Report Type:Periodic  
Age:29 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0554057A



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 36 DAY			Grand Mal Convulsion	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Initial or Prolonged 25MG Per day				Paxil Cr	C	Glaxosmithkline	ORAL
Other 5MG Twice per day				Klonopin	C		
5MG At night				Abilify	C		

Date:05/27/05ISR Number: 4676193-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0554081A  
Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Unknown 1 YR			Convulsion	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676194-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0554084A  
Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1 YR			Convulsion	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676195-1Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0554203A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Parosmia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676196-3Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0554207A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Contusion		Wellbutrin Xl	PS	Glaxosmithkline	
UNKNOWN				Effexor Xr	C		

Date:05/27/05ISR Number: 4676197-5Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554210A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Discomfort		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day 2 DAY		Dry Mouth		Homeopathic Medicines	C		
		Hyperhidrosis					
		Lacrimation Increased					
		Nausea					
		Sinus Congestion					
		Throat Irritation					

Date:05/27/05ISR Number: 4676198-7Report Type:Periodic  
Age:17 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0554214A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus Generalised		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 3 WK		Rash Generalised		Zoloft	C		

Urticaria

Date:05/27/05ISR Number: 4676199-9Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0554223A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676200-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554224A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Coeliac Disease		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Drug Ineffective					

Date:05/27/05ISR Number: 4676201-4Report Type:Periodic  
Age:30 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0554227A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Galactorrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	7 MON						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676202-6Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554232A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day				Advair	C	Glaxosmithkline	
				Albuterol	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676203-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0554238A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Migraine		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	6 WK			No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676204-XReport Type:Periodic  
Age:80 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0515873A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Skin		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	4 DAY	Erythema		Effexor	C		
		Oedema		Aricept	C		
		Pain		Metoprolol	C		
		Pruritus		Neurontin	C		
		Rash		Colchicine	C		
		Skin Exfoliation		Lipitor	C		
				Terazosin	C		
				Proscar	C		
				Indomethacin	C		

Date:05/27/05ISR Number: 4676205-1Report Type:Periodic  
Age:44 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0525050A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day	9	MON	Dysgeusia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Pharmaceutical Product Complaint	Paroxetine Synthroid	C C	Glaxosmithkline Glaxosmithkline	

Date:05/27/05ISR Number: 4676206-3Report Type:Periodic  
Age:50 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0531285A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300MG Per day	5	DAY	Dizziness	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	300MG Per day	5	MON	Palpitations	Lamictal	SS	Glaxosmithkline	ORAL
				Skin Hyperpigmentation	Synthroid	C	Glaxosmithkline	
	TOPICAL	45Z Unknown		Swollen Tongue	Sunscreen	C		
					Levothyroxine	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676207-5Report Type:Periodic  
Age:22 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0532862A

Outcome	PT
	Agitation
	Chills
	Diarrhoea
	Dissociation

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dyskinesia Erectile Dysfunction Fear					
150MG Per day	6 MON	Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
12.5MG Per day	4 DAY	Insomnia		Paxil Cr	SS	Glaxosmithkline	ORAL
		Muscle Tightness					
		Muscle Twitching		Aciphex	C		
		Nausea		Lopressor	C		
		Sleep Disorder		Antibiotic	C		
UNKNOWN				Nexium	C		
UNKNOWN				Neurontin	C		

Date:05/27/05ISR Number: 4676208-7Report Type:Periodic Company Report #US-GLAXOSMITHKLIN-A0535736A  
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	3 WK	Bowel Movement		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Irregularity		No Concurrent Medication	C		
		Drug Ineffective					
		Fatigue					
		Hunger					
		Loss Of Libido					
		Weight Increased					

Date:05/27/05ISR Number: 4676209-9Report Type:Periodic Company Report #US-GLAXOSMITHKLIN-A0538242A  
 Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	150MG Per day	3 DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Blood Glucose Decreased		Nexium	C		
		Chest Pain		Estrace	C		
		Dry Mouth					
		Dyspnoea					

Heart Rate Increased  
Muscular Weakness  
Nausea  
Nervousness  
Rash  
Thinking Abnormal  
Tinnitus  
Weight Decreased

Date:05/27/05ISR Number: 4676210-5Report Type:Periodic  
Age:32 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0539064A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	10	MON	Affect Lability	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Drug Ineffective	Glucophage	C		
			Sleep Disorder	Oral Contraceptives	C		
150MG As				Diflucan	C		ORAL
required							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676211-7Report Type:Periodic  
Age:62 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0540784A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2	MON		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Agitation					
		Anxiety		Prozac	C		
		Dry Mouth		Librium	C		
		Feeling Jittery		Ambien	C		
		Insomnia		Lopressor	C		
		Tremor					

Date:05/27/05ISR Number: 4676212-9Report Type:Periodic  
Age:57 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0541953A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Tinnitus					
				Hormone Replacement	C		

Date:05/27/05ISR Number: 4676213-0Report Type:Periodic  
Age:61 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0541963A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Anaphylactic Reaction		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	16	DAY					
Hospitalization -		Dyspnoea		Depakote	C		
UNKNOWN	500MG	Twice					
Initial or Prolonged		Pruritus Generalised					
per day							
Other		Swelling Face		Percocet	C		
UNKNOWN	325MG	As					
required		Swollen Tongue					
				Lorazepam	C		ORAL
2MG Four							
times per day							
				Klonopin	C		
UNKNOWN	1MG	At night					



100MG Twice		Zoloft	C		ORAL
per day					
UNKNOWN	40MG Twice	Furosemide	C	Glaxosmithkline	
per day					
UNKNOWN		Lotrel	C		
UNKNOWN		Klor-Con	C	Glaxosmithkline	
UNKNOWN	40MG Three	Oxycontin	C		
times per day					
20MG Twice		Isosorbide	C		
per day					

Date:05/27/05ISR Number: 4676214-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0542478A  
 Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hallucination, Auditory		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Unknown	20 DAY						

Date:05/27/05ISR Number: 4676215-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0542853A  
 Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	14 MON	Pharmaceutical Product Complaint Weight Decreased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676216-6Report Type:Periodic  
Age:50 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0542863A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 YR	Grand Mal Convulsion Tongue Biting		Anti-Viral Medication	C		
UNKNOWN				Toprol	C		
UNKNOWN							

Date:05/27/05ISR Number: 4676217-8Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543284A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Arrhythmia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676218-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543287A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	2 WK	Feeling Jittery		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Insomnia		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676219-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543288A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	6 MON	Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676220-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0543319A  
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - WK		Hypersensitivity		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Initial or Prolonged Other		Swelling Face Urticaria					

Date:05/27/05ISR Number: 4676221-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0543322A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Headache Nausea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676222-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0543326A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				Lexapro	C		
				Risperdal	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676223-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543332A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tachycardia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 DAY						

Date:05/27/05ISR Number: 4676224-5Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543333A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK	Gingival Pruritus Local Swelling Mood Altered Mouth Ulceration Sensitivity Of Teeth		Verapamil	C		

Date:05/27/05ISR Number: 4676225-7Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543336A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Paraesthesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	7 DAY			Xanax	C		
				Prilosec	C	Glaxosmithkline	
				Cymbalta	C		

Date:05/27/05ISR Number: 4676226-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543338A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676227-0Report Type:Periodic  
Age:45 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543485A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Feeling Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice		Memory Impairment					
per day		Panic Reaction		Synthroid	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676228-2Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543489A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Aggression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	6 WK	Agitation		Zoloft	C		
		Depression		Xanax	C		
		Heart Rate Increased		Aciphex	C		
		Irritability					
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676229-4Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543500A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:05/27/05ISR Number: 4676230-0Report Type:Periodic  
Age:65 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0543502A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Deafness Bilateral		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Tinnitus		Advair	C	Glaxosmithkline	
300MG Per day	5 MON			Albuterol	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676231-2Report Type:Periodic  
Age:20 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543506A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK			Birth Control Pills	C		

Date:05/27/05ISR Number: 4676232-4Report Type:Periodic  
Age:43 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543513A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6 MON	Insomnia		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
				Ambien	C		
				Naprosyn	C		

Date:05/27/05ISR Number: 4676233-6Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543516A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Pharyngolaryngeal Pain	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2	YR	Tongue Discolouration	Actonel	C		
				Prempro	C		
				Calcium	C		
				Vitamin D	C		
				Vitamin C	C	Glaxosmithkline	
				Colestid	C		
				Nexium	C		

Date:05/27/05ISR Number: 4676234-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543528A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Joint Swelling	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2	WK	Throat Tightness	Zyrtec	C	Glaxosmithkline	
			Urticaria				

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Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676235-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543699A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	Confusional State		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Food Allergy		Tylenol	C	Glaxosmithkline	
		Headache		Benadryl	C	Glaxosmithkline	
		Nausea					
		Nicotine Dependence					
		Panic Attack					

Date:05/27/05ISR Number: 4676236-1Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0543701A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysmenorrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676237-3Report Type:Periodic  
 Age:23 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0543716A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day 1 WK	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
	Initial or Prolonged	Incontinence					
		Weight Decreased					

Date:05/27/05ISR Number: 4676238-5Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543721A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day 3 WK	Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Anxiety					
		Hypoacusis					
		Tinnitus					



Date:05/27/05ISR Number: 4676239-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0543729A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:05/27/05ISR Number: 4676240-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543730A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Euphoric Mood		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Concerta	C		

Date:05/27/05ISR Number: 4676241-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543744A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other				No Concurrent Medication	C		
300MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676242-7Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543953A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	5 DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Anxiety					
		Irritability		Levoxyl	C	Glaxosmithkline	
				Lasix	C	Glaxosmithkline	
				Trazodone	C		
				Morphine	C		
				Klonopin	C		

Date:05/27/05ISR Number: 4676243-9Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543962A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Heart Rate Increased					
		Ventricular Extrasystoles					

Date:05/27/05ISR Number: 4676244-0Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543976A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	
Other		Convulsion					

Date:05/27/05ISR Number: 4676245-2Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0543978A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG See			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Agitation					
		Anger					
dosage text	8 WK	Dry Mouth					

Date:05/27/05ISR Number: 4676246-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0543986A  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	2 WK						

Date:05/27/05ISR Number: 4676247-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0543990A  
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sensation Of Heaviness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Haldol	C		

Date:05/27/05ISR Number: 4676248-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0544008A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Unknown							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676249-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544027A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams Nightmare		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676250-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544040A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676251-8Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544221A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 MON	Therapeutic Response		Ditropan Xl	C		
5MG per day		Unexpected		Azathioprine	C	Glaxosmithkline	
75MG per day							

Date:05/27/05ISR Number: 4676252-XReport Type:Periodic  
Age:18 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544247A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		White Blood Cell Count Decreased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 YR			No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676253-1Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544251A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Blood Pressure Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Feeling Abnormal					
		Tension Headache					
		Therapeutic Response					
		Unexpected					

Date:05/27/05ISR Number: 4676254-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0544258A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	24 WK						

Date:05/27/05ISR Number: 4676255-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0544272A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Wellbutrin Sr	SS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676256-7Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544308A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Purpura		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK						

Date:05/27/05ISR Number: 4676257-9Report Type:Periodic  
 Age:33 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0544498A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dyspnoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day	5 MON						
		Nausea		Nuvaring	C		

Date:05/27/05ISR Number: 4676258-0Report Type:Periodic  
 Age:31 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544524A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	4 DAY						
UNKNOWN	40MG Per day	Disturbance In Attention		Prozac	SS		
		Insomnia					

Date:05/27/05ISR Number: 4676259-2Report Type:Periodic  
 Age:77 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544530A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dry Eye		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	5 DAY						
		Dry Mouth		Namenda	C		

Date:05/27/05ISR Number: 4676260-9Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544544A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676261-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0544661A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
6 WK		Anger		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Crying Depression Drug Ineffective		Celexa	C		

Date:05/27/05ISR Number: 4676262-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0544758A  
 Age:41 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG Per day		Hallucination, Auditory		Wellbutrin	PS	Glaxosmithkline	ORAL
				Anti-Psychotic Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676263-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544760A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
13 DAY		Hyperchlorhydria Hypersensitivity Oedema Peripheral Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676264-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544783A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Dysphagia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Klonopin	C		
				Neurontin	C		

Date:05/27/05ISR Number: 4676265-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544787A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Twice per day	10 DAY	Overdose Tremor		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Paxil Cr	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676266-XReport Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0544936A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day		Blood Pressure Diastolic Increased Blood Pressure Systolic		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		



Increased

Date:05/27/05ISR Number: 4676267-1Report Type:Periodic  
Age:21 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544940A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	6	MON		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676268-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544953A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Nausea Palpitations Paraesthesia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676269-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545083A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676270-1Report Type:Periodic  
Age:20 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545149A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 YR	Feeling Drunk		Birth Control	C		

Date:05/27/05ISR Number: 4676271-3Report Type:Periodic  
Age:68 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545167A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Deafness		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Hypertension		Lithium	C	Glaxosmithkline	
150MG Per day		Tinnitus		Levothyroxine	C	Glaxosmithkline	
				Benicar	C		

Date:05/27/05ISR Number: 4676272-5Report Type:Periodic  
Age:49 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0545168A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day		Anxiety		Xanax	C		ORAL
.5U Twice per		Burning Sensation					
day		Euphoric Mood		Trazodone	C		ORAL
50MG As							

required

Feeling Hot

Nervousness

Therapeutic Response

Unexpected

Date:05/27/05ISR Number: 4676273-7Report Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0545178A

Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	5 DAY			Aciphex	C		

Date:05/27/05ISR Number: 4676274-9Report Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0545180A

Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice		Somnolence		Xanax	C		
per day	5 WK			Sinemet	C		
				Vitamins	C		
				Metoprolol	C		
				Resperidol	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676275-0Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0545183A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	3 DAY		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Convulsion		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676276-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545329A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Memory Impairment					

Date:05/27/05ISR Number: 4676277-4Report Type:Periodic  
Age:83 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545448A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	1 WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dry Mouth		Lipram	C		
		Dyspepsia		Zocor	C		
				Surfak	C		
				Bextra	C		
				Norvasc	C		
				Protonix	C		
				Doxazosin	C		
				Antacid	C		

Date:05/27/05ISR Number: 4676278-6Report Type:Periodic  
Age:77 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545458A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	6 DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Anorexia		Lipitor	C		
		Depression					

Insomnia

Norvasc	C	
Vitamins	C	
Gabapentin	C	
Tylenol	C	Glaxosmithkline

Date:05/27/05ISR Number: 4676279-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545478A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK	Swelling Urticaria					

Date:05/27/05ISR Number: 4676280-4Report Type:Periodic  
 Age:52 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545480A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	5 WK	Anxiety Tachycardia		Plavix Lipitor Diovan	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676281-6Report Type:Periodic  
Age:46 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545487A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
450MG In the		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
morning	2 WK	Dry Mouth					
		Insomnia					
		Tremor					
		Weight Decreased					

Date:05/27/05ISR Number: 4676282-8Report Type:Periodic  
Age:53 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0545543A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination, Auditory		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day							

Date:05/27/05ISR Number: 4676283-XReport Type:Periodic  
Age:41 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0545678A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Muscle Rigidity		Toprol Xl	C		
		Suffocation Feeling		Clonazepam	C		
				Ortho Tri Cyclen	C		

Date:05/27/05ISR Number: 4676284-1Report Type:Periodic  
Age:61 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545686A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	25 DAY	Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Swelling Face		Alprazolam	C		
				Lorazepam	C		

Date:05/27/05ISR Number: 4676285-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545693A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	6	DAY	Headache	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Insomnia	No Concurrent			
			Rash Pruritic	Medication	C		
			Urticaria				

Date:05/27/05ISR Number: 4676286-5Report Type:Periodic  
Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545701A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	1	YR	Agitation	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Pruritus	Bupropion	SS	Glaxosmithkline	ORAL
75MG Per day	1	WK	Sleep Disorder	Lexapro	C		
			Vision Blurred	Toprol Xl	C		
				Lipitor	C		
				Allegra	C		
				Atacand	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676287-7Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545705A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3 WK		Rash Generalised		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676288-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545711A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Irritability		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676289-0Report Type:Periodic  
Age:39 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0545719A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2TAB Three times per day 9 DAY		Feeling Jittery		Wellbutrin Xl Ultram	PS C	Glaxosmithkline	ORAL ORAL

Date:05/27/05ISR Number: 4676290-7Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0545736A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 MON 150MG Unknown MON		Drug Ineffective		Wellbutrin Xl Wellbutrin Sr	PS SS	Glaxosmithkline	ORAL ORAL

Date:05/27/05ISR Number: 4676291-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0545746A



Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Diastolic		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Increased					

Date:05/27/05ISR Number: 4676292-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0545766A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other		Agitation					

Date:05/27/05ISR Number: 4676293-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0545769A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Unknown	6 WK	Drug Ineffective					

Date:05/27/05ISR Number: 4676294-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0545803A  
 Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Crying		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	16 DAY	Insomnia		Nicotine Patch	C	Glaxosmithkline	
UNKNOWN		Tinnitus					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676295-6Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0545805A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG In the morning		Anorexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
12.5MG At night		Anxiety		Paxil Cr	SS	Glaxosmithkline	ORAL
UNKNOWN	.5MG At night	Dizziness		Ativan	SS		
		Nausea					
		Tremor					

Date:05/27/05ISR Number: 4676296-8Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545917A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	1 WK	Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Eating Disorder		No Concurrent Medication	C		
		Tinnitus					

Date:05/27/05ISR Number: 4676297-XReport Type:Periodic  
 Age:38 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545932A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG		Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Alternate days	8 WK			Synthroid	C	Glaxosmithkline	
				Xanax	C		

Date:05/27/05ISR Number: 4676298-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0545935A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Heart Rate Increased		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676299-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0545946A  
Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 DAY			Neurontin	C		
300MG At							
night							
10MG At night				Nortriptyline	C		

Date:05/27/05ISR Number: 4676300-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0545954A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Orgasmic Sensation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	8 MON	Decreased Vulvovaginal Dryness		Birth Control Pills	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676301-9Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0546033A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin	PS	Glaxosmithkline	

Date:05/27/05ISR Number: 4676302-0Report Type:Periodic  
Age:38 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0546035A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination		Wellbutrin	PS	Glaxosmithkline	ORAL
Disability							
150MG Per day	5 DAY						
Other		Hyperhidrosis		Effexor Xr	SS		

Date:05/27/05ISR Number: 4676303-2Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546044A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Wellbutrin	PS	Glaxosmithkline	
Other							
300MG Per day							

Date:05/27/05ISR Number: 4676304-4Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0546046A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day							

Date:05/27/05ISR Number: 4676305-6Report Type:Periodic  
Age:61 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0546049A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	10 MON						

Energy Increased  
Insomnia  
Thyroid Function Test  
Abnormal

Xanax C  
Ambien C

Date:05/27/05ISR Number: 4676306-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0546050A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	10 MON			Glucophage	C		
				Atenolol	C		

Date:05/27/05ISR Number: 4676307-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0546051A  
Age:22 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Pain Upper		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 DAY			Adderall	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676308-1Report Type:Periodic  
Age:52 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546074A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Per day	4 WK	Burning Sensation	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Lopressor	C		
				Lipitor	C		
				Xanax	C		

Date:05/27/05ISR Number: 4676309-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546076A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Cough	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676310-XReport Type:Periodic  
Age:35 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546081A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	150MG Per day	9 DAY	Eye Pain	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Headache	Amoxicillin	C	Glaxosmithkline	
			Malaise				

Date:05/27/05ISR Number: 4676311-1Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0546291A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
	300MG Unknown		Rash	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676312-3Report Type:Periodic  
Age:56 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546318A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	300MG Per day 8	MON	Condition Aggravated	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged			Pollakiuria	Ativan	C		
			Weight Decreased	Geodon	C		

Date:05/27/05ISR Number: 4676313-5Report Type:Periodic  
Age:52 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546323A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day 6	MON	Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Crying		Levoxyl	C	Glaxosmithkline	
		Irritability		Ambien	C		
				Topamax	C		
				Tricor	C		
				Ibuprofen	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676314-7Report Type:Periodic  
Age:46 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546325A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day 3	DAY	Pruritus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Rash		Synthroid	C	Glaxosmithkline	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676315-9Report Type:Periodic  
Age:49 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546332A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				No Concurrent Medication	C		
				Meridia	C		

Date:05/27/05ISR Number: 4676316-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546340A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 1 YR		Drug Ineffective		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
		Feeling Jittery					

Date:05/27/05ISR Number: 4676317-2Report Type:Periodic  
Age:39 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0546347A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 79 DAY				Effexor	SS		ORAL
75MG Unknown				Zonegran	C		
UNKNOWN				Cozar	C		
				Lipitor	C		
				Glucotrol	C		
				Cialis	C		

Date:05/27/05ISR Number: 4676318-4Report Type:Periodic  
Age:47 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0546505A



Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Pharmaceutical Product		Wellbutrin	PS	Glaxosmithkline	ORAL
	150MG	Per day	Complaint		Prozac	C		
			Poor Quality Drug		Lotensin	C	Glaxosmithkline	
			Administered		Seroquel	C		
					Clonazepam	C		
					Celebrex	C		

Date:05/27/05ISR Number: 4676319-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0546521A  
Age:61 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Hypotrichosis		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	300MG	Per day						

Date:05/27/05ISR Number: 4676320-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0546721A  
Age:55 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Fatigue		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	300MG	Per day 4 WK	Influenza Like Illness		Synthroid	C	Glaxosmithkline	
			Insomnia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676321-4Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546722A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day		Urticaria					

Date:05/27/05ISR Number: 4676322-6Report Type:Periodic  
 Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546729A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chills		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK	Dizziness Headache Hyperhidrosis Vomiting		Antibiotic	C		

Date:05/27/05ISR Number: 4676323-8Report Type:Periodic  
 Age:13 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546734A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	10 MON	Rheumatoid Factor Positive		Neurontin Topamax	C C		

Date:05/27/05ISR Number: 4676324-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546761A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	1 WK						

Date:05/27/05ISR Number: 4676325-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0546768A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation Feeling Abnormal Feeling Jittery		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676326-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0546812A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
3	WK						

Date:05/27/05ISR Number: 4676327-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0546926A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	30 MON			No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676328-7Report Type:Periodic  
Age:21 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546931A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Therapeutic Response Unexpected		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676329-9Report Type:Periodic  
Age:44 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546932A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Fatigue		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Paraesthesia		Klonopin	C		
				Glucovance	C		
				Vasotec	C		
				Prometrium	C		

Date:05/27/05ISR Number: 4676331-7Report Type:Periodic  
Age:37 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0546939A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Urticaria		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
per day	1 YR			Omeprazole	SS	Glaxosmithkline	ORAL
20MG Per day				Prilosec	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676333-0Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546942A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Stevens-Johnson Syndrome Wellbutrin PS Glaxosmithkline  
UNKNOWN  
Initial or Prolonged

Date:05/27/05ISR Number: 4676334-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0540242A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	
UNKNOWN			Nuclear Magnetic Resonance Imaging Abnormal				

Date:05/27/05ISR Number: 4676335-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0546943A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin	PS	Glaxosmithkline	ORAL
Other			Hepatic Enzyme Increased				
300MG Per day	12 WK			No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676336-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0541134A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG In the morning			Dystonia				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Topomax SS  
 Geodon C  
 Prozac C

Date:05/27/05ISR Number: 4676337-8Report Type:Periodic  
 Age:55 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546944A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	1 WK			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dyspepsia					
		Insomnia		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676338-XReport Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0542666A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	

Date:05/27/05ISR Number: 4676339-1Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546948A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
450MG Per day		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676340-8Report Type:Periodic  
 Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0543964A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day							

Date:05/27/05ISR Number: 4676341-XReport Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546964A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3 WK	Constipation					
		Drug Ineffective					
		Insomnia					
		Paraesthesia					

Date:05/27/05ISR Number: 4676342-1Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0550316A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Pharmaceutical Product					
		Complaint					
		Stool Analysis Abnormal					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676343-3Report Type:Periodic  
Age:23 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546968A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	4 DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Somnolence		Abilify	C		

Date:05/27/05ISR Number: 4676345-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546971A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	2 DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Dizziness		No Concurrent Medication	C		
		Stomach Discomfort					

Date:05/27/05ISR Number: 4676347-0Report Type:Periodic  
Age:46 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0546999A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	300MG Per day	78 DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	.5MG As			Ativan	C		
	required						

Date:05/27/05ISR Number: 4676348-2Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0547034A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Hypersensitivity					



Date:05/27/05ISR Number: 4676349-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0547135A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK	Chest Pain		Xanax	C		
		Tinnitus		Alprazolam	C		

Date:05/27/05ISR Number: 4676350-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0547149A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK			Blood Pressure Medication	C		

Date:05/27/05ISR Number: 4676351-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0547157A  
Age:22 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day				Marijuana	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676352-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547174A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
18 MON		Gingival Disorder Tooth Disorder Tooth Infection		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676353-6Report Type:Periodic  
Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547184A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Per day	1 WK	Anorexia Anxiety Dry Mouth Dyspnoea Flashback Insomnia Nausea Rash Tremor		Wellbutrin Xl Voltaren	PS C	Glaxosmithkline Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676354-8Report Type:Periodic  
Age:45 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0547185A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
450MG Unknown	WK	Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676355-XReport Type:Periodic  
Age:60 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0547186A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
450MG Unknown	WK	Tinnitus		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676356-1Report Type:Periodic  
Age:65 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0547354A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	2 WK	Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Nausea		Coumadin	C	Glaxosmithkline	
				Ultram	C		

Date:05/27/05ISR Number: 4676357-3Report Type:Periodic  
Age:55 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547361A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day	Erectile Dysfunction		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	2 YR			Allegra	C		

Date:05/27/05ISR Number: 4676358-5Report Type:Periodic  
Age:59 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547371A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	150MG Per day	Affect Lability		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	7 WK	Drug Ineffective		Lipitor	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676359-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547372A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	3	WK		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Clarinet	C		

Date:05/27/05ISR Number: 4676360-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547375A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	21	DAY		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Nexium	C		

Date:05/27/05ISR Number: 4676361-5Report Type:Periodic  
Age:80 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0547376A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -				Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	1	MON					
Initial or Prolonged							

Date:05/27/05ISR Number: 4676362-7Report Type:Periodic  
Age:17 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0547378A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -				Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day							
Initial or Prolonged				Lexapro	C		

Date:05/27/05ISR Number: 4676363-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547381A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Cholecystectomy		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 YR	Drug Ineffective		Alprazolam Quitex	C C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676364-0Report Type:Periodic  
Age:37 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547387A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Galactorrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 MON						

Date:05/27/05ISR Number: 4676365-2Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0547397A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Amnesia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Medication Error		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
		Pharmaceutical Product		No Concurrent			
		Complaint		Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676366-4Report Type:Periodic  
Age:42 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547555A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day	4 DAY	Depression		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Sleep Disorder					

Date:05/27/05ISR Number: 4676367-6Report Type:Periodic  
Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547564A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	74 WK	Headache		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Multivitamins	C		

Date:05/27/05ISR Number: 4676368-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547566A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day		Adverse Event		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	10 MON	Bruxism		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
		Feeling Hot		Dexedrine	SS	Glaxosmithkline	
		Insomnia		Paxil	C	Glaxosmithkline	
		Rash		Progesterone Cream	C		
				Tenormin	C		
				Klonopin	C		
				Fioricet	C		
				Vicodin	C		
				Acyclovir	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676369-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547570A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

150MG Per day	6	MON	Abdominal Discomfort	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Cymbalta	C		
				Protonix	C		

Date:05/27/05ISR Number: 4676370-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0547580A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Euphoric Mood		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	7	DAY	Sleep Disorder	Nexium	C		
				Tricor	C		

Date:05/27/05ISR Number: 4676371-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0547582A  
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day		Drug Ineffective		Nexium	C		
		Irritability		Allegra	C		
		Libido Decreased		Multivitamin	C		
		Pruritus					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676372-XReport Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547592A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Discomfort		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676373-1Report Type:Periodic  
Age:44 YR Gender:Female I/FU:F

Company Report #05-01-0123

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hospitalization - 300MG Per day		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Initial or Prolonged 400MG Per day 5 YR		Clozapine	SS		ORAL

Date:05/27/05ISR Number: 4676374-3Report Type:Periodic  
Age:37 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0547637A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Galactorrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		300MG Per day 2 MON					

Date:05/27/05ISR Number: 4676375-5Report Type:Periodic  
Age: Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0547759A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Wellbutrin	PS	Glaxosmithkline	ORAL
		UNKNOWN		Wellbutrin	SS	Glaxosmithkline	

Date:05/27/05ISR Number: 4676376-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547767A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							



Other Convulsion Wellbutrin PS Glaxosmithkline ORAL  
450MG Per day  
60MG per day Prozac C

Date:05/27/05ISR Number: 4676377-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0547772A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ejaculation Failure Libido Increased		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676378-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0547782A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Stool Analysis Abnormal		Wellbutrin Xl Prednisone	PS C	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676379-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0547790A  
Age: Gender:Female I/FU:I

Outcome PT  
Drug Ineffective  
Dry Mouth  
Palpitations  
Tinnitus

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vertigo

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Per day			Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Paxil	SS	Glaxosmithkline	ORAL
			Synthroid	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676380-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0547804A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day		Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Drug Withdrawal Syndrome					
		Hunger					
		Nicotine Dependence					
		Weight Increased					

Date:05/27/05ISR Number: 4676381-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0547972A  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Muscle Twitching		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676382-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0547979A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day Initial or Prolonged		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
				Prozac	C		

Date:05/27/05ISR Number: 4676383-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547984A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	2 DAY	Chest Discomfort	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Dizziness	Xanax	C		
			Dyspnoea	Geodon	C		
			Heart Rate Increased				
			Paraesthesia				
			Throat Tightness				
			Urine Abnormality				

Date:05/27/05ISR Number: 4676384-6Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0548002A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Convulsion	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676385-8Report Type:Periodic  
Age:36 YR Gender: I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0548150A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300MG Per day Initial or Prolonged Other	Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676386-XReport Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548173A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration 150MG Per day	Paranoia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	Rash Generalised Thinking Abnormal Urticaria		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676387-1Report Type:Periodic  
Age:53 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548180A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration 300MG Per day	Back Pain		Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676388-3Report Type:Periodic  
Age:20 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548184A

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration 300MG Per day	Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	Heart Rate Increased		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676389-5Report Type:Periodic  
Age:57 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548187A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	52 WK	Flushing Swelling Face		Synthroid	C	Glaxosmithkline	

Date:05/27/05ISR Number: 4676390-1Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548192A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	10 DAY	Insomnia Mouth Ulceration Rash Tinnitus Urticaria		No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676391-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548197A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Unknown	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676392-5Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548207A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	7 MON	Constipation	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Drug Administration Error	Clonidine	C		
				Xanax	C		

Date:05/27/05ISR Number: 4676393-7Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548230A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day	2 DAY	Musculoskeletal Chest	Wellbutrin	PS	Glaxosmithkline	
			Initial or Prolonged	Paxil	C	Glaxosmithkline	
				Topomax	C		

Date:05/27/05ISR Number: 4676394-9Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548231A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Hospitalization -	4 WK	Dizziness	Wellbutrin	PS	Glaxosmithkline	ORAL
			Initial or Prolonged	Alcohol	C		

Date:05/27/05ISR Number: 4676395-0Report Type:Periodic  
Age:27 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0548259A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Life-Threatening	450MG Per day	Convulsion	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	Other	1MG Twice per		Ativan	SS		ORAL

day

7.5MG Per day				Remeron	SS		ORAL
8MG As				Zofran	SS	Glaxosmithkline	ORAL
required							
150MG At				Seroquel	SS		ORAL
night							
UNKNOWN	100MG Per day			Prozac	SS		

Date:05/27/05ISR Number: 4676396-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0548361A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Feeling Abnormal		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Per day	6 WK	Lethargy		Valium	C		

Date:05/27/05ISR Number: 4676397-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0548374A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Constipation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
450MG Per day	4 WK	Flatulence Tremor		Paxil	C	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676398-6Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548380A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscular Weakness		Wellbutrin Xl Unknown	PS C	Glaxosmithkline	ORAL

Date:05/27/05ISR Number: 4676399-8Report Type:Periodic  
Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548381A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pyrexia		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 YR			Ortho Novum 777	C		

Date:05/27/05ISR Number: 4676400-1Report Type:Periodic  
Age:11 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0548387A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Grand Mal Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Hospitalization - 300MG Per day	2 WK			Benadryl	C	Glaxosmithkline	
Initial or Prolonged Other							

Date:05/27/05ISR Number: 4676401-3Report Type:Periodic  
Age:36 YR Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548388A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
150MG Twice per day	8 DAY	Nausea		Minocycline Lipitor	C C		



Date:05/27/05ISR Number: 4676402-5Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548407A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl Effexor	PS SS	Glaxosmithkline	ORAL
		Blood Pressure Increased					

Date:05/27/05ISR Number: 4676403-7Report Type:Periodic  
Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0548443A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG In the morning		Irritable Bowel Syndrome Stool Analysis Abnormal					

Date:05/27/05ISR Number: 4676404-9Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548598A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Wellbutrin Xl No Concurrent Medication	PS C	Glaxosmithkline	ORAL
		Alopecia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676405-0Report Type:Periodic  
Age:31 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548602A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -	6 MON		Grand Mal Convulsion	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged				Valium	C		

Date:05/27/05ISR Number: 4676406-2Report Type:Periodic  
Age:32 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548617A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Per day	3 WK		Eye Pain	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
			Muscle Twitching	No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676407-4Report Type:Periodic  
Age:56 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548618A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Paraesthesia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				Sinemet	C		
				Comtan	C		
				Mirapex	C		
				Zerit	C		
				Viread	C		
				Sustiva	C		
				Zoloft	C		
				Calcium	C		
				Fosamax	C		
				Valium	C		

Date:05/27/05ISR Number: 4676408-6Report Type:Periodic  
Age:40 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548648A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Palpitations  
150MG Per day 6 MON  
Wellbutrin Xl PS Glaxosmithkline ORAL  
No Concurrent Medication C

Date:05/27/05ISR Number: 4676409-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0548651A  
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Bruxism		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day				Buspar	C		

Date:05/27/05ISR Number: 4676410-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0548696A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day 1 MON							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4676411-6Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0548706A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300MG Three			Accidental Overdose	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
times per day 2	DAY						
			Dysphemia				
			Feeling Jittery				
			Influenza Like Illness				
			Medication Error				
			Nausea				
			Pain				
			Restlessness				
			Tremor				
			Vomiting				

Date:05/27/05ISR Number: 4676412-8Report Type:Periodic  
 Age:33 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0548812A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Per day 7	DAY		Myalgia	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:05/27/05ISR Number: 4676763-7Report Type:Expedited (15-DaCompany Report #US-MERCK-0505USA02903  
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Adverse Event	Cogentin	PS	Merck & Co., Inc	ORAL
Initial or Prolonged			Drug Interaction	Wellbutrin Sr	SS		ORAL
				Trazodone Hydrochloride	SS		
UNKNOWN				Seroquel	SS		
UNKNOWN				Klonopin	SS		
UNKNOWN							

UNKNOWN

Trilafon

SS

Date:05/27/05ISR Number: 4677806-7Report Type:Direct  
Age: Gender:Female I/FU:I

Company Report #CTU 249801

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Bupropion Sr 150 Mg	PS		
ONE TAB BID		Feeling Jittery					

Date:05/31/05ISR Number: 4677473-2Report Type:Expedited (15-DaCompany Report #200511975US  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Exposure During Pregnancy		Allegra	PS	Aventis Pharmaceuticals Inc.	ORAL
		Haemorrhage		Bupropion	SS		
		Umbilical Cord Abnormality		Celexa	SS		ORAL
				Zantac	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/05ISR Number: 4677864-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0389788A

Age:81 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blister		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Twice		Toxic Epidermal					
per day		Necrolysis		Paxil	C	Glaxosmithkline	
60MG per day				Nexium	C		
				Premarin	C		
YR							

Date:05/31/05ISR Number: 4677867-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553362A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Atrophic Vulvovaginitis		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:05/31/05ISR Number: 4677887-0Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0378887A

Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
1TAB Twice		Aggression		Zyban	PS	Glaxosmithkline	ORAL
per day	6 DAY	Anxiety					
		Disturbance In Attention					
		Dizziness					
		Drug Withdrawal Syndrome					
		Dry Mouth					
		Headache					
		Insomnia					
		Irritability					
		Tremor					

Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety		Zyban	PS	Glaxosmithkline	ORAL
1TAB Twice		Feeling Abnormal					
per day	4 DAY	Panic Attack		No Concurrent Medication	C		
		Serotonin Syndrome					

Date:05/31/05ISR Number: 4678432-6Report Type:Direct Company Report #CTU 249981

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Sedation		Bupropion 150mg Bid	PS		ORAL
150MG PO BID							

Date:05/31/05ISR Number: 4678779-3Report Type:Expedited (15-DaCompany Report #S05-USA-01449-01

Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bleeding Peripartum	Health Professional	Celexa (Citalopram Hydrobromide)	PS		ORAL
40 MG QD PO		Drug Exposure During					
		Pregnancy		Wellbutrin			
		Pregnancy		(Bupropion			
		Premature Separation Of		Hydrochloride)	SS		ORAL
300 MG QD PO		Placenta		Wellbutrin(Bupropion			
				Hydrochloride)	SS		ORAL
150 MG QD PO							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

300 MG QD PO	Wellbutrin(Bupropion Hydrochloride)	SS	ORAL
60 MG BID PO	Allegra(Pexofenadine Hydrochloride)	SS	ORAL
150 MG QD PO	Zantac(Ranitidine Hydrochloride)	SS	ORAL

Date:05/31/05ISR Number: 4678867-1Report Type:Expedited (15-DaCompany Report #2005075887

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200 MG (200 MG , 1 IN 1 D),ORAL		Abdominal Pain	Health	Zoloft (Sertraline)	PS		ORAL
(0.5 MG AS NECESSARY), ORAL		Autism Spectrum Disorder	Professional				
		Caesarean Section					
		Drug Exposure During Pregnancy		Lorazepam (Lorazepam)	SS		ORAL
		Neonatal Disorder					
		Pregnancy					
(100 MG, AS NECESSARY), ORAL		Small For Dates Baby		Wellbutrin Sr (Bupropion Hydrochloride)	SS		ORAL
(10 MG, AS NECESSARY), ORAL				Ambien (Zolpidem Tartrate)	SS		ORAL



Date:05/31/05ISR Number: 4678965-2Report Type:Expedited (15-DaCompany Report #A0559763A  
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional	Bupropion Hydrochloride Tablet (Generic) (Bupropion Hydrochloride)	PS		ORAL
75 MG/ SEE							
IMAGE/ORAL	1	DAY		Wellbutrin Xl Tablet-Extended Release (Bupropion Hydrochloride)	SS		ORAL
SEE							
IMAGE/ORAL				Olanzapine (Olanzapine)	SS		

Date:05/31/05ISR Number: 4678972-XReport Type:Expedited (15-DaCompany Report #A0559375A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Drug Exposure During Pregnancy	Study Health Professional	Wellbutrin (Bupropion Hydrochloride)	PS		
TRANSPLACENTAL	TRANSPLACENTA	Dysphagia	Company				
RY		Neonatal Disorder	Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/05ISR Number: 4678973-1Report Type:Expedited (15-DaCompany Report #A0559357A  
 Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Aggression Anxiety Psychomotor Hyperactivity	Consumer	Wellbutrin Xl Tablet-Extended Release (Bupropion Hydrochloride)	PS		ORAL
150 MG / PER DAY/ ORAL				Verapamil	C		

Date:05/31/05ISR Number: 4678980-9Report Type:Expedited (15-DaCompany Report #A0560141A  
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Disability PER DAY/ORAL Required Intervention to Prevent Permanent Impairment/Damage		Accidental Overdose Amnesia Confusional State Disorientation Therapy Non-Responder	Health Professional	Wellbutrin Xl Tablet-Extended Release (Bupropion Hydrochloride)	PS		ORAL
				Ketorolac Trometamol Metoprolol Succinate Tramadol Hydrochloride Lortab	C C C C		

Date:05/31/05ISR Number: 4678982-2Report Type:Expedited (15-DaCompany Report #A0560085A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Road Traffic Accident	Consumer	Wellbutrin Tablet (Bupropion Hydrochloride)	PS		ORAL
150 MG/TWICE PER DAY/ ORAL							

Date:05/31/05ISR Number: 4679615-1Report Type:Expedited (15-DaCompany Report #B0380967A  
Age:63 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150 MG/ PER DAY / ORAL	Confusional State Disorientation Dizziness Malaise Pneumonia Serotonin Syndrome Sleep Disorder	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)  Amitriptyline Hcl	PS  C		ORAL

Date:05/31/05ISR Number: 4679813-7Report Type:Expedited (15-DaCompany Report #B0381635A  
Age:46 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required Intervention to Prevent Permanent 150 MG/TWICE Impairment/Damage PER DAY/ORAL	Anaphylactic Reaction	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/05ISR Number: 4679816-2Report Type:Expedited (15-DaCompany Report #A0559952A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Thrombophlebitis Thrombosis	Foreign Health Professional Company	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
ORAL			Representative	Hydromorphone Hcl Risperidone Amitriptyline	C C C		

Date:05/31/05ISR Number: 4679954-4Report Type:Expedited (15-DaCompany Report #B0382344A  
 Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required Intervention to Prevent Permanent 150 MG / Impairment/Damage		Epilepsy Facial Bones Fracture Lower Limb Fracture	Foreign Consumer Other	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		

Date:05/31/05ISR Number: 4679976-3Report Type:Expedited (15-DaCompany Report #B0381636A  
 Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required Intervention to Prevent Permanent 150 MG/TWICE Impairment/Damage PER DAY/ ORAL		Dermatitis Allergic Drug Abuser Drug Administration Error Joint Swelling	Foreign Health Professional	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
				Cannabis	C		

Date:06/02/05ISR Number: 4679353-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0559798A  
 Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Induced		Bupropion	PS	Glaxosmithkline	ORAL
300MG Per day							

Date:06/02/05ISR Number: 4679354-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0559798B  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Anencephaly		Bupropion	PS	Glaxosmithkline	
300MG Per day							
Congenital Anomaly		Drug Exposure During Pregnancy					

Date:06/02/05ISR Number: 4679379-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0381871A  
 Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Emotional Distress		Zyban	PS	Glaxosmithkline	ORAL
50 DAY							
Other		Psoriasis		Acitretin	C		
UNKNOWN	50MG per day						
				Ciclosporin	C		
UNKNOWN	400MG per day						
				Amlodipine	C		
UNKNOWN	10MG per day						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/02/05ISR Number: 4679842-3Report Type:Periodic  
 Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540083A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus		Augmentin	PS	Glaxosmithkline	ORAL
1000MG Twice							
per day	3	DAY		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
300MG Per day	3	WK		No Concurrent Medication	C		

Date:06/02/05ISR Number: 4680005-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0560408A  
 Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disturbance In Sexual		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Arousal		No Concurrent Medication	C		
300MG Per day	27	DAY					
		Loss Of Libido					
		Prostate Cancer					

Date:06/02/05ISR Number: 4680033-0Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0380967A  
 Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization -		Disorientation		Endep	C	Glaxosmithkline	
150MG Per day		50MG Variable					
Initial or Prolonged		Dizziness					
UNKNOWN		Fall		Androcur	C		
dose		50MG Per day		Bricanyl Turbohaler	C		
UNKNOWN		Malaise					
RESPIRATORY		Pneumonia					
(INHALATION)		2PUFF Four					

	Serotonin Syndrome				
times per day					
UNKNOWN	Sleep Disorder		Diamicron	C	
	80MG Twice				
per day					
UNKNOWN			Ditropan	C	
	5MG Twice per				
day					
UNKNOWN			Epilim	C	
	500MG Three				
times per day					
UNKNOWN			Naprosyn Sr	C	
	1000MG See				
dosage text					
UNKNOWN			Panadeine Forte	C	Glaxosmithkline
UNKNOWN			Progynova	C	
	2MG Twice per				
day					
RESPIRATORY			Seretide	C	Glaxosmithkline
(INHALATION)					
	1PUFF Twice				
per day					
UNKNOWN			Temazepam	C	
	10MG At night				

Date:06/02/05ISR Number: 4680044-5Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0383114A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Crying		Zyban	PS	Glaxosmithkline	
UNKNOWN		Depression		No Concurrent			
		Eating Disorder		Medication	C		
		Insomnia					
		Suicidal Ideation					
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/02/05ISR Number: 4681499-2Report Type:Expedited (15-DaCompany Report #IMP\_0865\_2005  
Age:57 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Drug Interaction	Literature	Bupropion	PS		
Initial or Prolonged	Haemodynamic Instability	Health	Linezolid	SS		
	Hypertension	Professional				
	Hypertensive Crisis					

Date:06/02/05ISR Number: 4681820-5Report Type:Expedited (15-DaCompany Report #IMP\_0864\_2005  
Age:62 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Alanine Aminotransferase	Foreign	Bupropion	PS		
300 MG QDAY 3 WK						
Initial or Prolonged	Increased	Literature	Sertraline	SS		
50 MG QDAY 17 DAY						
Required	Aspartate	Health	Piracetam	SS		
2400 MG QDAY 17 DAY						
Intervention to	Aminotransferase	Professional	Venlafaxine	SS		
75 MG QDAY 17 DAY						
Prevent Permanent Impairment/Damage	Increased Autonomic Nervous System Imbalance Balance Disorder Blood Pressure Fluctuation Cerebral Atrophy Clumsiness Confusional State Coordination Abnormal Depressed Level Of Consciousness Drug Interaction Inhibition Electrocardiogram Qt Prolonged Electrocardiogram Repolarisation Abnormality Faecal Incontinence Fall Gait Disturbance	Other				



Hallucinations, Mixed  
Heart Rate Irregular  
Hyperhidrosis  
Insomnia  
Lethargy  
Memory Impairment  
Mental Impairment  
Myoclonus  
Psychomotor Agitation  
Pyrexia  
Serotonin Syndrome  
Tremor  
Urinary Incontinence

Date:06/03/05ISR Number: 4681063-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553777A  
Age:59 YR Gender:Female I/FU:F

Outcome PT  
Other Atrial Flutter  
Eye Pain  
Headache

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
4	MON	Heart Rate Increased Hyperhidrosis Hypertension				
		Nausea	Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Neuralgia	Benicar	C		
		Photopsia	Skelaxin	C		
		Red Blood Cell Count Decreased	Protonix	C		
		Vitreous Floaters	Potassium Supplement	C		
		Vomiting	Celebrex	C		
		White Blood Cell Count Decreased	Vicodin	C		
			Maxalt	C		
			Prednisolone Acetate	C		
			Betanol 0.3%	C		
			Alphagan	C		
			Ditropan	C		
			Benadryl	C	Glaxosmithkline	
			Zyrtec	C	Glaxosmithkline	
			Gerovit (Multivitamin)	C		
			Tylenol	C	Glaxosmithkline	

Date:06/03/05ISR Number: 4681066-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0558861A  
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	450MG Per day	Suicidal Ideation		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Hospitalization -	125MG Per day	Suicide Attempt		Lamictal	C	Glaxosmithkline	ORAL
Initial or Prolonged	100MG Per day			Topamax	C		ORAL
	.5MG Twice			Clonazepam	C		
	per day						

Date:06/03/05ISR Number: 4681069-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0560822A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Disability Other		Diarrhoea Pancreatic Enzymes Increased Weight Decreased		Bupropion	PS	Glaxosmithkline	ORAL

Date:06/03/05ISR Number: 4681080-5Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0381635A  
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day	15 DAY	Anaphylactic Reaction Joint Swelling Oedema Peripheral Orbital Oedema Pruritus Rash Generalised Urticaria		Zyban	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/03/05ISR Number: 4684598-4Report Type:Expedited (15-DaCompany Report #2005017220  
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	25 MG (25 MG, 1 IN 1 D)	Abdominal Distension	Consumer	Zoloft (Sertraline)	PS		
		Alopecia	Health				
		Body Height Decreased	Professional	Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
		Dry Skin					
		Eructation					
		Flatulence		Klonopin (Clonazepam)	C		
		Glossodynia					
		Movement Disorder					
		Nervous System Disorder					
		Self Mutilation					

Date:06/06/05ISR Number: 4682634-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0560943A  
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day 3 MON	General Physical Health		Wellbutrin	PS	Glaxosmithkline	ORAL
		Deterioration		No Concurrent Medication	C		

Date:06/06/05ISR Number: 4682689-5Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0383467A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	UNKNOWN	Angina Pectoris		Zyban	PS	Glaxosmithkline	

Date:06/06/05ISR Number: 4685803-0Report Type:Direct Company Report #CTU 250458  
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin Sr 150 Mg (Unk If Generic Dispensed)	PS		ORAL
150 MG QD X 7							
THEN BID PO							

Date:06/06/05ISR Number: 4686476-3Report Type:Expedited (15-DaCompany Report #2005079514  
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Pressure Increased	Consumer	Bextra (Valdecoxib)	PS		ORAL
20 MG(10 MG,		Cellulitis					
2 IN 1 D),		Erysipelas					
ORAL		Eye Allergy		Benadryl Injection	SS		
		Hypersensitivity		(Diphenhydramine)			
INTRAVENOUS	INTRAVENOUS	Pain		Lisinopirl	SS		ORAL
20 MG (20 MG,		Pharyngeal Oedema		(Lisinopril)			
1 IN 1 D),		Pharyngolaryngeal Pain					
ORAL		Urticaria					
				Cortisone	SS		
INTRAVENOUS	INTRAVNEOUS			(Cortisone)			
300 MG (150				Wellbutrin	SS		ORAL
MG, 2 IN 1				(Bupropion			
D), ORAL				Hydrochloride)			

Freedom Of Information (FOI) Report

ORAL Toprol (Metoprolol) SS ORAL

Date:06/06/05ISR Number: 4686519-7Report Type:Expedited (15-DaCompany Report #2005080150  
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Consumer	Celebrex (Celecoxib)	PS		
		Feeling Abnormal		Paxil (Paroxetine			
		Laboratory Test Abnormal		Hydrochloride)	SS		
		Nasopharyngitis		Wellbutrin			
		Pain		(Bupropion			
		Personality Disorder		Hydrochloride)	SS		
		Retching		Aleve (Naproxen			
		Vomiting		Sodium)	SS		
				Avalide			
				(Hydrochloride,			
				Irbesartan)	C		
				Acetylsalicylic Acid			
				(Acetylsalicylic			
				Acid)	C		
				Multivitamins			
				(Multivitamins)	C		
				Atenolol (Atenolol)	C		

Date:06/07/05ISR Number: 4684149-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0560956A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability							
300MG Per day	3 YR			Wellbutrin	PS	Glaxosmithkline	ORAL
		Convulsion					
		Head Injury		Alcohol	C		

Date:06/07/05ISR Number: 4684181-0Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0380880A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 150MG Per day Initial or Prolonged	Anxiety Blood Pressure Increased Chest Pain Palpitations Sense Of Oppression Tachycardia Tension	Zyban	PS	Glaxosmithkline	ORAL
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Date:06/08/05ISR Number: 4685361-0Report Type:Expedited (15-DaCompany Report #13296  
 Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 300MG Per day Hospitalization - Initial or Prolonged Other	Apnoea Cyanosis Drug Exposure During Pregnancy Drug Withdrawal Syndrome Respiratory Distress		Wellbutrin Tricyclic Antidepressant Protonix Multivitamins	PS  C C C	Glaxosmithkline	

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Freedom Of Information (FOI) Report

Date:06/08/05ISR Number: 4685392-0Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0378949A  
Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Twice per day UNKNOWN	13 DAY	Anxiety Hallucination Insomnia Panic Reaction Paranoia Paraplegia Rash		Zyban Alprazolam	PS C	Glaxosmithkline	ORAL

Date:06/08/05ISR Number: 4685393-2Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0380967A  
Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Per day Initial or Prolonged UNKNOWN dose UNKNOWN RESPIRATORY (INHALATION) times per day UNKNOWN per day UNKNOWN day UNKNOWN	3 WK 50MG Variable 50MG Per day 2PUFF Four 80MG Twice 5MG Twice per 500MG Three	Confusional State Disorientation Dizziness Fall Malaise Pneumonia Serotonin Syndrome Sleep Disorder		Zyban Endep Androcur Bricanyl Turbohaler Diamicron Ditropan Epilim	PS C C C C	Glaxosmithkline Glaxosmithkline	ORAL



times per day

UNKNOWN 1000MG See

Naprosyn Sr C

dosage text

UNKNOWN

Panadeine Forte C Glaxosmithkline

UNKNOWN 2MG Twice per

Progynova C

day

RESPIRATORY

Seretide C Glaxosmithkline

(INHALATION) 1PUFF Twice

per day

UNKNOWN 10MG At night

Temazepam C

Date:06/09/05ISR Number: 4686230-2Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0561473A

Age:18 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Paxil	PS	Glaxosmithkline	
Other		Drug Interaction		Wellbutrin Sr	SS	Glaxosmithkline	
		Drug Withdrawal Syndrome					
15	DAY	Hyperhidrosis					
		Hypoaesthesia					
		Paraesthesia					
		Tremor					

Date:06/09/05ISR Number: 4687440-0Report Type:Direct Company Report #CTU 250815

Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Ultram 50 Mg	PS		
Other		Fall					
50 MG 2 TID		Grand Mal Convulsion					
PRN		Loss Of Consciousness		Wellbutrin Sr 150 Mg	SS		
150 MG BID							



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/10/05ISR Number: 4687979-8Report Type:Expedited (15-DaCompany Report #PHEH2005US06302

Age:10 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Trileptal	PS	Novartis Sector: Pharma	ORAL
475 mg, QD		Electroencephalogram					
		Abnormal		Wellbutrin (Xl)	SS		ORAL
300 mg, QD				Wellbutrin (Xl)	SS		ORAL
150 mg, QD				Dexedrine			
				"Smithkline Beecham"	C		ORAL
5 mg, BID				Abilify	C		
5 mg, UNK							

Date:06/10/05ISR Number: 4688018-5Report Type:Expedited (15-DaCompany Report #2005UW08423

Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anger		Seroquel	PS	Zeneca Pharmaceutical	ORAL
		Crying					ORAL
		Depression		Wellbutrin Xl	SS		ORAL
		Intentional Self-Injury		Risperdal	C		
		Laceration		Klonopin	C		
		Poisoning		Paxil	C		ORAL
9 YR		Psychotic Disorder					
		Suicide Attempt					

Date:06/10/05ISR Number: 4688211-1Report Type:Expedited (15-DaCompany Report #AU-ABBOTT-05P-008-0302257-00

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Epilim	PS		
UNKNOWN							
		Depression		Bupropion			
		Overdose		Hydrochloride	SS		
UNKNOWN							

UNKNOWN

Temazepam SS

Diazepam SS

Ipratropium Bromide SS

UNKNOWN

Fluticasone

Propionate SS

UNKNOWN

Hydrochlorothiazide SS

UNKNOWN

Salbutamol SS

UNKNOWN

Date:06/13/05ISR Number: 4688800-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0548205A

Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	300MG Per day 74 DAY	Amnesia		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged		Confusional State Grand Mal Convulsion Incorrect Dose Administered Medication Error Overdose Temporal Lobe Epilepsy		Lexapro	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/13/05ISR Number: 4688811-9Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0561484A  
 Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Affect Lability	Paroxetine	PS	Glaxosmithkline	ORAL
20MG Per day							
			Agitation	Bupropion	SS	Glaxosmithkline	ORAL
150MG Per day	1 WK		Blood Pressure Increased	Trazodone	SS		ORAL
50MG Unknown							
			Carotid Artery Thrombosis	Contraceptive	C		
			Cerebrovascular Accident	Lithium Carbonate	C	Glaxosmithkline	
			Delirium				
			Disorientation				
			Hyperreflexia				
			Muscular Weakness				
			Pyrexia				
			Serotonin Syndrome				
			Tachycardia				

Date:06/13/05ISR Number: 4688817-XReport Type:Expedited (15-DaCompany Report #13148  
 Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Tachycardia	Bupropion	PS	Glaxosmithkline	
150MG Per day							
				Unspecified Ssri	C		

Date:06/13/05ISR Number: 4688818-1Report Type:Expedited (15-DaCompany Report #13149  
 Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Abortion Spontaneous	Bupropion	PS	Glaxosmithkline	
300MG Per day							
				Unspecified Ssri	C		

Date:06/13/05ISR Number: 4688833-8Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0380012A  
 Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema		Zyban	PS	Glaxosmithkline	ORAL
10 DAY		Dry Mouth		Thyroxine	C	Glaxosmithkline	
		Hyperhidrosis		Diphenhydramine			
		Ill-Defined Disorder		Hydrochloride	C	Glaxosmithkline	
10MG per day		Mouth Ulceration		Prednisolone	C	Glaxosmithkline	
5MG per day		Oral Pain		Furosemide	C	Glaxosmithkline	
20MG per day		Swelling Face					
		Tremor					

Date:06/13/05ISR Number: 4688834-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0380740A

Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Analgesic Effect		Zyban	PS	Glaxosmithkline	ORAL
300MG per day		Drug Abuser		Imiject	SS	Glaxosmithkline	
SUBCUTANEOUS	4INJ per day	Therapeutic Response		No Concurrent			
		Unexpected		Medications	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/13/05ISR Number: 4688854-5Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0384025A  
 Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema Cholestasis Cytolytic Hepatitis		Zyban	PS	Glaxosmithkline	ORAL

Date:06/13/05ISR Number: 4690182-9Report Type:Expedited (15-DaCompany Report #2005-01-1690  
 Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Depression	Health	Peg-Intron			
Hospitalization - Initial or Prolonged		Haemorrhagic Stroke	Professional Company	(Peginterferon Alfa-2b) Redipen	PS		
SUBCUTANEOUS	150 MCG QWK		Representative				
SUBCUTANEOUS				Copegus (Ribavirin) Capsules	SS		ORAL
1200 MG QD							
ORAL							
SUBCUTANEOUS	180 MCG QWK			Pegasys (Pegylated Interferon Alfa-2a) Injectable	SS		
SUBCUTANEOUS							
100 MG TID				Wellbutrin (Bupropion)	SS		ORAL
ORAL							
				Percocet	C		
				.....	C		
				Protonix (Pantoprazole Sodium)	C		
				Percodan	C		
				Procrit Injectable	C		
				Diazepam	C		
				Lorazepam	C		

Date:06/14/05ISR Number: 4690738-3Report Type:Direct  
 Age:33 YR Gender:Male I/FU:I

Company Report #CTU 251070

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 100MG	Convulsion		Bupropion	100mg	PS	ORAL
Hospitalization - TWICE DAILY	Life Support					
Initial or Prolonged ORAL	Treatment Noncompliance					
Required Intervention to Prevent Permanent Impairment/Damage			Cyclobenzaprine Hcl		C	
			Diazepam		C	
			Fluoxetine Hcl		C	
			Gabapentin		C	
			Hydrochlorothiazide		C	
			Irbesartan		C	
			Loratadine		C	
			Omeprazole		C	
			Topiramate		C	
			Verapamil Hcl		C	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/15/05ISR Number: 4690270-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0561914A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	100MG Unknown	1 WK	Muscle Spasms	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
Other	150MG Unknown		Muscle Twitching	Wellbutrin Xl	SS	Glaxosmithkline	ORAL

Date:06/15/05ISR Number: 4690305-1Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0384155A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	UNKNOWN	7 DAY	Anxiety	Zyban	PS	Glaxosmithkline	
			Depression				
			Smoker				
			Suicidal Ideation				

Date:06/15/05ISR Number: 4690626-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0511243A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG See		Completed Suicide	Wellbutrin Sr	PS	Glaxosmithkline	
dosage text			Decreased Appetite				
			Depressed Mood				
			Disturbance In Attention				
			Feelings Of Worthlessness				
			Gun Shot Wound				
			Head Injury				
			Major Depression				
			Sleep Disorder				
			Social Avoidant Behaviour				
			Suicidal Ideation				
			Treatment Noncompliance				
			Tremor				

Date:06/15/05ISR Number: 4690628-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0535145A  
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Constipation		Wellbutrin XL	PS	Glaxosmithkline	
150MG Per day	8 DAY	Proctalgia		Concerta	C		
		Rectal Haemorrhage					
		Stool Analysis Abnormal					

Date:06/15/05ISR Number: 4690630-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0552342A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Amnesia		Bupropion	PS	Glaxosmithkline	ORAL
150MG Twice							
Initial or Prolonged		Aura					
per day	21 MON	Convulsion		Paxil	C	Glaxosmithkline	
				Clonazepam	C		
1MG Per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/15/05ISR Number: 4690633-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0557288A  
 Age:14 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Contusion		Wellbutrin	PS	Glaxosmithkline	ORAL
306 DAY		Grand Mal Convulsion		Zyprexa	C		ORAL
10MG Per day		Incontinence		Zoloft	C		ORAL
50MG Per day		Scratch					
		Somnolence					

Date:06/15/05ISR Number: 4690638-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0562131A  
 Age:25 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Consciousness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day				Depakote	C		
				Adderall	C		
				Xanax	C		

Date:06/15/05ISR Number: 4690639-0Report Type:Expedited (15-DaCompany Report #CL-GLAXOSMITHKLINE-A0562265A  
 Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Laryngeal Oedema		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
150MG See		Overdose					
Hospitalization - dosage text		Suicide Attempt					
Initial or Prolonged							
Other							

Date:06/15/05ISR Number: 4691416-7Report Type:Expedited (15-DaCompany Report #2005051512  
 Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 400 MG (200 Initial or Prolonged MG, 2 IN 1 Other D), ORAL		Anxiety	Consumer	Celebrex (Celecoxib)	PS		ORAL
		Back Pain	Health				
		Depression	Professional				
		Nervousness Suicidal Ideation		Wellbutrin (Bupropion Hydrochloride)	SS		ORAL
300 MG (150 MG, 2 IN 1 D), ORAL				Lexapro (Escitalopram)	C		
				Trazadone (Trazadone)	C		
				Advair (Fluticasone Propionate, Salmeterol Xinafoate)	C		
				All Other Therapeutic Products (All Othr Therapeutic Products)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/15/05ISR Number: 4691865-7Report Type:Direct  
 Age: Gender:Male I/FU:I

Company Report #CTU 251230

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anger		Bupropriion Sr 200			
PO BID		Condition Aggravated		Mg	PS		ORAL
		Therapeutic Response		Zanaflex	C		
		Unexpected With Drug		Neurontin	C		
		Substitution		Catapress	C		
				Tetracycline	C		
				Topical Clotrimazole	C		
				Levaquin	C		
				Macrodantin	C		
				Suprofen	C		
				Envlose Fibercon	C		

Date:06/15/05ISR Number: 4693851-XReport Type:Expedited (15-DaCompany Report #S05-USA-02982-01  
 Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -							
Initial or Prolonged		Convulsion	Consumer	Lexapro			
30 MG QD PO		Dizziness		(Escitalopram)	PS		ORAL
		Dyskinesia		Lexapro			
10 MG QD PO		Fall		(Escitalopram)	SS		ORAL
		Fatigue		Lexapro			
		Head Injury		(Escitalopram)	SS		
		Headache		Lexapro			
40 MG QD PO				(Escitalopram)	SS		ORAL
				Wellbutrin			
				(Bupropion			
				Hydrochloride)	SS		

Date:06/16/05ISR Number: 4694005-3Report Type:Expedited (15-DaCompany Report #IMP\_0847\_2005  
 Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Anxiety	Consumer	Bupropion	PS	ORAL
1 TAB QDAY PO					
	Crying		Lexapro	C	
	Depression		Lorazepam	C	
	Paranoia		Calcium Citrate	C	
	Pharmaceutical Product		Glucosamine		
	Complaint		Chondrotin	C	
	Suicidal Ideation		Multivitamin	C	

Date:06/17/05ISR Number: 4692725-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0558531A  
Age:48 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Alcohol Withdrawal		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 23 DAY						
Hospitalization -	Syndrome		Alcohol	SS		
Initial or Prolonged	Amnesia		Benicar	C		
Other	Extradural Haematoma					
	Fall					
	Grand Mal Convulsion					
	Head Injury					
	Skull Fracture					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/17/05ISR Number: 4693277-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0558376A

Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eating Disorder		Wellbutrin	PS	Glaxosmithkline	ORAL
450MG Per day		Emotional Disorder		Crack Cocaine	SS		
		Flight Of Ideas		Alcohol	SS		
		Grand Mal Convulsion		Valium	C		
		Incontinence		Xanax	C		
1MG Twice per		Logorrhoea					
day		Overdose					
		Stress					
		Tearfulness					
		Tongue Biting					

Date:06/17/05ISR Number: 4693278-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0558504A

Age:16 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
1800MG Per		Intentional Misuse					
day	1 DAY	Somnolence		Albuterol	C	Glaxosmithkline	
				Advair	C	Glaxosmithkline	
				Singulair	C		

Date:06/17/05ISR Number: 4693279-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0558505A

Age:17 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
900MG Per day	1 DAY	Intentional Misuse		Albuterol	C	Glaxosmithkline	
		Nausea		Advair	C	Glaxosmithkline	
		Overdose					
		Tremor					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Antepartum Haemorrhage	Health	Celexa (Citalopram			
40 MG QD PO		Drug Exposure During	Professional	Hydrobromide)	PS		ORAL
		Pregnancy		Wellbutrin			
		Haemorrhage		(Bupropion			
300 MG QD PO		Placental Disorder		Hydrochloride)	SS		ORAL
		Pregnancy		Wellbutrin			
		Umbilical Cord		(Bupropion			
150 MG QD PO		Abnormality		Hydrochloride)	SS		ORAL
				Wellbutrin			
				(Bupropion			
300 MG QD PO				Hydrochloride)	SS		ORAL
				Allegra			
				(Fexofenadine			
60 MG BID PO				Hydrochloride)	SS		ORAL
				Zantac (Ranitidine			
150 MG QD PO				Hydrochloride)	SS		ORAL



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Freedom Of Information (FOI) Report

Date:06/20/05ISR Number: 4694661-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0562517A  
 Age:19 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Injury		Ethanol	C		ORAL

Date:06/20/05ISR Number: 4694665-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0563187A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Eskalith	PS	Glaxosmithkline	ORAL
		Road Traffic Accident		Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Per day	1	MON		Unknown	C		

Date:06/20/05ISR Number: 4694692-XReport Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0384877A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cerebral Haemorrhage		Zyban	PS	Glaxosmithkline	
UNKNOWN	150MG	Twice					
		Cerebrovascular Accident					
per day	6	DAY					
		Convulsion					
		Death					

Date:06/21/05ISR Number: 4695995-5Report Type:Expedited (15-DaCompany Report #2005-01-1690  
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Haemorrhagic Stroke		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Three							
Hospitalization -							
times per day	413	DAY					
Initial or Prolonged				Pegylated Interferon			

SUBCUTANEOUS 180MCG Weekly 396 DAY  
 1200MG Per  
 day 413 DAY  
 SUBCUTANEOUS 150MCG Weekly  
 .175MG Per  
 day  
 1MG Three  
 times per day  
 40MG Per day  
 325MG Twice  
 per day 101 DAY  
 SUBCUTANEOUS 40000U Weekly 277 DAY  
 5MG Twice per  
 day 413 DAY

Alpha 2a SS  
 Copegus SS ORAL  
 Pegylated Interferon  
 Alpha 2b SS  
 Percocet C  
 Benzodiazepine C  
 Levoxyl C Glaxosmithkline ORAL  
 Lorazepam C ORAL  
 Protonix C ORAL  
 Percodan C ORAL  
 Procrit C  
 Diazepam C ORAL

Date:06/21/05ISR Number: 4696002-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0563000A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150MG Per day 2 DAY	Dystonia		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Feeling Abnormal Speech Disorder		No Concurrent Medication	C		

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Freedom Of Information (FOI) Report

Date:06/21/05ISR Number: 4696030-5Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0384576A

Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspnoea		Zyban	PS	Glaxosmithkline	
11	DAY						
		Heart Rate Increased					
		Hunger					
		Loss Of Consciousness					
		Panic Attack					

Date:06/21/05ISR Number: 4697783-2Report Type:Direct Company Report #CTU 251631

Age:81 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anorexia		Bupropion Sa 100 Mg	PS		
1 TAB BID							
Initial or Prolonged		Confusional State		Fluoxetine 20 Mg	SS		
2 CAPS DAILY							
		Dehydration					
		Serotonin Syndrome					
		Urinary Tract Infection					

Date:06/22/05ISR Number: 4697237-3Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0385105A

Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Chest Pain		Zyban	PS	Glaxosmithkline	
UNKNOWN	5						DAY
Initial or Prolonged		Dizziness		Valerian	C	Glaxosmithkline	
		Heart Alternation					
		Hypertension					
		Palpitations					

Date:06/22/05ISR Number: 4697683-8Report Type:Expedited (15-DaCompany Report #2005080150

Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depressed Mood	Consumer	Celebrex (Celecoxib)	PS		
		Dizziness		Paxil (Paroxetine			
		Feeling Abnormal		Hydrochloride)	SS		
		Flat Affect		Wellbutrin			
		Multiple Sclerosis		(Bupropion			
		Nasopharyngitis		Hydrochloride)	SS		
		Pain		Aleve (Naproxen			
		Personality Disorder		Sodium)	SS		
		Retching		Avalide			
		Vomiting		(Hydrochlorothiazide			
				, Irbesartan)	C		
				Acetylsalicylic Acid			
				(Acetylsalicylic			
				Acid)	C		
				Multivitamins			
				(Multivitamins)	C		
				Atenolol (Atenolol)	C		

Date:06/23/05ISR Number: 4698146-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0559188A  
Age: Gender:Female I/FU:F

Outcome	PT
Other	Drug Exposure During Pregnancy

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Jaundice Neonatal  
Premature Baby

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
450MG per day			Wellbutrin	PS	Glaxosmithkline	
20MG per day			Lexapro	C		

Date:06/23/05ISR Number: 4698147-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0559798A  
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Abortion Induced	Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day			Stillbirth				

Date:06/23/05ISR Number: 4698148-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0559798B  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Abortion Induced	Wellbutrin	PS	Glaxosmithkline	
300MG Per day			Anencephaly				
Congenital Anomaly			Drug Exposure During Pregnancy				
			Stillbirth				

Date:06/23/05ISR Number: 4698149-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0562405A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening			Adverse Event	Wellbutrin	PS	Glaxosmithkline	ORAL
2 YR			Anoxic Encephalopathy	Xanax	SS		
Hospitalization -			Cardiac Arrest	Alcohol	SS		
Initial or Prolonged			General Physical Health				
Disability			Deterioration				
Other							

Grand Mal Convulsion  
Hypotension  
Myocardial Infarction

Date:06/23/05ISR Number: 4698150-8Report Type:Expedited (15-DaCompany Report #2005-06-0342  
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dehydration		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice		Diarrhoea					
per day		Haematochezia		Ativan	SS		
UNKNOWN		Hallucination, Visual		Pegylated Interferon			
		Headache		Alpha 2b	SS		
SUBCUTANEOUS		Neck Pain		Rebetol	SS		ORAL
		Vomiting					

Date:06/23/05ISR Number: 4698153-3Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0563226A  
Age:50 YR Gender:Female I/FU:I

Outcome	PT
Other	Convulsion
	Dizziness
	Drug Withdrawal Syndrome
	Dyskinesia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fatigue Loss Of Control Of Legs Malaise					
150MG Twice per day		Nausea		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
300MG per day		Tremor		Topamax	C		ORAL
10MG At night	3 YR			Clonazepam	C		ORAL

Date:06/23/05ISR Number: 4698155-7Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0563485A  
Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 150MG Single dose	1 DAY	Convulsion Pallor Sensory Disturbance		Zyban	PS	Glaxosmithkline	ORAL
				Psorex	C	Glaxosmithkline	ORAL
				Triptanol	C		ORAL
				Aspirin	C	Glaxosmithkline	ORAL
				Buscopan	C		
				Lisador	C		

Date:06/23/05ISR Number: 4698165-XReport Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0383114A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other UNKNOWN		Crying		Zyban	PS	Glaxosmithkline	
		Depression Eating Disorder Insomnia Suicidal Ideation Tremor		No Concurrent Medication	C		

Date:06/23/05ISR Number: 4698166-1Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0383245A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
Other		Crying Depression Eating Disorder Insomnia Nausea Suicidal Ideation Tremor		No Concurrent Medication	C		

Date:06/23/05ISR Number: 4698171-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0384598A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
Death		Cardiac Arrest					
750MG per day							
Hospitalization - Initial or Prolonged		Depressed Level Of Consciousness Grand Mal Convulsion Hypothermia Intentional Misuse Status Epilepticus		Alcohol	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/23/05ISR Number: 4699383-7Report Type:Direct  
Age:43 YR Gender:Female I/FU:I

Company Report #CTU 251868

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia		Wellbutrin Sr 150			
		Confusional State		Mgm Q Am, 75 Mgm Q			
		Depressed Level Of		Pm	PS		
		Consciousness		Lexapro 20 Mgm Q Am,			
		Disorientation		10 Mg Mgm Q Pm	SS		
		Road Traffic Accident					
		Status Epilepticus					

Date:06/24/05ISR Number: 4699282-0Report Type:Expedited (15-DaCompany Report #13611  
Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Deficiency Anaemia		Bupropion	PS	Glaxosmithkline	
Other		Gastrointestinal		Prenatal Vitamins	C		
1TAB Per day		Infection		Calcium	C		
1200MG Twice		Polyhydramnios					
per day							

Date:06/24/05ISR Number: 4699283-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0559914A  
Age:16 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Other		Fall		Geodon	C		
54 DAY		Grand Mal Convulsion		Provigil	C		ORAL
200MG Per day				Ambien	C		ORAL

Date:06/24/05ISR Number: 4699288-1Report Type:Expedited (15-DaCompany Report #S05-USA-02982-01  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 30MG Per day		Convulsion Dizziness		Wellbutrin Lexapro	PS SS	Glaxosmithkline	ORAL ORAL
		Dyskinesia Epilepsy Fall Fatigue Head Injury Postictal Headache					

Date:06/24/05ISR Number: 4699289-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0563222A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Feeling Abnormal Suicidal Ideation		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:06/24/05ISR Number: 4699323-0Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0383467A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN		Angina Pectoris Chest Discomfort Pregnancy		Zyban No Concurrent Medication	PS C	Glaxosmithkline	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/24/05ISR Number: 4699329-1Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0384603A

Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis Bullous		Zyban	PS	Glaxosmithkline	ORAL
23 DAY				Co-Amoxiclav	C	Glaxosmithkline	ORAL
2 DAY				Omeprazole	C	Glaxosmithkline	
UNKNOWN	20MG per day	3 WK		Diclofenac Sodium	C	Glaxosmithkline	ORAL
50MG Per day							

Date:06/24/05ISR Number: 4699330-8Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0384604A

Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Orthopnoea		Zyban	PS	Glaxosmithkline	ORAL
150MG per day	1 MON			Ibuprofen	C	Glaxosmithkline	ORAL
400MG per day							
		Sleep Disorder					

Date:06/24/05ISR Number: 4699334-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0384922A

Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Coordination Abnormal		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	11 DAY						
		Road Traffic Accident					

Date:06/24/05ISR Number: 4700967-8Report Type:Expedited (15-DaCompany Report #B0384487A

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Hypersensitivity	Study Literature	Zyban Tablet-Zyban (Bupropion			

150 MG / PER Health Hydrochloride) PS ORAL  
DAY / ORAL WK Professional

Date:06/27/05ISR Number: 4700047-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0563807A  
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Drug Exposure During Pregnancy Nervous System Disorder		Wellbutrin No Concurrent Medication	PS C	Glaxosmithkline	

Date:06/27/05ISR Number: 4700550-4Report Type:Direct Company Report #CTU 252005  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG BID ORAL		Dysgeusia Feeling Abnormal Nausea Pharmaceutical Product Complaint		Bupropion Sr 150mg Teva Pharma	PS	Teva Pharma	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/27/05ISR Number: 4701458-0Report Type:Expedited (15-DaCompany Report #HQWYE382713JUN05

Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest Diarrhoea Drug Interaction	Literature	Effexor (Venlafaxine Hydrochloride, Tablet)	PS		
OVERDOSE		Dyspnoea					
AMOUNT		Hepatic Congestion Hepatic Steatosis		Manerix (Moclobemide)	SS		
OVERDOSE		Hyperhidrosis Multiple Drug Overdose Pulmonary Congestion Pulmonary Oedema Respiratory Arrest Serotonin Syndrome		Trimipramine (Trimipramine) Wellbutrin (Bupropion Hydrochloride)	SS SS SS		

Date:06/27/05ISR Number: 4701808-5Report Type:Expedited (15-DaCompany Report #2005SP001183

Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1X; ORAL		Multiple Drug Overdose	Health	Lunesta (3 Mg)	PS		ORAL
Initial or Prolonged IX,			Professional	Trandolapril	SS		
IX;				Verapamil	SS		
IX,				Avandia	SS		
IX;				Metformin	SS		
IX,				Lasix	SS		
IX,				Wellbutrin Xl	SS		
IX;				Alcohol	SS		

Date:06/28/05ISR Number: 4701131-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0563630A  
Age:16 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Per day 4 MON Initial or Prolonged	Abnormal Behaviour		Wellbutrin	PS	Glaxosmithkline	ORAL
	Blood Creatine Phosphokinase Increased Dementia Hot Flush		Methylphenidate	C		

Date:06/28/05ISR Number: 4701133-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0563834A  
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Feeling Abnormal		Wellbutrin	PS	Glaxosmithkline	ORAL
			Trazodone	SS		

Date:06/28/05ISR Number: 4701134-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0563842A  
Age:60 YR Gender:Female I/FU:F

Outcome	PT
Other	Anaemia Anorexia Asthenia Chest Pain Colitis Eating Disorder Gastritis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Malaise Malnutrition Red Blood Cell				
		Sedimentation Rate Increased Weight Decreased	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:06/28/05ISR Number: 4701135-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0563865A  
Age: Gender:Male I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose Other		PT Anxiety Faecal Incontinence Psychotic Disorder	Wellbutrin	PS	Glaxosmithkline	ORAL

Date:06/28/05ISR Number: 4701145-9Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0381636A  
Age:21 YR Gender:Male I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose Disability 150MG Twice Other per day	18 DAY	PT Dermatitis Allergic Joint Swelling Pharmaceutical Product Complaint Rash Stevens-Johnson Syndrome Urticaria	Zyban   Marijuana Cannabis	PS   C C	Glaxosmithkline	ORAL

Date:06/28/05ISR Number: 4701953-4Report Type:Expedited (15-DaCompany Report #US-MERCK-0304USA00834  
Age:63 YR Gender:Male I/FU:F

Outcome	Report Source
Hospitalization - Initial or Prolonged Disability Other	PT Acute Myocardial Infarction Angina Unstable Anxiety Arthralgia

Arthropathy  
Back Pain  
Benign Prostatic  
Hyperplasia  
Cardiac Failure  
Congestive  
Chest Pain  
Coronary Artery Disease  
Coronary Artery  
Restenosis  
Depression  
Diabetes Mellitus  
Non-Insulin-Dependent  
Diabetic Nephropathy  
Diabetic Neuropathy  
Diabetic Retinopathy  
Drug Hypersensitivity  
Drug Ineffective  
Fall  
Hepatitis  
Herpes Zoster  
Hypercholesterolaemia



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Infection				
		Injury				
		Intervertebral Disc				
		Degeneration	Vioxx	PS	Merck & Co., Inc	ORAL
10 DAY		Intervertebral Disc	Vioxx	SS	Merck & Co., Inc	ORAL
146 DAY		Disorder	Vioxx	SS	Merck & Co., Inc	ORAL
		Ischaemic Cardiomyopathy	Vioxx	SS	Merck & Co., Inc	ORAL
		Oedema Peripheral	Zyban	SS		
UNKNOWN		Pneumonia	Vioxx	SS	Merck & Co., Inc	ORAL
10 DAY		Renal Cyst	Vioxx	SS	Merck & Co., Inc	ORAL
146 DAY		Sepsis	Zestril	C		
UNKNOWN	2170 DAY	Sexual Dysfunction	Zestril	C		
UNKNOWN	2170 DAY	Weight Increased	Tenormin	C		
UNKNOWN			Tenormin	C		
UNKNOWN			Glucophage	C		
UNKNOWN			Micronase	C		
UNKNOWN			Lanoxin	C		
UNKNOWN	1051 DAY		[Therapy Unspecified]	C		
UNKNOWN			Aciphex	C		
UNKNOWN						

Date:06/28/05ISR Number: 4703071-8Report Type:Direct  
 Age:52 YR Gender:Female I/FU:I

Company Report #CTU 252109

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Asthenia		Wellbutrin	PS		ORAL
75MG X1 PO	1 DAY	Condition Aggravated		Wellbutrin	SS		ORAL
100 MG X1 PO	1 DAY						

Gout  
Headache  
Muscle Spasms  
Paraesthesia

Date:06/29/05ISR Number: 4702350-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0509984A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eye Pain		Ventolin	PS	Glaxosmithkline	
RESPIRATORY		Muscle Twitching					
(INHALATION)		Vision Blurred		Wellbutrin Xenical	SS C	Glaxosmithkline	

Date:06/29/05ISR Number: 4702623-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0102274A  
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Rash Pruritic					
per day	22 DAY						

Date:06/29/05ISR Number: 4702624-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0508929A  
Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Zyban	PS	Glaxosmithkline	
Other		Urticaria					
24 DAY							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/29/05ISR Number: 4702625-2Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0509455A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:06/29/05ISR Number: 4702626-4Report Type:Periodic  
Age:31 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0510661A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Insomnia		Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702627-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0510662A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702628-8Report Type:Periodic  
Age:27 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0511093A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain		Zyban	PS	Glaxosmithkline	ORAL
150TAB Twice		Dizziness					
per day	3 DAY			No Concurrent Medication	C		

Date:06/29/05ISR Number: 4702629-XReport Type:Periodic  
Age:30 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0511110A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	11 DAY	Anxiety Palpitations Urticaria		Zyban  Oral Contraceptives	PS  C	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702630-6Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0511623A  
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 98 DAY		Asthenia Musculoskeletal Stiffness Syncope Vertigo		Zyban Birth Control Pepcid	PS C C	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702631-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0512884A  
 Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation Insomnia		Zyban	PS	Glaxosmithkline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/29/05ISR Number: 4702632-XReport Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0513565A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Unknown	Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL
Pharmaceutical Product Complaint							

Date:06/29/05ISR Number: 4702633-1Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0515321A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Life-Threatening	Stevens-Johnson Syndrome		Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702634-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515529A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	2 DAY	Hypoaesthesia		Zyban	PS	Glaxosmithkline	

Date:06/29/05ISR Number: 4702635-5Report Type:Periodic  
 Age:54 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0515586A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose	150MG Per day 1 WK	Insomnia		Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702636-7Report Type:Periodic  
 Age: Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0515900A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Insomnia  
Zyban PS Glaxosmithkline ORAL  
150MG Twice  
per day 7 DAY  
No Concurrent Medication C

Date:06/29/05ISR Number: 4702637-9Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0516245A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	1 WK	Euphoric Mood					

Date:06/29/05ISR Number: 4702638-0Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0517786A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adverse Drug Reaction		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Dyspnoea					
per day		Pharyngolaryngeal Pain					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/29/05ISR Number: 4702639-2Report Type:Periodic  
 Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0518977A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702640-9Report Type:Periodic  
 Age:25 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520109A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Cold Sweat		Zyban	PS	Glaxosmithkline	ORAL
per day	10 DAY	Dizziness					
		Heart Rate Increased		Oral Contraceptive	C		
		Panic Attack					
		Paranoia					

Date:06/29/05ISR Number: 4702641-0Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0520184A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Unknown	WK	Pruritus		Zyban	PS	Glaxosmithkline	ORAL
				Hydrochlorothiazide	C		

Date:06/29/05ISR Number: 4702642-2Report Type:Periodic  
 Age:63 YR Gender:Female I/FU:I

Company Report #2065362

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Anger		Zyban	PS	Glaxosmithkline	ORAL
TOPICAL				Nicoderm Cq	C	Glaxosmithkline	

Date:06/29/05ISR Number: 4702643-4Report Type:Periodic  
Age:82 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0520974A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG Twice		Coordination Abnormal		Zyban	PS	Glaxosmithkline	ORAL
per day	10 DAY	Dizziness					
		Dysgeusia		Toprol	C		
UNKNOWN		Dysphonia		Lanoxin	C	Glaxosmithkline	
		Insomnia		Coumadin	C	Glaxosmithkline	

Date:06/29/05ISR Number: 4702644-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0521425A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
1 WK		Pruritus Generalised		Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702645-8Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0522144A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Tinnitus		Zyban	PS	Glaxosmithkline	ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/29/05ISR Number: 4702646-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523210A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nervousness		Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702647-1Report Type:Periodic  
 Age:33 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523453A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG Twice per day	8 DAY	Abnormal Dreams Drug Ineffective Feeling Jittery Myalgia		Zyban  No Concurrent Medication	PS  C	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702648-3Report Type:Periodic  
 Age:47 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0523538A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Local Swelling Obsessive Thoughts Pruritus		Zyban Nicoderm	PS C	Glaxosmithkline Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702649-5Report Type:Periodic  
 Age:20 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0523656A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 MON	Bacterial Infection Drug Interaction		Zyban	PS	Glaxosmithkline	ORAL
500MG Unknown	4 DAY	Flushing Tachycardia		Metronidazole	SS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702650-1Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524074A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
150MG	Unknown		Blood Pressure Increased	Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702651-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0524373A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
			Insomnia	Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702652-5Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525191A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
2	MON		Blood Pressure Increased	Zyban	PS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/29/05ISR Number: 4702653-7Report Type:Periodic  
Age:66 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525192A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression		Zyban	PS	Glaxosmithkline	ORAL
		Gastric Disorder		Unknown Medication	C		
		Heart Rate Increased		Unknown Medication	C		
		Pruritus					
		Rash					
		Urticaria					

Date:06/29/05ISR Number: 4702654-9Report Type:Periodic  
Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525197A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness		Zyban	PS	Glaxosmithkline	ORAL
		Tremor					

Date:06/29/05ISR Number: 4702655-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0525992A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 WK		Pruritus		Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702656-2Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526192A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MCG Per day	1 WK	Euphoric Mood		Zyban	PS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:06/29/05ISR Number: 4702657-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0526797A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Intolerance		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown				Wellbutrin Sr	C	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702658-6Report Type:Periodic  
Age:63 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0527947A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Herpes Zoster		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Rash					
per day	4 WK			Toprol Xl	C		
				Lisinopril	C		

Date:06/29/05ISR Number: 4702659-8Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0528396A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown	2 MON	Smoker		Wellbutrin	SS	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/29/05ISR Number: 4702660-4Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #IMP\_0823\_2004

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Zyban	PS	Glaxosmithkline	ORAL
				Bupropion			
				Hydrochloride	SS	Glaxosmithkline	ORAL

150MG Per day

Date:06/29/05ISR Number: 4702661-6Report Type:Periodic  
 Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0529992A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702662-8Report Type:Periodic  
 Age:65 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530010A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypertrophy Breast		Zyban	PS	Glaxosmithkline	ORAL
		Sexual Dysfunction		Avodart	SS	Glaxosmithkline	ORAL
				Atenolol	C		
				Lipitor	C		

150MG Per day

15 MON

Date:06/29/05ISR Number: 4702663-XReport Type:Periodic  
 Age:41 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0530372A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Zyban	PS	Glaxosmithkline	ORAL
		Insomnia		Levoxyl	C	Glaxosmithkline	

1TAB Per day 4 DAY

Date:06/29/05ISR Number: 4702664-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0531196A  
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Mouth Ulceration		Zyban	PS	Glaxosmithkline	ORAL
300MG Per day	12 DAY	Oedema Peripheral Urticaria					

Date:06/29/05ISR Number: 4702665-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0531250A  
Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Hypervigilance		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Tremor					
per day		Vision Blurred					

Date:06/29/05ISR Number: 4702666-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0531550A  
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Abnormal Faeces		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Confusional State					
per day	1 MON	Irritability		Ambien	C		

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Freedom Of Information (FOI) Report

Date:06/29/05ISR Number: 4702668-9Report Type:Periodic  
 Age:76 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0532370A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL
150MG	Unknown						

Date:06/29/05ISR Number: 4702669-0Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0532920A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702670-7Report Type:Periodic  
 Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533053A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Balance Disorder		Zyban	PS	Glaxosmithkline	ORAL
150MG	Twice						
per day	1	Constipation					
		Dizziness					

Date:06/29/05ISR Number: 4702671-9Report Type:Periodic  
 Age:55 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0533264A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia		Zyban	PS	Glaxosmithkline	ORAL
150MG	Twice						
per day	13						
				Prempro	C		

Date:06/29/05ISR Number: 4702672-0Report Type:Periodic  
 Age:32 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0535043A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Chills Insomnia Night Sweats Nightmare		Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702673-2Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535588A  
 Age:48 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1 MON		Convulsion		Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702674-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0535633A  
 Age:31 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2TAB Twice per day	2 WK	Hypersensitivity Pruritus		Zyban	PS	Glaxosmithkline	ORAL



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Freedom Of Information (FOI) Report

Date:06/29/05ISR Number: 4702675-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0535782A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
		Balance Disorder Difficulty In Walking Mobility Decreased					

Date:06/29/05ISR Number: 4702676-8Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538231A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
150MG Per day		Neck Pain		Amoxicillin	C	Glaxosmithkline	

Date:06/29/05ISR Number: 4702677-XReport Type:Periodic  
Age:32 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0539708A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	ORAL
150MG Twice per day		Urticaria					

Date:06/29/05ISR Number: 4702678-1Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540277A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Zyban	PS	Glaxosmithkline	
UNKNOWN	2 WK	Dysgeusia					

Date:06/29/05ISR Number: 4702679-3Report Type:Periodic  
Age:25 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0542846A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Oedema					
per day	28	DAY		Nicorette Gum	C	Glaxosmithkline	
		Rash		Oral Contraceptive	C		
		Rash Maculo-Papular					

Date:06/29/05ISR Number: 4702680-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0543061A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day	3	WK		Nicorette Gum	C	Glaxosmithkline	

Date:06/29/05ISR Number: 4702681-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0543948A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Palpitations		Zyban	PS	Glaxosmithkline	ORAL
2	DAY			No Concurrent Medication	C		

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Freedom Of Information (FOI) Report

Date:06/29/05ISR Number: 4702682-3Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544038A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Zyban	PS	Glaxosmithkline	ORAL
		Drug Interaction		Xenical	SS		

Date:06/29/05ISR Number: 4702683-5Report Type:Periodic  
 Age:45 YR Gender:Female I/FU:I

Company Report #2005003216

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Zyban	PS	Glaxosmithkline	ORAL
TOPICAL	21MG Per day	Fatigue		Nicoderm	SS	Glaxosmithkline	
		Insomnia					
		Psychomotor Hyperactivity					

Date:06/29/05ISR Number: 4702684-7Report Type:Periodic  
 Age:60 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0544543A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	20 DAY	Dizziness		Luvox	C		
		Heart Rate Increased					
		Nausea					
		Palpitations					
		Rash					
		Tinnitus					

Date:06/29/05ISR Number: 4702685-9Report Type:Periodic  
 Age:50 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545155A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							

per day 1 WK

Advair	C	Glaxosmithkline
Spiriva	C	
Nicoderm	C	Glaxosmithkline
Reglan	C	Glaxosmithkline
Nexium	C	
Singulair	C	

Date:06/29/05ISR Number: 4702686-0Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0546355A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased		Zyban	PS	Glaxosmithkline	ORAL
1	MON						

Date:06/29/05ISR Number: 4702687-2Report Type:Periodic  
Age:41 YR Gender:Male I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0548191A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization - 150MG Twice							
Initial or Prolonged per day	6 DAY	Asthenia					
Other		Rash Serum Sickness		Prednisone	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/29/05ISR Number: 4702688-4Report Type:Periodic  
Age:55 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549404A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal		Zyban	PS	Glaxosmithkline	ORAL
150MG	Unknown			Blood Pressure Medication	C		
				Unknown Medication	C		

Date:06/29/05ISR Number: 4702689-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0549904A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia		Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702690-2Report Type:Periodic  
Age:36 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550651A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nervousness		Zyban	PS	Glaxosmithkline	ORAL
150MG	Per day 6 WK			No Concurrent Medication	C		

Date:06/29/05ISR Number: 4702691-4Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0551345A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria		Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702692-6Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0555386A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
per day							

Date:06/29/05ISR Number: 4702693-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0555428A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity Therapeutic Response Unexpected		Zyban	PS	Glaxosmithkline	ORAL

Date:06/29/05ISR Number: 4702694-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0555458A  
 Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Pruritus					
per day							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/29/05ISR Number: 4703108-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0558978A  
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 40 DAY		Asthenia		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged Other		Dysarthria Facial Dysmorphism Memory Impairment					

Date:06/30/05ISR Number: 4704378-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0556917A  
 Age:18 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 900MG Single Initial or Prolonged dose	2 HR	Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
		Fatigue Heart Rate Increased Hypoaesthesia Overdose Posture Abnormal		No Concurrent Medication	C		

Date:06/30/05ISR Number: 4704379-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0557281A  
 Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 100MG Per day 1 DAY		Aspiration		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization - Initial or Prolonged Other		Depressed Level Of Consciousness Grand Mal Convulsion Incontinence Pulse Absent Staring Vomiting		Seroquel	C		

Date:06/30/05ISR Number: 4704381-0Report Type:Expedited (15-DaCompany Report #13148  
Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tachycardia		Wellbutrin	PS	Glaxosmithkline	
150MG Per day				Unspecified Ssri	C		

Date:06/30/05ISR Number: 4704383-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0563942A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death		Wellbutrin	PS	Glaxosmithkline	
RESPIRATORY Life-Threatening (INHALATION)		Drug Abuser					
Other		Incorrect Route Of Drug Administration		No Concurrent Medication	C		

Date:06/30/05ISR Number: 4704384-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0563979A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Paxil	PS	Glaxosmithkline	
		Pituitary Tumour		Wellbutrin	SS	Glaxosmithkline	ORAL



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Freedom Of Information (FOI) Report

Date:06/30/05ISR Number: 4704386-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0564272A  
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG Per day	Asthma		Wellbutrin	PS	Glaxosmithkline	ORAL
	3 MON			Synthroid	C	Glaxosmithkline	
				Lithium	C	Glaxosmithkline	

Date:06/30/05ISR Number: 4704387-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0564273A  
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	6 WK	Grand Mal Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Hypoaesthesia		Amitriptyline	C		
		Staring		Flexeril	C		
		Unresponsive To Verbal		Vasotec	C		
		Stimuli		Carbidopa	C		
				Darvocet	C		
				Enalapril	C		
				Verapamil	C		
				Clavix	C		
				Cyclobenzaprine	C		
				Meclizine	C		

Date:06/30/05ISR Number: 4704390-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0564635A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS	Glaxosmithkline	

Date:06/30/05ISR Number: 4704431-1Report Type:Direct Company Report #CTU 252264  
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other  
150 MG PO BID

Condition Aggravated

Wellbutrin Sr PS ORAL

Depression  
Therapeutic Response  
Unexpected With Drug  
Substitution

Date:07/01/05ISR Number: 4705378-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0557786A  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 241 DAY		Anger		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Grand Mal Convulsion Intentional Self-Injury Memory Impairment		Alcohol	SS		ORAL

Date:07/01/05ISR Number: 4705382-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0560141A  
Age:42 YR Gender:Male I/FU:F

Outcome  
Life-Threatening  
Hospitalization -  
Initial or Prolonged  
Disability

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Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
7 DAY		Accidental Overdose		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
10MG Four times per day		Amnesia		Toradol	C		ORAL
		Blood Pressure Increased					
		Confusional State		Toprol	C		
		Drug Ineffective		Ultram	C		
		Medication Error		Biaxin Xl	C		
				Lortab	C		

Date:07/01/05ISR Number: 4705388-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0564276A  
Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day 4 WK Initial or Prolonged		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
		Feeling Abnormal		No Concurrent Medication	C		
		Muscle Tightness					

Date:07/01/05ISR Number: 4705392-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0564494A  
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day		Glaucoma		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice per day				Wellbutrin Sr	SS	Glaxosmithkline	ORAL
				Sam-E	C		
				Melatonin	C		
				Hctz	C		
				Lovastatin	C		

Date:07/01/05ISR Number: 4705396-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0564755A  
Age:86 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 200MG Twice	Confusional State		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day 6 MON	Dehydration					
Disability	Dementia		Digoxin	C	Glaxosmithkline	
	Hallucination		Norvasc	C		
			Glipizide	C		
			Avandia	C	Glaxosmithkline	
			Prandin	C		
			Toprol Xl	C		
			Lipitor	C		
			Celexa	C		
			Megace	C		
			Reminyl	C		

Date:07/05/05ISR Number: 4706451-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0556337A  
Age:32 YR Gender:Female I/FU:F

Outcome	PT
Other	Dizziness
	Heart Rate Increased
	Hyperhidrosis
	Night Sweats

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
25MG Per day	7 MON	Paraesthesia Psoriasis		Paxil Cr	PS	Glaxosmithkline	ORAL
10 DAY				Wellbutrin	SS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:07/05/05ISR Number: 4706459-4Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0564879A  
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Lenticular Opacities		Paxil	PS	Glaxosmithkline	ORAL
Other		Myopia		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
				Bromazepam	C		
				Losec	C	Glaxosmithkline	
				Starnoc	C		
				Zopiclone	C		

Date:07/05/05ISR Number: 4706474-0Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0381636A  
Age:21 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Arthralgia		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Blister					
Other	18 DAY	Chest Pain		Marijuana	C		
per day		Constipation		Cannabis	C		
		Dermatitis Allergic					
		Disorientation					
		Drug Administration Error					
		Eye Swelling					
		Fear					
		Joint Swelling					
		Loss Of Consciousness					
		Mood Altered					
		Oral Pain					

Palpitations  
Panic Reaction  
Pharmaceutical Product  
Complaint  
Pharyngitis Streptococcal  
Rash  
Skin Exfoliation  
Stevens-Johnson Syndrome  
Urticaria

Date:07/05/05ISR Number: 4706504-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0386598A  
Age: Gender:Female I/FU:F

Outcome PT  
Other Dizziness  
Eyelid Ptosis  
Hypoaesthesia  
Insomnia  
Malaise  
Nightmare  
Pain In Extremity  
Skin Discolouration

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Thrombosis

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
UNKNOWN			Zyban	PS	Glaxosmithkline	

Date:07/05/05ISR Number: 4706505-8Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0386645A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN				Zyban	PS	Glaxosmithkline	
Initial or Prolonged		Bedridden Fatigue Stress Urticaria		No Concurrent Medication	C		

Date:07/05/05ISR Number: 4706799-9Report Type:Expedited (15-DaCompany Report #2005-DE-02334GD  
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Abuser Drug Toxicity	Literature	Paracetamol (Paracetamol)	PS		
				Oxazepam (Oxazepam)	SS		
				Diazepam (Diazepam)	SS		
				Doxylamine (Doxylamine Succinate)	SS		
				Morphine (Morphine)	SS		
				Oxycodone (Oxycodone)	SS		
				Salicylates (Salicylates)	SS		
				Promethazine (Promethazine)	SS		
				Heroin (Diamorphine)	SS		
				Bupropion (Bupropion)	SS		
				Carbamazepine (Carbamazepine)	SS		

Metoprolol  
(Metoprolol)

SS

Date:07/05/05ISR Number: 4708147-7Report Type:Direct  
Age: Gender: I/FU:I

Company Report #USP 57224

Outcome Dose	Duration	PT Drug Dispensing Error Medication Error	Report Source	Product	Role	Manufacturer	Route
TABLET , EXTENDED RELEASE				Burpropion Extended Release	PS	Eon Labs	
TABLET, EXTENEDED RELEASE				Wellbutrin Xl	SS	Glaxosmithkline	



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/05/05ISR Number: 4709237-5Report Type:Expedited (15-DaCompany Report #2005090868

Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Urticaria Generalised	Consumer	Rogaine Fragrance For Women - Spring Bloom (Minoxidil)	PS		
TOPICAL	1 ML, BID,						
TOPICAL				Bupropion Hydrochloride (Bupropoi Hydrochloride)	SS		
	150 MG ONCE,						
DAILY							

Date:07/06/05ISR Number: 4707459-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0564736A

Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG Per day	2 WK	Medication Error		Marijuana	C		

Date:07/06/05ISR Number: 4708527-XReport Type:Expedited (15-DaCompany Report #S05-USA-03304-01

Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Dizziness Epilepsy Fall Fatigue Head Injury Memory Impairment Postictal Headache	Consumer	Citalopram Wellbutrin (Bupropion Hydrochloride)	PS  SS		

Date:07/07/05ISR Number: 4708512-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0546080A

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Wellbutrin	PS	Glaxosmithkline	ORAL
Life-Threatening		Medication Error		Wellbutrin	SS	Glaxosmithkline	ORAL
Other				Bupropion	SS	Glaxosmithkline	
				No Concurrent Medication	C		

Date:07/07/05ISR Number: 4708543-8Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0386598A

Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dizziness		Zyban	PS	Glaxosmithkline	ORAL
8 DAY		Eyelid Ptosis					
		Hypoaesthesia					
		Insomnia					
		Lip Disorder					
		Malaise					
		Nightmare					
		Pain In Extremity					
		Sensation Of Heaviness					
		Skin Discolouration					
		Thrombosis					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/07/05ISR Number: 4709065-0Report Type:Expedited (15-DaCompany Report #IMP\_0847\_2005

Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1 TAB QDAY PO	Anxiety	Health	Bupropion	PS		ORAL
		Condition Aggravated	Professional	Lexapro	C		
		Crying		Lorazepam	C		
		Disease Recurrence		Calcium Citrate	C		
		Feeling Abnormal		Glucosamine			
		Major Depression		Chondrotin	C		
		Mental Disorder		Multivitamin	C		
		Paranoia					
		Pharmaceutical Product Complaint					
		Post-Traumatic Stress Disorder					
		Suicidal Ideation					

Date:07/08/05ISR Number: 4709442-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0563942A

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Wellbutrin	PS	Glaxosmithkline	
RESPIRATORY							
Life-Threatening		Drug Abuser					
(INHALATION)							
Other				No Concurrent Medication	C		

Date:07/08/05ISR Number: 4709443-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0564491A

Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Life-Threatening		Overdose		Cocaine	SS		
Hospitalization -				Advil	SS	Glaxosmithkline	
Initial or Prolonged							
Other							

Date:07/08/05ISR Number: 4709446-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0565250A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropion	PS	Glaxosmithkline	
Other		Hypertension Renal Failure		Lithium	SS	Glaxosmithkline	

Date:07/08/05ISR Number: 4709459-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0386419A  
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 36 DAY		Angina Pectoris		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Bradycardia Chest Pain Chest Wall Pain Cholelithiasis Constipation Hepatic Pain					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/08/05ISR Number: 4711041-9Report Type:Direct  
Age:17 YR Gender:Male I/FU:I

Company Report #CTU 252844

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Abnormal Behaviour		Wellbutrin 150 Mg			
Hospitalization -		Mood Swings		Brand	PS		
Initial or Prolonged		Suicidal Ideation		Wellbutrin 100 Mg			
Other				Generic	SS		
				Depakote	C		
				Tenex	C		
				Adderal	C		

Date:07/08/05ISR Number: 4711322-9Report Type:Expedited (15-DaCompany Report #2001058760US  
Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Headache	Health	Solu-Medrol			
Initial or Prolonged		Pruritus	Professional	(Methylprednisolone			
Other		Rash		Sodium Succinate)	PS		
INTRAVENOUS	500 MG	(125					
MG, 4 IN 1		Urticaria					

D),

INTRAVENOUS;

SEE IMAGE

				Hydroxyzine			
				Hydrochloride			
				(Tablet)			
				(Hydroxyzine			
ORAL				Hydrochloride)	SS		ORAL
				Prednisone Tablet			
				(Prednisone)	SS		ORAL
				Wellbutrin			
				(Bupropion			
				Hydrochloride)	SS		ORAL

300 MG (150

MG, 2 IN 1

D), ORAL

Ortho Tri-Cyclen  
(Ethinylestradiol,  
Norgestimate) C  
Propranolol  
(Propranolol) C  
Imitrex (Sumatriptan  
Succinate) C

Date:07/11/05ISR Number: 4710829-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0551834A  
Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Convulsion Depressed Level Of Consciousness Dizziness Drug Toxicity Fall Headache Loss Of Consciousness Malaise Respiratory Arrest Tachycardia		Wellbutrin	PS	Glaxosmithkline	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/11/05ISR Number: 4710831-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0559557A

Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypothyroidism		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	8 MON						
20MG Per day		Myopathy		Lexapro	C		ORAL

Date:07/11/05ISR Number: 4710837-7Report Type:Expedited (15-DaCompany Report #S05-USA-02982-01

Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Dizziness		Citalopram	SS		ORAL
		Dyskinesia					
		Epilepsy					
		Fall					
		Fatigue					
		Head Injury					
		Postictal Headache					

Date:07/11/05ISR Number: 4710841-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0565030A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anger		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
MON							
		Condition Aggravated					
		Depressed Mood					
		Feeling Abnormal					
		Irritability					
		Psychiatric Symptom					
		Suicidal Ideation					

Date:07/11/05ISR Number: 4710871-7Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0384296A

Age:70 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Burning Sensation		Zyban	PS	Glaxosmithkline	
UNKNOWN		300MG	Per day 13 DAY		Celebrex	C		
200U Per day			Chills		Norvasc	C		
			Difficulty In Walking					
			Erythema					
			Feeling Cold					
			Hyperaesthesia					
			Pain					
			Pain In Extremity					
			Psoriasis					
			Rash					
			Rash Generalised					
			Skin Exfoliation					
			Skin Laceration					

Date:07/11/05ISR Number: 4710874-2Report Type:Expedited (15-DaCompany Report #SE-GLAXOSMITHKLINE-B0386513A  
Age:58 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Aggression		Zyban	PS	Glaxosmithkline	ORAL
1 DAY			Anger		Valeriana	C	Glaxosmithkline	
Initial or Prolonged			Anxiety		Antidepressant	C		
			Drug Interaction		Alcohol	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/11/05ISR Number: 4710875-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0386556A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Asthma		Zyban	PS	Glaxosmithkline	ORAL
				Atenolol	C		
				Beclomethasone	C	Glaxosmithkline	

Date:07/11/05ISR Number: 4710887-0Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0387151A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 10 DAY		Aggression		Zyban	PS	Glaxosmithkline	ORAL
		Agitation					
		Anger					
		Crying					
		Dry Mouth					
		Emotional Disorder					
		Insomnia					
		Irritability					
		Stress					

Date:07/11/05ISR Number: 4710888-2Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0046030A

Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 150MG Per day		Cardiac Failure		Zyban	PS	Glaxosmithkline	ORAL
Hospitalization - 10MG Per day		Dyspnoea		Simvastatin	C		ORAL
Initial or Prolonged UNKNOWN		Flank Pain		Metoprolol	C		
UNKNOWN		Oxygen Saturation		Delix	C		
UNKNOWN		Decreased		Aldactone	C		
UNKNOWN		Supraventricular		Sortis	C		

UNKNOWN	Tachycardia	Torem	C	
UNKNOWN		Ass	C	Glaxosmithkline
UNKNOWN		Omeprazol	C	Glaxosmithkline

Date:07/11/05ISR Number: 4711504-6Report Type:Direct Company Report #CTU 252956  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 40 MG QD		Anger		Paxil 40 Mg.	PS		ORAL
ORAL		Emotional Disorder					
450 MG QD		Hostility Muscle Twitching		Wellbutrin Xl 450 Mg.	SS		ORAL
ORAL		Nervousness		Prozac	C		

Date:07/11/05ISR Number: 4712021-XReport Type:Direct Company Report #CTU 252969  
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1 TABLET		Formication Therapeutic Response		Bupropion 150 Mg Sr Ta Watson	PS	Watson	ORAL
BID ORAL		Unexpected With Drug Substitution					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/12/05ISR Number: 4712239-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0565550A  
Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	18 MON	Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL
		Overdose					

Date:07/12/05ISR Number: 4712240-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0565551A  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	150MG Per day	Rash		Wellbutrin	PS	Glaxosmithkline	ORAL
		Skin Exfoliation		Unknown	C		
		Stevens-Johnson Syndrome					
		White Blood Cell Count Increased					

Date:07/12/05ISR Number: 4712251-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0380373A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1TAB Twice per day	Anxiety		Zyban	PS	Glaxosmithkline	ORAL
	4 DAY	Feeling Abnormal					
		Panic Attack		No Concurrent Medication	C		
		Serotonin Syndrome					

Date:07/12/05ISR Number: 4712268-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-B0387160A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Condition Aggravated		Zyban	PS	Glaxosmithkline	ORAL
		Depressed Mood					
		Disability					

Hostility  
Overdose  
Suicidal Ideation

Date:07/13/05ISR Number: 4713304-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0565250A  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Lithium	PS	Glaxosmithkline	
Other		Drug Exposure During Pregnancy Hypertension Pregnancy Renal Failure Renal Function Test Abnormal Renal Impairment		Bupropion	SS	Glaxosmithkline	

Date:07/13/05ISR Number: 4714340-XReport Type:Direct Company Report #CTU 253163  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Bupropion 150 (Generic)	PS		
BID		Anxiety Fatigue					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/14/05ISR Number: 4714382-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0556917A  
 Age:18 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 900MG Single Initial or Prolonged dose	Anxiety		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
15 DAY	Convulsion					
	Discomfort		Lexapro	SS		ORAL
	Fatigue Heart Rate Increased Hypoaesthesia Intentional Misuse Muscle Contracture Paraesthesia Posture Abnormal Suicide Attempt Tremor		No Concurrent Medication	C		

Date:07/14/05ISR Number: 4714383-6Report Type:Expedited (15-DaCompany Report #S05-USA-02393-01  
 Age:18 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 20MG Per day 15 DAY	Convulsion		Wellbutrin Xl	PS	Glaxosmithkline	ORAL
	Discomfort		Lexapro	SS		ORAL
	Fatigue Hypoaesthesia Ill-Defined Disorder Intentional Misuse Muscle Contractions Involuntary Paraesthesia Tremor					

Date:07/14/05ISR Number: 4714384-8Report Type:Expedited (15-DaCompany Report #2005-01-1690  
 Age:55 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Death	Haemorrhagic Stroke	Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Three					
Hospitalization -					
times per day 413 DAY					
Initial or Prolonged		Pegylated Interferon			
		Alpha 2a	SS		
SUBCUTANEOUS 180MCG Weekly 396 DAY		Copegus	SS		ORAL
1200MG Per					
day 413 DAY					
		Pegylated Interferon			
SUBCUTANEOUS 150MCG Weekly		Alpha 2b	SS		
		Percocet	C		
		Benzodiazepine	C		
		Levoxyl	C	Glaxosmithkline	ORAL
.175MG Per					
day					
1MG Three		Lorazepam	C		ORAL
times per day					
40MG Per day		Protonix	C		ORAL
325MG Twice		Percodan	C		ORAL
per day 101 DAY					
SUBCUTANEOUS 40000U Weekly 277 DAY		Procrit	C		
5MG Twice per		Diazepam	C		ORAL
day 413 DAY					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/14/05ISR Number: 4714388-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0566115A  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Vitreous Detachment		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Twice							
per day				No Concurrent Medication	C		

Date:07/14/05ISR Number: 4714389-7Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0338473A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Consciousness		Zyban	PS	Glaxosmithkline	
				Ethanol	SS		

Date:07/14/05ISR Number: 4714390-3Report Type:Expedited (15-DaCompany Report #NO-GLAXOSMITHKLINE-B0349044A  
 Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Insomnia		Zyban	PS	Glaxosmithkline	ORAL
		Sleep Disorder					
		Visual Disturbance					

Date:07/14/05ISR Number: 4714510-0Report Type:Expedited (15-DaCompany Report #200511975US  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Exposure During Pregnancy		Allegra	PS	Aventis Pharmaceuticals Inc.	ORAL
		Haemorrhage		Bupropion	SS		
		Pregnancy		Celexa	SS		ORAL
		Umbilical Cord Abnormality		Zantac	SS		ORAL

Date:07/15/05ISR Number: 4715141-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0566214A  
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthma		Wellbutrin XL	PS	Glaxosmithkline	
150MG Per day	3	WK	Chest Discomfort	Singulair	C		
				Zyrtec	C	Glaxosmithkline	
				Advair	C	Glaxosmithkline	
				Beconase	C	Glaxosmithkline	NASAL

Date:07/15/05ISR Number: 4715153-5Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0386645A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anxiety		Zyban	PS	Glaxosmithkline	
UNKNOWN							
Initial or Prolonged		Bedridden		No Concurrent			
		Fatigue		Medication	C		
		Stress					
		Urticaria					



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/15/05ISR Number: 4715162-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0387289A  
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	150MG per day 3 DAY	Sudden Death		Zyban	PS	Glaxosmithkline	ORAL
				Topalgic (France)	C	Other	ORAL
				Effergal	C	Other	ORAL
				Percutalgine	C	Other	
				Brexin	C	Other	ORAL

Date:07/15/05ISR Number: 4717312-4Report Type:Expedited (15-DaCompany Report #2005080150  
 Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dizziness Feeling Abnormal Flat Affect Multiple Sclerosis Nasopharyngitis Pain Personality Change Retching Visual Disturbance Vomiting	Consumer	Celebrex (Celecoxib)	PS		
				Paxil (Paroxetine Hydrochloride)	SS		
				Wellbutrin (Bupropion Hydrochloride)	SS		
				Aleve (Naproxen Sodium)	SS		
				Avalide (Hydrochlorothiazide , Irbesartan)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Multivitaminis (Multivitaminis)	C		
				Atenolol (Atenolol)	C		

Date:07/18/05ISR Number: 4716003-3Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0385105A  
 Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization -	Chest Pain	Zyban	PS	Glaxosmithkline	ORAL
1TAB Per day 5 DAY					
Initial or Prolonged	Dizziness	Valerian	C	Glaxosmithkline	ORAL
3U per day 11 DAY					
	Heart Alternation				
	Hypertension				
	Palpitations				

Date:07/18/05ISR Number: 4716019-7Report Type:Expedited (15-DaCompany Report #EE-GLAXOSMITHKLINE-B0387910A  
 Age:36 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day 39 DAY						
	Loss Of Consciousness		Seroquel	C		ORAL
.2G Per day						
	Screaming					

Date:07/18/05ISR Number: 4717865-6Report Type:Direct Company Report #CTU 253484  
 Age:51 YR Gender:Female I/FU:I

Outcome  
 Life-Threatening  
 Hospitalization -  
 Initial or Prolonged  
 Required

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Intervention to Prevent Permanent Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Grand Mal Convulsion		Wellbutrin Initial Dosage Glaxosmithkline	PS	Glaxosmithkline	ORAL
ORAL				Ms	C		

Date:07/18/05ISR Number: 4718062-0Report Type:Direct  
Age: Gender: I/FU:I Company Report #CTU 253475

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 20 ML ONCE A Hospitalization - DAY ORAL		Bipolar Disorder		Zoloft	PS		ORAL
Initial or Prolonged 350/400 ML Required ONCE A DAY		Depression		Wellbutrin	SS		ORAL
Intervention to Prevent Permanent Impairment/Damage		Disturbance In Attention Headache Intentional Self-Injury Suicide Attempt Tremor Vision Blurred Weight Decreased		Triliptal Lamictal Topomax Geoddon Prozac	C C C C C		

Date:07/18/05ISR Number: 4719224-9Report Type:Direct  
Age:41 YR Gender:Male I/FU:I Company Report #CTU 253453

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG BID ORAL		Depression Pharmaceutical Product Complaint		Bupropion 150mg Watson Pharmaceuticals	PS	Watson Pharmaceuticals	ORAL

Lipitor C  
 Avandia C  
 Lisinopril C  
 Depakote C

Date:07/19/05ISR Number: 4717130-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0551252A  
 Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Vision Blurred		Paxil	PS	Glaxosmithkline	
		Visual Acuity Reduced		Wellbutrin	SS	Glaxosmithkline	ORAL
				Trazodone	C		ORAL
5 YR				Risperdal	C		ORAL
2 YR				Cogentin	C		ORAL
2MG Twice per							
day	2 YR			Unknown Medication	C		
				Vitamin B12	C	Glaxosmithkline	
				Multivitamin	C		
				Vitamin C	C	Glaxosmithkline	
				Magnesium	C		
				Iron	C		
				Calcium	C		
				Vitamin E	C		
				Colace	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/19/05ISR Number: 4717132-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0560956A  
 Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	300MG Per day	3 YR	Confusional State	Wellbutrin	PS	Glaxosmithkline	ORAL
Other			Dehydration	Alcohol	SS		
			Fall	Lexapro	C		ORAL
	20MG Per day		Grand Mal Convulsion	Yasmin	C		ORAL
			Head Injury	Multivitamin	C		
			Hypertension				
			Hypoxia				
			Poisoning				
			Pupil Fixed				
			Vomiting				

Date:07/19/05ISR Number: 4717133-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0561914A  
 Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	100MG Per day	1 WK	Muscle Spasms	Wellbutrin Sr	PS	Glaxosmithkline	ORAL
Other	150MG Per day	24 DAY	Muscle Twitching	Wellbutrin Xl	SS	Glaxosmithkline	ORAL
				Ambien	C		
				Lexapro	C		

Date:07/19/05ISR Number: 4717138-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0566516A  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Drug Abuser	Wellbutrin	PS	Glaxosmithkline	
				Prozac	SS		
				Alcohol	SS		

Date:07/19/05ISR Number: 4717146-0Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0386645A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN		Anxiety		Zyban	PS	Glaxosmithkline	
Initial or Prolonged		Bedridden Fatigue Stress Urticaria		No Concurrent Medication	C		

Date:07/19/05ISR Number: 4717159-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0387710A  
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1TAB Twice		Chest Pain		Zyban	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day	38 DAY	Cholelithiasis Constipation		Trimethoprim	C	Glaxosmithkline	ORAL

Date:07/19/05ISR Number: 4717162-9Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0387737A  
Age:23 YR Gender:Female I/FU:F

Outcome	PT
Other	Depression Dry Mouth Emotional Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Emotional Distress Hallucination, Auditory Headache				
		Inappropriate Affect Insomnia Lethargy Night Sweats Nightmare	Zyban	PS	Glaxosmithkline	ORAL

Date:07/19/05ISR Number: 4717165-4Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0388012A  
Age: Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
Hospitalization - Initial or Prolonged		Asthma Dyspnoea Pneumonia	Zyban Prednisolone	PS C	Glaxosmithkline Glaxosmithkline	ORAL

Date:07/20/05ISR Number: 4718609-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0566773A  
Age: Gender:Male I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
Death		Death	Wellbutrin	PS	Glaxosmithkline	

Date:07/20/05ISR Number: 4718642-2Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0388098A  
Age:45 YR Gender:Male I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
Other UNKNOWN		Feeling Abnormal 3 DAY Hallucination Heart Rate Increased Nightmare Panic Attack Somnolence	Zyban	PS	Glaxosmithkline	

Date:07/20/05ISR Number: 4723534-9Report Type:Direct  
Age:53 YR Gender:Male I/FU:I

Company Report #CTU 253756

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anaphylactic Shock Obstructive Airways Disorder Swelling		Lexapro Wellbutrin	PS SS		

Date:07/20/05ISR Number: 4723710-5Report Type:Direct  
Age:74 YR Gender:Male I/FU:I

Company Report #CTU 253789

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 30 MG PO BID Initial or Prolonged 150 MG Q AM 2 TSP Q4HR PRN		Abnormal Behaviour Aggression Hallucinations, Mixed Psychotic Disorder Serotonin Syndrome		Paroxetine Bupropion Guaifenesin Dextromethorphan	PS SS SS SS		ORAL



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/20/05ISR Number: 4724272-9Report Type:Direct  
Age: Gender:Male I/FU:I

Company Report #CTU 253815

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated		Wellbutrin Sr	PS		ORAL
150 MG PO BID		Depression		Lipitor	C		
		Drug Effect Decreased		Ultram	C		
		Fatigue		Lorazepam	C		
		Social Avoidant Behaviour					
		Therapeutic Response					
		Unexpected With Drug					
		Substitution					

Date:07/21/05ISR Number: 4720718-0Report Type:Expedited (15-DaCompany Report #CA-GLAXOSMITHKLINE-A0566496A  
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death		Wellbutrin Sr	PS	Glaxosmithkline	ORAL
				Ethanol	C		ORAL

Date:07/21/05ISR Number: 4720719-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0566707A  
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other		Drug Abuser		Seroquel	C		

Date:07/21/05ISR Number: 4720720-9Report Type:Expedited (15-DaCompany Report #13031  
Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous		Wellbutrin	PS	Glaxosmithkline	
300MG per day				Unspecified Ssri	C		
				Mood Medication	C		
				Tricyclic			

Date:07/21/05ISR Number: 4721458-4Report Type:Periodic  
Age:63 YR Gender:Female I/FU:I

Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-12708301

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Coumadin Tabs	PS	Bristol-Myers Squibb Company	ORAL
Start 5mg qd; present daily dosage was 10mg, 10mg, 5mg,		Drug Interaction International Normalised Ratio Decreased					
				Zyprexa	C		
				Effexor	C		
				Multivitamin	C		
				Plavix	C	Regulatory Health Authority South Africa	
Previously on Rx				Klonopin	C		
				Wellbutrin	I		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/22/05ISR Number: 4722832-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0567026A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ventricular Tachycardia		Wellbutrin Unknown	PS C	Glaxosmithkline	ORAL

Date:07/22/05ISR Number: 4722837-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0567064A

Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other	150MG Per day 2 DAY	Blister Haemorrhage Local Swelling		Wellbutrin Xl  Allergen No Concurrent Medication	PS  SS C	Glaxosmithkline	ORAL

Date:07/22/05ISR Number: 4725665-6Report Type:Direct

Company Report #CTU 254066

Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other	200 MG TWICE A DAY ORAL	Amnesia Therapeutic Response Unexpected With Drug  Substitution		Bupropion Hcl Extended Release Tabs Global  Wellbutrin	PS  C	Global	ORAL

Date:07/25/05ISR Number: 4723460-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0566698A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -	20MG Per day 5 YR	Anaemia		Paxil	PS	Glaxosmithkline	ORAL
Initial or Prolonged	52 WK	Drug Exposure During		Wellbutrin	SS	Glaxosmithkline	ORAL

250MG Single	Pregnancy	Ibuprofen	SS	Glaxosmithkline	ORAL
dose	Initial Insomnia				
	Overdose	Iron	C		
	Somnolence				

Date:07/25/05ISR Number: 4723468-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0382344A  
 Age:31 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Epilepsy		Zyntabac	PS	Glaxosmithkline	
UNKNOWN	300MG per day					
Initial or Prolonged	Facial Bones Fracture					
Other	Fibula Fracture					
	Ill-Defined Disorder					

Date:07/25/05ISR Number: 4723481-2Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0388188A  
 Age:42 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Life-Threatening	Panic Attack		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	7 DAY					
TOPICAL	Suicidal Ideation		Estraderm	C		
	100MCG per					
day						
50MG Three			Diclofenac	C		ORAL
times per day						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/25/05ISR Number: 4725754-6Report Type:Expedited (15-DaCompany Report #A0499051A  
Age:6 MON Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Exposure Via Breast Milk	Literature Health Professional	Wellbutrin Sr Tablet-Controlled Release (Bupropion Hydrochloride)	PS		
TRANSMAMMARY	150 MG/	PER					
DAY/		Restlessness					
TRANSMAMMARY		Staring					
		Upper Respiratory Tract Infection					

Date:07/25/05ISR Number: 4726035-7Report Type:Expedited (15-DaCompany Report #D0039102A  
Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Acute Psychosis Disturbance In Attention Insomnia	Foreign Literature Health	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		ORAL
ORAL	5 DAY	Schizophreniform Disorder Suicidal Ideation Vision Blurred	Professional				

Date:07/26/05ISR Number: 4724742-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0567176A  
Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Grand Mal Convulsion Overdose		Wellbutrin Marijuana	PS C	Glaxosmithkline	ORAL
Other							

Date:07/26/05ISR Number: 4724743-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0567177A  
Age:19 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Blood Creatine		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -		Phosphokinase Increased		Ethylene Glycol	SS		
Initial or Prolonged		Convulsion		Marijuana	C		
Other		Drug Screen Positive					
		Dyskinesia					
		Loss Of Consciousness					
		Metabolic Acidosis					
		Overdose					
		Renal Failure Acute					
		Vomiting					

Date:07/26/05ISR Number: 4724745-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0567196A  
Age:77 YR Gender:Female I/FU:I

Outcome	PT
Other	Abdominal Pain Upper
	Anger
	Arthralgia
	Chest Pain
	Drug Ineffective
	Ear Haemorrhage
	Insomnia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150MG Per day	2 WK	Irritability Mouth Ulceration Myalgia Rash Generalised Somnolence		Wellbutrin	PS	Glaxosmithkline	ORAL
				Lipitor	C		
				Aspirin	C	Glaxosmithkline	
				Vitamins	C		
				Glucosamine	C		
				Calcium	C		
				Aleve	C		

Date:07/26/05ISR Number: 4724751-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0567576A  
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 40MG Per day		Chills		Paxil	PS	Glaxosmithkline	ORAL
Initial or Prolonged 300MG Per day		Confusional State		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Coordination Abnormal Diarrhoea Pyrexia Serotonin Syndrome Tremor		Dextromethorphan	C	Glaxosmithkline	

Date:07/26/05ISR Number: 4724779-4Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0388621A  
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG Twice per day	4 DAY	Condition Aggravated Myopia Vision Blurred		Zyntabac	PS	Glaxosmithkline	ORAL

Date:07/26/05ISR Number: 4724783-6Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0388660A  
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression Anger		Zyban	PS	Glaxosmithkline	ORAL

Date:07/26/05ISR Number: 4725119-7Report Type:Expedited (15-DaCompany Report #GXKR2005US01328  
 Age:15 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	61 DAY	Anxiety Confusional State	Literature Health	Bupropion (Ngx)(Bupropion)	PS		
150 MG, BID		Convulsion Depression	Professional	Risperidone (Ngx) (Risperidone)	SS		
0.25 MG, BID	213 DAY	Drug Interaction Dry Mouth		Atomoxetine (Atomoxetine)	SS		
INTENTIONAL OVERDOSE OF 1200 MG		Electrocardiogram Qt Corrected Interval Prolonged Heart Rate Increased Intentional Misuse Overdose Postictal State Tremor		Alprazolam (Alprazolam)	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/26/05ISR Number: 4726101-6Report Type:Direct  
Age:52 YR Gender:Male I/FU:I

Company Report #CTU 254348

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Tinnitus		Wellbutrin Xr	PS		ORAL
UP TO 225 MG							
ORAL				Numerous Products	C		

Date:07/26/05ISR Number: 4727185-1Report Type:Direct  
Age:52 YR Gender:Male I/FU:I

Company Report #CTU 254325

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Nausea Vomiting		Bupropion	PS		
				Docusate Na	C		
				Trazodone Hcl	C		
				Oxycodone Hcl	C		
				Gatifloxacin	C		
				Bupropion Hcl	C		
				Clotrimazole	C		
				Zinc Oxide	C		
				Meloxicam	C		

Date:07/26/05ISR Number: 4727412-0Report Type:Expedited (15-DaCompany Report #2005085576  
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Creatine	Health	Zoloft (Sertraline)	PS		ORAL
200 MG (200							
MG, 1 IN 1		Phosphokinase Increased	Professional				
D),ORAL		Myalgia					
				Wellbutrin Xl (Bupropion Hydrochloride)	SS		
				Wellbutrin (Bupropion Hydrochloride)	C		

Date:07/27/05ISR Number: 4725946-6Report Type:Expedited (15-DaCompany Report #BPC-ZN-05-419  
 Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Pruritus Generalised	Zyban	PS	Glaxosmithkline	ORAL
150MG Twice							
			Rash				
per day	2	WK	Sensation Of Foreign Body				

Date:07/27/05ISR Number: 4725947-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0567258A  
 Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other			Abnormal Dreams	Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day	3	DAY					
			Anorexia	Micardis	C	Glaxosmithkline	
			Anxiety	Copaxone	C		
			Homicidal Ideation				
			Insomnia				
			Irritability				
			Suicidal Ideation				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/27/05ISR Number: 4725948-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0567272A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Wellbutrin	PS	Glaxosmithkline	NASAL
		Euphoric Mood					
		Incorrect Route Of Drug Administration					

Date:07/27/05ISR Number: 4725949-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0567282A

Age:11 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amenorrhoea		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Depression		Trazodone	C		
		Suicide Attempt					

Date:07/27/05ISR Number: 4725953-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0567599A

Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypoacusis		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Stress		Tenormin	C		
		Syncope Vasovagal		Topomax	C		
				Klonopin	C		

Date:07/28/05ISR Number: 4727253-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0567739A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blindness		Wellbutrin	PS	Glaxosmithkline	ORAL
150MG Per day		Iritis					

Date:07/28/05ISR Number: 4727273-XReport Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0388927A  
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination		Bupropion			
		Nausea		Hydrochloride	PS	Glaxosmithkline	
UNKNOWN	150MG	Per day 5 DAY					
		Pharyngolaryngeal Pain					
		Pyrexia					

Date:07/28/05ISR Number: 4728116-0Report Type:Direct Company Report #CTU 254657  
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Anger		Wellbutrin Xl 300mg.			
Hospitalization -		Balance Disorder		Glax	PS	Glax	
1 A DAY							
Initial or Prolonged		Bipolar Disorder					
Disability		Dysarthria					
		Loss Of Employment					
		Mania					
		Memory Impairment					
		Mental Disorder					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/29/05ISR Number: 4728733-8Report Type:Expedited (15-DaCompany Report #2005-03-0065  
Age:46 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300MG Unknown	Collapse Of Lung		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged SUBCUTANEOUS	Convulsion		Peg-Intron	SS		
	Pneumonia		Ribavirin	SS		ORAL
25MG Single dose	Vomiting		Paxil Cr	C	Glaxosmithkline	ORAL
.05MG Unknown			Levoxyl	C	Glaxosmithkline	ORAL
30MG Unknown			Prevacid	C		ORAL

Date:07/29/05ISR Number: 4728744-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0567927A  
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 150MG Per day 8 DAY	Abdominal Discomfort		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged	Cholelithiasis		Coumadin	C	Glaxosmithkline	
	Constipation		Hydrochlorothiazide	C		
			Metoprolol	C		
			Prevacid	C		

Date:07/29/05ISR Number: 4728745-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0567939A  
Age:29 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability 1TAB Per day 3 WK	Grand Mal Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Other	Loss Of Consciousness		Zyrtec	C	Glaxosmithkline	
	Tremor					

Date:07/29/05ISR Number: 4728746-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0567951A  
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day	3 YR	Amnesia		Aricept	C		
		Anger					
		Anxiety					
		Back Disorder					
		Back Pain					
		Blood Glucose Increased					
		Confusional State					
		Crying					
		Delirium					
		Hallucination					
		Paranoia					

Date:07/29/05ISR Number: 4728748-XReport Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0321501A  
Age:49 YR Gender:Male I/FU:F

Outcome	PT
Other	Akinesia
	Dyskinesia
	Dysphonia
	Hypokinesia
	Masked Facies
	Monoplegia
	Musculoskeletal Stiffness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Parkinsonism

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300MG Per day	19 DAY		Zyntabac	PS	Glaxosmithkline	

Date:07/29/05ISR Number: 4728756-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0386419A  
 Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1TAB See Initial or Prolonged dosage text	36 DAY	Bradycardia		Zyban	PS	Glaxosmithkline	ORAL
		Chest Pain Chest Wall Pain Cholelithiasis Constipation Hepatic Pain					

Date:07/29/05ISR Number: 4728768-5Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0388674A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 150MG Per day	5 DAY	Psoriasis		Zyban	PS	Glaxosmithkline	ORAL
		Rash Skin Exfoliation					

Date:07/29/05ISR Number: 4728771-5Report Type:Expedited (15-DaCompany Report #NL-GLAXOSMITHKLINE-B0389067A  
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN	150MG Per day	Depression Suicidal Ideation		Bupropion Hydrochloride	PS	Glaxosmithkline	
	4 DAY			No Concurrent			

Date:07/29/05ISR Number: 4734457-3Report Type:Expedited (15-DaCompany Report #S05-USA-03304-01  
Age:28 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Dizziness	Consumer	Citalopram	PS		
Initial or Prolonged	Dyskinesia		Wellbutrin			
	Epilepsy		(Bupropion			
	Fall		Hydrochloride)	SS		
	Fatigue					
	Grand Mal Convulsion					
	Headache					

Date:08/01/05ISR Number: 4730191-4Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0387737A  
Age:23 YR Gender:Female I/FU:F

Outcome	PT
Other	Crying
	Depression
	Dry Mouth
	Emotional Disorder
	Emotional Distress
	Hallucination, Auditory



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Headache Inappropriate Affect Insomnia				
		Lethargy Night Sweats Nightmare	Zyban	PS	Glaxosmithkline	ORAL

Date:08/01/05ISR Number: 4730196-3Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0388647A  
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Death		Death Intentional Misuse	Zyban Propranolol Benzodiazepine Alcohol	PS C C C	Glaxosmithkline	ORAL

Date:08/02/05ISR Number: 4738145-9Report Type:Expedited (15-DaCompany Report #B0321501A  
Age:49 YR Gender:Male I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Other		Parkinsonism	Zyban Tablet-Zyban (Bupropion Hydrochloride)	PS		
300 MG PER DAY		Foreign Literature Health Professional				

Date:08/03/05ISR Number: 4732508-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0568344A  
Age: Gender:Male I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Wellbutrin	PS	Glaxosmithkline	

Date:08/03/05ISR Number: 4732533-2Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0384877A  
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebral Haemorrhage		Zyban	PS	Glaxosmithkline	ORAL
150MG Twice		Cerebrovascular Accident					
per day	6	DAY					
		Convulsion					
		Subarachnoid Haemorrhage					

Date:08/03/05ISR Number: 4732744-6Report Type:Expedited (15-DaCompany Report #S05-USA-02982-01  
Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Dizziness		Citalopram	SS		ORAL
		Dyskinesia					
		Epilepsy					
		Fall					
		Fatigue					
		Grand Mal Convulsion					
		Head Injury					
		Postictal Headache					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/03/05ISR Number: 4732748-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLIN-A0568350A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death				Wellbutrin Xl	PS	Glaxosmithkline	ORAL
300MG See dosage text			Cardio-Respiratory Arrest				
			Convulsion				
			Overdose				

Date:08/03/05ISR Number: 4732752-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLIN-A0568459A

Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death				Wellbutrin	PS	Glaxosmithkline	ORAL
3 YR Hospitalization - Initial or Prolonged			Ill-Defined Disorder	Ultram	SS		

Date:08/03/05ISR Number: 4732934-2Report Type:Periodic Company Report #US-GLAXOSMITHKLIN-A0562033A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
				Avandamet	PS	Glaxosmithkline	ORAL
150MG Per day				Wellbutrin Xl	SS	Glaxosmithkline	ORAL
			Stomach Discomfort	Glucophage	SS		ORAL
3 YR				No Concurrent Medication	C		

Date:08/04/05ISR Number: 4734761-9Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLIN-B0387737A

Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Other 150MG Twice per day	11	DAY	Crying Depression Dry Mouth Emotional Disorder Emotional Distress Hallucination, Auditory Headache Insomnia Lethargy Night Sweats Nightmare	Zyban  Levlen Ed	PS  C	Glaxosmithkline	ORAL  ORAL
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Date:08/04/05ISR Number: 4734773-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0389004A  
Age:59 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Per day	2 DAY	Depressed Mood		Zyban	PS	Glaxosmithkline	ORAL
40MCG Per day		Head Discomfort		Ipratropium Bromide	C	Glaxosmithkline	
200MCG Per day		Sensory Disturbance		Salmeterol Beclomethasone	C C	Glaxosmithkline Glaxosmithkline	
125MCG per day				Thyroxine	C	Glaxosmithkline	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/04/05ISR Number: 4734774-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0389007A

Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Menorrhagia		Zyban	PS	Glaxosmithkline	ORAL
4 DAY							
		Rash		Atorvastatin	C		ORAL
20MG per day				Gaviscon	C	Glaxosmithkline	ORAL

Date:08/04/05ISR Number: 4734778-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0389361A

Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Consciousness		Zyban	PS	Glaxosmithkline	ORAL
		Syncope					

Date:08/05/05ISR Number: 4735616-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0563979A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Paxil	PS	Glaxosmithkline	
		Pituitary Tumour		Wellbutrin	SS	Glaxosmithkline	ORAL

Date:08/05/05ISR Number: 4735620-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0568718A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Macular Degeneration		Wellbutrin	PS	Glaxosmithkline	ORAL
400MG Per day							

Date:08/05/05ISR Number: 4735633-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0382344A

Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Epilepsy		Zyntabac	PS	Glaxosmithkline	
UNKNOWN	300MG per day						
Initial or Prolonged		Facial Bones Fracture					
Other		Fibula Fracture					
		Ill-Defined Disorder					

Date:08/05/05ISR Number: 4735643-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0389012A  
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Headache		Zyban	PS	Glaxosmithkline	ORAL
150MG per day	5 DAY						
		Tremor					
		Visual Disturbance					

Date:08/05/05ISR Number: 4736567-3Report Type:Direct Company Report #CTU 255513  
Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Convulsion		Wellbutrin 300mg			
Hospitalization -		Headache		Glascosmithkline	PS	Glascosmithkline	
ONCE DAILY							
Initial or Prolonged		Vomiting					
Required							
Intervention to							
Prevent Permanent							
Impairment/Damage							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/08/05ISR Number: 4736856-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0551834A  
 Age:30 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion Depressed Level Of Consciousness Dizziness Drug Toxicity Fall Headache Loss Of Consciousness Malaise Respiratory Arrest Tachycardia		Wellbutrin	PS	Glaxosmithkline	

Date:08/08/05ISR Number: 4736857-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0557105A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Life-Threatening Other		Death Overdose		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:08/08/05ISR Number: 4736858-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0559798A  
 Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day Initial or Prolonged Other		Abortion Induced Stillbirth		Wellbutrin	PS	Glaxosmithkline	ORAL

Date:08/08/05ISR Number: 4736859-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0559798B  
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death  
300MG Per day  
Congenital Anomaly

Abortion Induced

Alpha 1 Foetoprotein  
Abnormal  
Anencephaly  
Drug Exposure During  
Pregnancy

Wellbutrin

PS

Glaxosmithkline

Date:08/08/05ISR Number: 4736860-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0565550A  
Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Wellbutrin	PS	Glaxosmithkline	ORAL
18 MON		Drug Screen Positive Overdose					

Date:08/08/05ISR Number: 4736901-4Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0389840A  
Age: Gender:Male I/FU:F

Outcome	PT
Other	Abdominal Pain Blood Glucose Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		Blood Pressure Increased Diarrhoea Hallucination Headache	Zyban	PS	Glaxosmithkline	
		Hypertension Impaired Driving Ability Vomiting				

Date:08/09/05ISR Number: 4738368-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0555664A  
Age: Gender:Male I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		PT Coloboma Drug Exposure During Pregnancy Microphthalmos	Wellbutrin Xl	PS	Glaxosmithkline	

Date:08/09/05ISR Number: 4738372-0Report Type:Expedited (15-DaCompany Report #BPC-ZN-05-419  
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
150MG Twice per day	2 WK	PT Pruritus Generalised Rash Sensation Of Foreign Body	Zyban	PS	Glaxosmithkline	ORAL

Date:08/09/05ISR Number: 4738378-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0568459A  
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Death 3 YR Hospitalization - Initial or Prolonged		PT Death Drug Interaction Drug Level Increased Fall	Wellbutrin Ultram	PS SS	Glaxosmithkline	ORAL

Date:08/09/05ISR Number: 4738379-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0569020A  
 Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Murder		Paxil	PS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	

Date:08/09/05ISR Number: 4738383-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0569132A  
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Distension		Wellbutrin	PS	Glaxosmithkline	
		Adrenal Insufficiency		Wellbutrin	SS	Glaxosmithkline	ORAL
300MG Twice							
per day	2	YR					
		Bacterial Infection					
		Bruxism		Armour Thyroid	C		
		Diarrhoea		Fosamax	C		
		Fatigue		Multivitamins +			
		Flatulence		Supplements	C		
		Headache					
		Medication Error					
		Neck Pain					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/09/05ISR Number: 4738386-0Report Type:Expedited (15-DaCompany Report #BR-GLAXOSMITHKLINE-A0569142A  
 Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150MG Three Initial or Prolonged times per day 21 Disability 6MG Per day 51	DAY	DAY	Attention Deficit/Hyperactivity Disorder Coordination Abnormal Dizziness Headache Hypoaesthesia Sedation Weight Decreased	Zyban    Bromazepam	PS    C	Glaxosmithkline	ORAL    ORAL

Date:08/09/05ISR Number: 4738387-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0569146A  
 Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG Per day 6	MON		Convulsion Ill-Defined Disorder	Wellbutrin Xl	PS	Glaxosmithkline	ORAL

Date:08/09/05ISR Number: 4738407-5Report Type:Expedited (15-DaCompany Report #ZA-GLAXOSMITHKLINE-B0389923A  
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death UNKNOWN			Completed Suicide	Zyban	PS	Glaxosmithkline	

Date:08/09/05ISR Number: 4738408-7Report Type:Expedited (15-DaCompany Report #LU-GLAXOSMITHKLINE-B0389926A  
 Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Life-Threatening 15MG per day 1 DAY	Convulsion	Zyban	PS	Glaxosmithkline	ORAL
	Depressed Level Of Consciousness Multiple Drug Overdose Suicide Attempt	Alcohol	C		

Date:08/09/05ISR Number: 4738411-7Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0390060A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
UNKNOWN		Electric Shock		Zyban	PS	Glaxosmithkline	
		Hypervigilance Middle Insomnia Shock Tinnitus Tremor					

Date:08/10/05ISR Number: 4739091-7Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0554003A

Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
25MG Per day 10 MON		Abnormal Dreams		Paxil Cr	PS	Glaxosmithkline	ORAL
52 DAY		Insomnia		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Weight Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/10/05ISR Number: 4739135-2Report Type:Periodic  
 Age:48 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0555368A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
12.5MG Per day	2 YR	Distractibility		Paxil Cr	PS	Glaxosmithkline	ORAL
UNKNOWN	0 DAY	Disturbance In Attention		Wellbutrin	SS	Glaxosmithkline	
		Irritability		Lipitor	C		
		Mood Altered					
		Pharmaceutical Product Complaint					
		Therapeutic Response Unexpected					

Date:08/10/05ISR Number: 4739191-1Report Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0558860A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
YR		Adverse Drug Reaction		Paxil Cr	PS	Glaxosmithkline	ORAL
				Wellbutrin Xl	SS	Glaxosmithkline	ORAL

Date:08/10/05ISR Number: 4739237-0Report Type:Periodic  
 Age:51 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0561374A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
12.5MG Per day	2 YR	Depression		Paxil Cr	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK	Dizziness		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Drug Withdrawal Syndrome		Activella	C		
		Fatigue		Oxycontin	C		
		Hot Flush		Zyrtec	C	Glaxosmithkline	
		Hyperhidrosis		Ativan	C		
		Panic Attack					
		Tremor					

Date:08/10/05ISR Number: 4739294-1Report Type:Periodic  
Age:39 YR Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0540282A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Paxil	PS	Glaxosmithkline	ORAL
20MG Unknown	13 YR	Chest Pain		Wellbutrin	SS	Glaxosmithkline	
UNKNOWN		Depression		Ativan	C		
		Drug Withdrawal Syndrome					
		Feeling Jittery					
		Hyperhidrosis					
		Nervousness					
		Palpitations					
		Paraesthesia					
		Social Phobia					

Date:08/10/05ISR Number: 4739323-5Report Type:Periodic  
Age:54 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0541523A

Outcome	PT
	Body Temperature
	Decreased
	Drug Ineffective
	Skin Odour Abnormal
	Thirst
	Urine Odour Abnormal

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Weight Increased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
8 WK			Paxil	PS	Glaxosmithkline	ORAL
			Wellbutrin	SS	Glaxosmithkline	ORAL
			Neurontin	C		
			Wellbutrin	C	Glaxosmithkline	
			Advair	C	Glaxosmithkline	

Date:08/10/05ISR Number: 4739443-5Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0542872A  
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Paxil	PS	Glaxosmithkline	ORAL
150MG Per day	1 MON			Wellbutrin Xl	SS	Glaxosmithkline	ORAL
				Klonopin	C		
				Neurontin	C		
				Vitamin E	C		
				Multivitamin	C		
				Vitamin C	C	Glaxosmithkline	

Date:08/10/05ISR Number: 4739470-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0551298A  
 Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Paroxetine	PS	Glaxosmithkline	ORAL
50MG Per day		Drug Ineffective		Paxil	SS	Glaxosmithkline	ORAL
300MG Per day	5 YR			Wellbutrin	SS	Glaxosmithkline	ORAL
				No Concurrent Medication	C		

Date:08/10/05ISR Number: 4739600-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0553572A  
 Age: Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Fatigue		Paxil	PS	Glaxosmithkline	ORAL
	1TAB	Per day	Irritability		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
	150MG	Twice	Weight Increased					
		per day			Cozaar	C		
					Ultram	C		
					Blood Thinner	C		

Date:08/10/05ISR Number: 4739602-1Report Type:Periodic  
Age:39 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553586A

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Anxiety		Paxil	PS	Glaxosmithkline	ORAL
			Drug Ineffective		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
	300MG	Per day 5 MON	Panic Reaction		No Concurrent			
			Stress		Medication	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/10/05ISR Number: 4739749-XReport Type:Periodic  
 Age:58 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0557762A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
20MG Per day	7 YR	Drug Ineffective		Paxil	PS	Glaxosmithkline	ORAL
300MG At night	2 WK	Insomnia		Wellbutrin	SS	Glaxosmithkline	ORAL
		Libido Decreased					
		Libido Increased		Crestor	C		
				Aspirin	C	Glaxosmithkline	
				Metoprolol	C		

Date:08/10/05ISR Number: 4739824-XReport Type:Periodic  
 Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0559025A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
4 YR		Nervousness		Paxil	PS	Glaxosmithkline	ORAL
1 MON		Panic Reaction		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Tremor					
		Weight Increased					

Date:08/10/05ISR Number: 4739952-9Report Type:Periodic  
 Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0429084A

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
20MG Per day	12 MON	Crying		Paxil	PS	Glaxosmithkline	ORAL
4 WK		Drug Withdrawal Syndrome		Wellbutrin	SS	Glaxosmithkline	ORAL
		Emotional Disorder		Ativan	C		
		Headache					
		Migraine					
		Nausea					
		Paraesthesia					
		Photophobia					

Date:08/10/05ISR Number: 4740183-7Report Type:Periodic  
Age: Gender: I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0545471A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor		Paxil	PS	Glaxosmithkline	ORAL
				Wellbutrin	SS	Glaxosmithkline	ORAL

Date:08/10/05ISR Number: 4740219-3Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0547647A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anger		Paxil	PS	Glaxosmithkline	ORAL
2 YR		Depression		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
2 WK		Sexual Dysfunction					

Date:08/10/05ISR Number: 4740280-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0550474A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation		Paxil	PS	Glaxosmithkline	ORAL
150MG Unknown	WK	Female Orgasmic Disorder		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Irritability					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/10/05ISR Number: 4740685-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0546519A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
4 YR			Abnormal Dreams	Lamictal	PS	Glaxosmithkline	ORAL
			Adverse Event	Wellbutrin	SS	Glaxosmithkline	
			Hypnagogic Hallucination	Lithium	SS	Glaxosmithkline	
			Nightmare	Zoloft	C		
			Weight Increased				

Date:08/10/05ISR Number: 4740979-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0560142A

Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
20MG Per day 1 MON			Blood Amylase Increased	Paxil	PS	Glaxosmithkline	
150MG Per day 1 MON			Lipase Increased	Wellbutrin	SS	Glaxosmithkline	ORAL
				Hyzaar	C		

Date:08/10/05ISR Number: 4740982-1Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0560255A

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
20MG Unknown			Blood Amylase Increased	Paxil	PS	Glaxosmithkline	ORAL
			Lipase Increased	Wellbutrin	SS	Glaxosmithkline	ORAL
			Pancreatitis				

Date:08/10/05ISR Number: 4741010-4Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0560966A

Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
40MG Per day 11 YR			Anxiety	Paxil	PS	Glaxosmithkline	ORAL

200MG per day	2	WK	Depression	Wellbutrin	SS	Glaxosmithkline	ORAL
			Drug Ineffective	Protonix	C		
			Headache	Temazepam	C		

Date:08/10/05ISR Number: 4741046-3Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0561811A  
 Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness		Paxil	PS	Glaxosmithkline	ORAL
		Insomnia		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Per day	3	DAY		Prilosec	C	Glaxosmithkline	
				Valium	C		
				Ativan	C		

Date:08/10/05ISR Number: 4741147-XReport Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0537773A  
 Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Asthenia		Paxil	PS	Glaxosmithkline	ORAL
6 YR		Drug Ineffective		Wellbutrin	SS	Glaxosmithkline	ORAL
100MG Twice		Lethargy					
per day	9	MON		No Concurrent Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/10/05ISR Number: 4741261-9Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0538126A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anger		Paxil	PS	Glaxosmithkline	ORAL
		Anxiety		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG In the morning		Depression					
		Dizziness					
		Drug Withdrawal Syndrome					
		Headache					
		Heart Rate Increased					
		Paraesthesia					
		Tearfulness					

Date:08/10/05ISR Number: 4741278-4Report Type:Periodic  
Age: Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538318A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Feeling Abnormal		Paxil	PS	Glaxosmithkline	
UNKNOWN				Wellbutrin	SS	Glaxosmithkline	ORAL
				Geodon	C		

Date:08/10/05ISR Number: 4741286-3Report Type:Periodic  
Age:54 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0538553A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abnormal Dreams		Paxil	PS	Glaxosmithkline	ORAL
40MG Per day 3 YR		Drug Ineffective		Wellbutrin	SS	Glaxosmithkline	ORAL
150MG Per day		Dry Mouth		Klonopin	C		
				Topamax	C		
				Amitriptyline	C		
				Aspirin	C	Glaxosmithkline	
				Verapamil	C		
				Vitamin B12	C	Glaxosmithkline	

TRANSDERMAL

Date:08/11/05ISR Number: 4742205-6Report Type:Periodic  
Age:58 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0550467A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
12.5MG Per				Paxil Cr	PS	Glaxosmithkline	ORAL
day	13	DAY					
150MG Unknown				Wellbutrin Xl	SS	Glaxosmithkline	ORAL
				Paxil	SS	Glaxosmithkline	

Date:08/11/05ISR Number: 4742252-4Report Type:Periodic  
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0551724A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
25MG Per day	1	YR		Paxil Cr	PS	Glaxosmithkline	ORAL
150MG Per day	4	DAY		Wellbutrin Xl	SS	Glaxosmithkline	ORAL

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Freedom Of Information (FOI) Report

Date:08/11/05ISR Number: 4742274-3Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0552172A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
25MG Per day	3	YR	Chest Pain	Paxil Cr	PS	Glaxosmithkline	ORAL
150MG Per day	9	MON	Depression	Wellbutrin Xl	SS	Glaxosmithkline	ORAL
			Dizziness	Xanax	C		
			Insomnia	Verapamil	C		
				Hct	C		
				Metformin	C		
				Amaryl	C		
				Actos	C		
				Adderall Xr	C		
				Theophylline	C		
				Albuterol	C	Glaxosmithkline	
				Tylenol With Codeine	C		
				Trazodone	C		

Date:08/11/05ISR Number: 4742342-6Report Type:Periodic  
Age: Gender:Female I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0553408A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
12.5MG Per			Pruritus Generalised	Paxil Cr	PS	Glaxosmithkline	ORAL
day	3	YR					
200MG Per day	3	YR		Wellbutrin Sr	SS	Glaxosmithkline	ORAL

Date:08/11/05ISR Number: 4742686-8Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20050801049  
Age: Gender:Female I/FU:I

Company Report #US-JNJFOC-20050801049

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death			Accidental Overdose	Ultram	PS		ORAL
			Respiratory Distress	Wellbutrin	SS		
				Medication Nos	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Agitation		Depakote	PS		
Initial or Prolonged	Anxiety		Bupropion			
Other	Depression		Hydrochloride	SS		ORAL
	Drug Ineffective		Bupropion			
	Hepatocellular Damage		Hydrochloride	SS		
	Social Phobia		Lamotrigine	SS		
	Suicidal Ideation		Lithium	C		
			Actos	C		
			Glipizide	C		
			Levothyroxine Sodium	C		
			Irbesartan	C		
			Clonazepam	C		
			Zolpidem Tartrate	C		
			Amitriptyline			
			Hydrochloride	C		
			Propranolol			
			Hydrochloride	C		
			Fluoxetine			
			Hydrochloride	C		
			Venlafaxine			
			Hydrochloride	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Estradiol	C
Cholestyramine	C
Irbesartan	C
Zolpidem Tartrate	C
Amitriptyline	
Hydrochloride	C
Propranolol	
Hydrochloride	C
Fluoxetine	
Hydrochloride	C
Venlafaxine	
Hydrochloride	C
Oestradiol	C

Date:08/12/05ISR Number: 4743715-8Report Type:Expedited (15-DaCompany Report #13148  
 Age:26 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Tachycardia		Wellbutrin	PS	Glaxosmithkline	
150MG Per day	30 DAY			Lexapro	SS		
10MG per day	134 DAY			Unspecified Ssri	C		

Date:08/12/05ISR Number: 4743737-7Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0356916A  
 Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous		Zyban	PS	Glaxosmithkline	ORAL
150MG See							
dosage text	50 DAY			Infertility Therapy			
				Unspecified	C		

Date:08/12/05ISR Number: 4743742-0Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0385105A  
 Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Chest Pain		Zyban	PS	Glaxosmithkline	ORAL
1TAB Per day	5 DAY						
Initial or Prolonged		Dizziness		Valerian	C	Glaxosmithkline	ORAL
3U per day	11 DAY						
		Hypertension					
		Palpitations					
		Supraventricular					
		Extrasystoles					

Date:08/12/05ISR Number: 4743753-5Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0389332A

Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Interaction		Zyban	PS	Glaxosmithkline	ORAL
150MG Per day	3 WK						
100MG Four		Hallucination, Visual		Tramadol	C		ORAL
times per day				Zopiclone	C		ORAL
7.5MG At							
night				Loratadine	C		ORAL
10MG Per day				Cetirizine	C	Glaxosmithkline	ORAL
10MG Per day				Co-Codamol	C		ORAL
				Diclofenac	C		ORAL
50MG Three							
times per day				Propranolol	C		ORAL
80MG Twice							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

per day  
 RESPIRATORY  
 (INHALATION) 400MCG Twice  
 Beclometasone C Glaxosmithkline

per day  
 RESPIRATORY  
 (INHALATION) 12MCG  
 Formoterol C  
 Variable dose

Date:08/12/05ISR Number: 4743760-2Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0390296A  
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Homicidal Ideation Paraesthesia		Zyban	PS	Glaxosmithkline	ORAL

Date:08/12/05ISR Number: 4743761-4Report Type:Expedited (15-DaCompany Report #AU-GLAXOSMITHKLINE-B0390305A  
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination Vision Blurred		Zyban	PS	Glaxosmithkline	ORAL

Date:08/15/05ISR Number: 4744822-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0553401A  
 Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Excoriation		Wellbutrin	PS	Glaxosmithkline	ORAL
11 DAY		Fall		Effexor Xr	C		ORAL
300MG In the		Grand Mal Convulsion					
morning							

45MG At night	Hand Fracture	Remeron	C	ORAL
UNKNOWN	Myalgia	Premarin	C	
	.125MG			
Unknown	Pain In Extremity			
	Periorbital Haematoma			

Date:08/15/05ISR Number: 4744826-3Report Type:Expedited (15-DaCompany Report #2005-07-0898  
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Depression		Wellbutrin	PS	Glaxosmithkline	ORAL
		Personality Change		Pegylated Interferon			
		Restless Legs Syndrome		Alpha 2b	SS		
UNKNOWN		Schizophrenia		Rebetol	SS		ORAL
		Suicidal Ideation					

Date:08/15/05ISR Number: 4744831-7Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0570056A  
Age:16 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Convulsion		Wellbutrin	PS	Glaxosmithkline	ORAL
Hospitalization -		Overdose					
Initial or Prolonged		Suicide Attempt					
Disability							
Other							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/16/05ISR Number: 4745915-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0565250A

Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Exposure During Pregnancy		Lithium	PS	Glaxosmithkline	
		Hypertension		Bupropion	SS	Glaxosmithkline	
300MG per day				Wellbutrin	SS	Glaxosmithkline	
		Renal Failure		Lexapro	C		
30MG per day							
		Renal Function Test		Nortriptyline	C		
100MG Per day							
		Abnormal		Topamax	C		
150MG per day							
		Renal Impairment					

Date:08/16/05ISR Number: 4745923-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0570086A

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Acne		Bupropion	PS	Glaxosmithkline	
89 DAY							
		Drug Exposure During Pregnancy		Terbutaline	C		
		Uterine Contractions During Pregnancy		Magnesium	C		

Date:08/16/05ISR Number: 4745925-2Report Type:Expedited (15-DaCompany Report #2005-08-0496

Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Wellbutrin	PS	Glaxosmithkline	
UNKNOWN							
		Condition Aggravated		Peg-Intron	SS		
SUBCUTANEOUS							
		Suicidal Ideation		Rebetol	SS		ORAL

Date:08/17/05ISR Number: 4746709-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0564635A  
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aggression Agitation Anxiety Completed Suicide Crying Emotional Disorder Irritability Social Avoidant Behaviour Tremor		Wellbutrin	PS	Glaxosmithkline	

Date:08/17/05ISR Number: 4746713-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0569266A  
Age:66 YR Gender:Male I/FU:F

Outcome	PT
Other	Aggression Altered Visual Depth Perception Apathy Asthenia Balance Disorder Confusional State Coordination Abnormal Dizziness Drug Ineffective

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Interaction	Report Source	Product	Role	Manufacturer	Route
3 MON		Early Morning Awakening Headache					
		Insomnia		Lamictal	PS	Glaxosmithkline	ORAL
150MG Per day		Temperature Intolerance		Wellbutrin Xl	SS	Glaxosmithkline	ORAL
		Visual Acuity Reduced		Clonazepam	SS		

Date:08/17/05ISR Number: 4746717-0Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0570240A  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Per day 5 DAY Initial or Prolonged		Asthenia		Wellbutrin	PS	Glaxosmithkline	ORAL
		Contusion		Ambien	C		
		Convulsion		Lotrel	C		
		Dysarthria		Depakote	C		
		Fall		Clonazepam	C		
		Feeling Drunk		Vitamins	C		
		Muscular Weakness					

Date:08/17/05ISR Number: 4746719-4Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0370023A  
Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 150MG Twice per day	18 DAY	Arthralgia		Zyban	PS	Glaxosmithkline	ORAL
		Discomfort					
		Hypersensitivity Scar					

Date:08/17/05ISR Number: 4746733-9Report Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0390517A  
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Abdominal Pain Upper	Zyban	PS	Glaxosmithkline	ORAL
150MG Unknown 10 DAY					
	Chest Pain				
	Constipation				
	Diplopia				
	Dry Mouth				
	Headache				
	Hemiparesis				
	Insomnia				
	Palpitations				
	Suicidal Ideation				

Date:08/18/05ISR Number: 4748055-9Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0570232A  
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability							
150MG Per day	4	MON	Gastrooesophageal Reflux	Wellbutrin	PS	Glaxosmithkline	ORAL
			Disease	Wellbutrin	SS	Glaxosmithkline	ORAL
75MG Per day	4	MON	Medication Error	No Concurrent Medication	C		



FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/18/05ISR Number: 4748058-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0570423A  
 Age:40 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 150MG Per day 4 MON		Anxiety	Wellbutrin	PS	Glaxosmithkline	ORAL
Initial or Prolonged		Blood Pressure Increased Chest Pain Feeling Jittery Insomnia	No Concurrent Medication	C		

Date:08/18/05ISR Number: 4748059-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0570427A  
 Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 450MG Per day		Bipolar Disorder	Wellbutrin	PS	Glaxosmithkline	ORAL
		Muscle Twitching	Prozac	C		

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Summary report for FOI selections:

Selection by inexact search of active ingredient:

BUPROPION%

Selection by inexact search of Tradename/Verbatim:

WELLBUTRIN%

Total number of reports: 18,667

From: 01-NOV-1997

To: Present

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